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Pattern of antibiotic prescription in the management of endodontic infections amongst oral and maxillofacial surgeons and endodontists in Gujarat

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Abstract

Aim
To identify antibiotic prescription in the treatment of endodontic infections amongst Oral Surgeons and Endodontists in Gujarat.

Methodology
The Oral Surgeons & Endodontists registered with the Gujarat State Dental Council were surveyed on antibiotic prescription for six different pulpal and periapical diagnoses. A total of 70 questionnaires were delivered with 59 returned (84%).

Results
The average duration of antibiotic therapy was 7.0 ± 1.0 days. Ninety five percent of respondents selected amoxicillin as the first choice of antibiotic in patients with no medical allergies, alone (34%) or associated to clavulanate (61%). The first drug of choice for patients with an allergy to penicillins was clindamycin (65%), followed by azithromycin (15%) and metronidazole (13%). For cases of irreversible pulpitis, 86% of respondents prescribed antibiotics. For the scenario of a necrotic pulp, acute apical periodontitis and no swelling, 71% prescribed antibiotics. Almost 60% of respondents prescribed antibiotics for necrotic pulps with chronic apical periodontitis and a sinus tract.

Conclusions
The majority of the Oral Surgeons & Endodontists in Gujarat were selecting the appropriate antibiotic for use in endodontic infections, but there are still many who are prescribing antibiotics inappropriately. The use of antibiotics for minor infections, or in some cases in patients without infections, could be a major contributor to the world problem of antimicrobial resistance.

Keywords
Apical periodontitis, endodontic infections, irreversible pulpitis, orofacial infections, pharmacology.

Introduction
Antibiotics are prescribed by the dentists for treatment as well as prevention of infection. Indications for the use of systemic antibiotics in dentistry are limited, since most of the dental and periodontal diseases are best managed by operative intervention and oral hygiene measures. However, the literature provides evidence of inadequate prescribing practices by the dentists, due to a number of factors ranging from inadequate knowledge to certain social factors.

Dentists prescribe medications for the management of a number of oral conditions, mainly orofacial infections. Since most human orofacial infections originate from odontogenic infections, the prescription of antibiotics by dental practitioners has become an important aspect of dental practice. For this reason, antibiotics account for the vast majority of medicines prescribed by the dentists. Dentists prescribe between 7% and 11% of all common antibiotics (beta-lactams, macrolides, tetracyclines, clindamycin, and metronidazole). In the UK, for instance, dentists accounted for 7% of all community prescriptions of antimicrobials. On the other hand, the National Centre for Disease Control and Prevention estimated that approximately one-third of all outpatient antibiotic prescriptions were unnecessary. Al-Haroni & Skau in 2007 analysed 268,834 prescriptions issued by 4,765 dentists and showed that the dentists' prescriptions of antibiotics contributed 8% of the total national consumption in Norway.

Yingling et al. in 2002 concluded that the majority of the members of the AAE were selecting the appropriate antibiotic for use in orofacial infections, but there were still many who are prescribing antibiotics inappropriately. Antibiotic prescribing may be associated with unfavourable side effects ranging from gastrointestinal disturbances to fatal anaphylactic shock and development of resistance. Cars et al. in 2001, concluded that Spain had one of the highest antibiotic consumption rate in Europe, therefore, has the highest percentages of bacterial resistance.

The increasing resistance problems of recent years are probably related to misuse of broad-spectrum agents such as cephalosporins and fluoroquinolones. We now have entered an era where some of the bacterial species are resistant to the full range of antibiotics. The methicillin-
resistant Staphylococcus aureus being the most widely known example of extensive resistance\footnote{Dentistry’ contributes to the problem of antibiotic resistance substantially, because dentists prescribe approximately 10% of all common antibiotics.}

To our best knowledge, data of antibiotic prescription by Gujarati Oral Surgeons & Endodontists is scarce. Therefore, this study was carried out to determine antibiotic prescription trends among them.

**Methodology**

A cross sectional survey was conducted in the state of Gujarat which included all the Oral Surgeons & Endodontists. Candidates were requested to answer a one-page questionnaire which surveyed the pattern of use of antibiotics in the treatment of endodontic infections. The questions were based on the previous surveys developed in the USA (Whitten et al. 1996, Yingling et al. 2002) and Spain (Rodriguez-Nunez et al. 2009). Seventy questionnaires were delivered to the Oral Surgeons & Endodontists of Gujarat. Only 59 questionnaires were returned completed (84%).

A database was created for further analysis, using version 15.0 of the Statistical Package for Social Sciences (SPSS; SPSS Inc., Chicago, IL, USA). Data description was carried out by frequency tables. When obtaining the numerical representation by percentages, the total number of answers for each query was taken into account. Data was analyzed using descriptive statistics, chi square test of independence and logistic regression. Statistically significant differences were considered for P < 0.05.

**Results**

The demographics of the respondents are described in the Table 1. Male respondents accounted for 49% and females 51% of the total. Seventy six percent of the respondents were less than 36 years old and 11% more than 45 years old. The mean age of the respondents was 34 years. The average duration of antibiotic therapy was 7.0 ± 1.0 days (Fig. 2). The standard deviation in this response indicated that majority prescribed for a period of 6 to 8 days. There were no significant differences amongst respondents in relation with age and gender. Most of respondents (95%) chose amoxicillin in non-allergic patients (Table 2), alone (40%) or associated to clavulanic acid (61%). Amoxicillin/Clavulanic acid 875/125 mg was prescribed as first choice antibiotic by 42% of respondents, where as 18%, 9%, 3% and 1% selected amoxicillin 750 mg, amoxicillin 500 mg, clindamycin and metronidazole, respectively. The first drug of choice for patients with an allergy to penicillin was clindamycin 300 mg (65%), followed by azithromycin (15%) and metronidazole (13%) (Table 3). Table 4 lists the percentage of respondents who prescribed antibiotics for various pulpual and periapical diagnoses. For cases of irreversible pulps with moderate/severe symptoms and irreversible pulps with acute apical periodontitis and moderate/severe symptoms, 32% and 54% of respondents, respectively, prescribed antibiotics. In cases of a necrotic pulp, chronic apical periodontitis, no swelling and no other symptoms, antibiotics were prescribed by 31%. In the scenario of necrotic pulp, acute apical periodontitis, moderate/severe symptoms but no swelling, 71% prescribed antibiotics. For a case of necrotic pulp, chronic apical periodontitis, asymptomatic but with sinus tract, 60% prescribe antibiotics. In the case of a necrotic pulp, acute apical periodontitis, swelling and other moderate/severe symptoms, 95% of respondents prescribed antibiotics.

**Discussion**

In our study, the questions and the six endodontic treatment situations proposed were based on those asked in previous surveys developed in the USA (Whitten et al. 1996, Yingling et al. 2002) and Spain (Rodriguez-Nunez et al. 2009). The overall response rate of 84% can be considered to be an acceptable rate of return for surveys. In relation with antibiotic therapy, an endodontic infection must be persistent or systemic to justify the need for antibiotics, i.e. fever, swelling, lymphadenopathy, trismus or malaise in a healthy patient (Yingling et al. 2002). Endodontic infections typically have a rapid onset and short duration, 2–7 days or less, particularly if the cause is treated or eliminated (Pallasch 1993). The average length of antibiotic prescriptions in this study was 7.0 ± 1.0 days, in accordance with the result (6.8 days) reported previously by Rodriguez-Nuñez et al. (2009) amongst Spanish Endodontists. The proper dose and duration of an antibiotic are enough when there is sufficient evidence that the patient host defenses have gained control of the infection. When the infection is resolving or has resolved, then the drug should be terminated (Pallasch 1993, Yingling et al. 2002). A 6- to 7-day course would probably be appropriate for most endodontic infections. An antibiotic loading dose should be used whenever the half-life of the antibiotics longer than 3 h or whenever a delay of 12 h or more is unacceptable to achieve therapeutic blood levels (Montgomery & Kroeger 1984). Confusion about prescribing antibiotics and inappropriate prescribing practices, however, were reported by respondent dentists. The majority of endodontic infections resolve in 3–7 days (Epstein et al. 2000); thus, the 18.0% of respondents who routinely prescribe antibiotics for more than 7 days should reassess how they prescribe antibiotics.

The first drug of choice for patients with an allergy to penicillins was clindamycin (65%), in accordance with the result previously found amongst Spanish Endodontists (63%) (Rodriguez-Nuñez et al. 2009). In the United States, the study of Whitten et al. (1996) reported a 21.6% for clindamycin as first choice antibiotic, but the study
carried out by Yingling et al. (2002) found a percentage (57.03%) similar to that reported in this study.

**Conclusions**

The majority of the Oral Surgeons & Endodontists in Gujarat were selecting the appropriate antibiotic for use in endodontic infections, but there are still many who are prescribing antibiotics in appropriately. The use of antibiotics for minor infections, or in some cases inpatients without infections, could be a major contributor to the world problem of antimicrobial resistance.

**References**

Histoplasmosis of the periodontium – An uncommon condition

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Abstract
Histoplasmosis is caused by the fungus Histoplasma capsulatum, a dimorphic fungus that grows in the yeast form in infected tissue. The Periodontium consists of investing and supporting tissue of the tooth: gingival, periodontal ligament, cementum and alveolar bone. This report describes an unusual form of presentation of Histoplasmosis of gingiva including palate and retromolar region, which could have been mistaken as an inflammatory swelling or tubercular ulcer unless histological examination was carried out. The man was non-diabetic & Enzyme-Linked – Immunosorbent assay (ELISA) test for Human Immunodeficiency Virus was negative. During further evaluation the patient was found to have hyponatraemia and bilateral adrenal mass. He developed adrenal crisis and was managed with hydrocortisone and sodium supplementation. He has been successfully treated with Amphotericin B followed by Itraconazole. Periodontal treatment includes proper brushing instruction, 0.2%, 10 ml chlorhexidine oral rinse twice daily for 30 days.

Key words
Periodontium, Histoplasmosis, Histoplasma capsulatum.

Introduction
Histoplasmosis, a systemic mycosis resulting from inhalation of the spores of the fungus Histoplasma capsulatum, was first described in 1906 in Panama. The organism is found in acidic soils.
Histoplasmosis is clinically classified as a primary acute pulmonary form (usually asymptomatic); a chronic pulmonary form (occurs in the presence of underlying pulmonary disease) and a disseminated form (exclusively in infants, the elderly, and in debilitated or immunocompromised patients). The latter is characterized by the progressive spread of infection to extra pulmonary sites, and the lesions in this form may be extra pulmonary in the oral cavity or intestine.

The disseminated form of histoplasmosis (DH) is most common1 (affected age group over 54 years). In 1985, DH was added to the spectrum of ‘AIDS-defining’ diseases2. Rare oral lesions are present as ulcer, granulomas or verrucous and plaque like lesion 3 and have been detected up to 50% of individuals with DH and HIV infection4.

Case Report
A 51-year-old man reported to the Department of Periodontics of Dr. R. Ahmed Dental College and Hospital, with the chief complain of inability to take food for about last 2-3 months due to excessive swollen gums and burning sensation and painful ulceration on palate which started insidiously and increased gradually.

Clinical examination revealed, scattered red and white granulomatous lesions on the gingiva in the right maxillary region (Fig.1) and in the both retromolar regions showing the most extensive involvement (Fig. 2 & 3). The labial gingiva exhibited patches of sloughing epithelium (Fig. 1). The palatal side of right maxillary region showing (Fig. 4) erythematos, ulcerative lesion. Patient’s oral hygiene was poor with moderate to heavy plaque and generalized inflammation.

Patient gave history of smoking and beetle-nut chewing for past 10 years.

Medical history revealed that he was suffering with intermittent low grade fever (100ºF/99ºF) coming every
week for last one month. There was gradual loss of weight and generalized weakness, cough and cramps in both lower extremities for about last 3 months. Patient gave history of fever and cough 1 year ago, which was treated by local physician. Investigation showed Hb-8.9 gm%, TLC-4200, N-71%, L-24%, M-2%, E-3%, ESR-103 mm/1st hour. His Post Prandial blood sugar was 102 mg%. His Post Prandial blood sugar was 102 mg%. Sputum for AFB was negative.

Routine periodontal treatment include proper brushing instruction, 10 ml, 0.2% chlorhexidine oral rinse twice daily for 1 month and supragingival scaling showed no improvement of the gingival and palatal lesion.

Incisional biopsy was taken from palatal lesion under local anesthesia. Histological examination of the biopsy specimen revealed section showed multiple fragments of tissue infiltrated by sheets of histiocytes admixed with chronic inflammatory cells. Some of the histiocytes contain fungal bodies resembling Histoplasma capsulatum. Each of these microorganisms was composed of an eosinophilic core surrounded by a clear halo and showed positive staining with Giemsa, methenamine silver consistent with Histoplasma capsulatum(Fig. 9).

The patient was referred to the School of Tropical Medicine (STM), Kolkata for HIV – Screening test as the diagnosis of Histoplasmosis often follows the diagnosis of HIV – infection.

The patient was diagnosed as HIV Negative by ELISA.

During further evaluation at STM, the patient was found to have bilateral cervical lymphadenopathy, enlargement of liver up to 4 cms, hyponatremia and bilateral adrenal
mass in Ultrasonography of abdomen. Bone marrow has been aspirated and sent for fungal culture which showed growth of fungus.

The patient developed adrenal crisis and was managed with hydrocortisone and sodium supplementation. He has been successfully treated with Amphotericin B followed by Itraconazole (Fig 5 to 8).

Discussion

Histoplasmosis is a potentially serious condition. The occurrence of oral manifestation of histoplasmosis in non-HIV infected patients has been demonstrated in several case reports and in case series. Padhye et al. (1994) described oral histoplasmosis cases from India. Studies demonstrate the extremely low prevalence of oral histoplasmosis in South – East Asia.

In conclusion, histoplasmosis, particularly oral histoplasmosis, seems to be rare in HIV-negative individuals of south and southeast region of Asia. Diagnosis depends on demonstration of the organisms by histology and culture of tissue specimens. Definitive diagnosis is usually made by a combination of culture, detection of the organism in tissues, measurement of antibodies, or detection of antigen. (last two tests are not available here.) Therapy is achieved by administration of Amphotericin B followed by Itraconazole or Itraconazole6.

References

Operationalisation and Utilisation of AYUSH Clinics in Chandigarh, India: A cross sectional evaluation study

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Abstract
National Rural Health Mission (NRHM) strategy of mainstreaming AYUSH health facilities, require evaluations for continual improvement.

Objectives
To assess operationalisation and utilization of AYUSH clinics, to ascertain the level of patient satisfaction, to explore the perceptions of AYUSH practitioners regarding implementation of AYUSH scheme and to assess the factors influencing AYUSH scheme implementation.

Material and Methods
Cross-sectional study design and mixed methodology was followed that included record review, and interviews with AYUSH service users and the service providers, of both Allopathic and AYUSH streams.

Results
AYUSH system has been implemented successfully in Chandigarh and people are using these services. Females, prior users of allopathic medicine and suffering from chronic problems were the main users. Client satisfaction with service provision by AYUSH clinics was high. However, allopathic practitioners are not much enthusiastic about recognizing these systems as scientific therapies. More publicity about AYUSH in the region is required as the main goal of integration is not being fulfilled.

Key Words
AYUSH, Ayurveda, NRHM, Health, Evaluation.

Introduction
National Rural Health Mission (NRHM) envisaged total functional integration between AYUSH dispensaries/ hospitals with the allopathic system to improve access to wider spectrum of treatments at affordable costs¹. There is a need for evaluation and conduct surveys to understand the lacunae in service delivery. Therefore this study was conducted with the objectives to know how successful Chandigarh health sector has been in achieving mainstreaming of AYUSH systems under the NRHM umbrella, what are the satisfaction levels of patients regarding these services, how allopathic as well as AYUSH practitioners perceive the scheme and what are the underlying factors affecting the implementation of this scheme in Chandigarh.

Material and Methods
Study design and sampling: Cross sectional study was done during August 2009 – December 2010, in two community health centres, where both allopathic clinics and AYUSH clinics (NRHM) were operational under one roof. Study participants were a) The AYUSH practitioners employed under NRHM by Chandigarh administration and their collocated allopathic counterparts and b) patients attending AYUSH clinics. Sample size of 96 patients was calculated for patients interview, based on the assumption of 60% patient satisfaction level 3, 95% confidence level, and beta error of 5%. A total of 100 patients available at the institutional level were interviewed.

Study tools and their administration: Self-administered questionnaires were given to the doctors of both the streams to explore their perceptions regarding the AYUSH scheme. Questionnaire for AYUSH doctors had questions on overview about various issues pertaining to the work pattern, constraints faced, if any, etc. For Allopathic doctors, 5-item semi-structured questionnaire was administered to know their perceptions on the integration of AYUSH systems, their attitude towards cross-referral between two systems, trainings and their role in health service provision to the community.

Interview schedule was administered to the patient respondents. It covered the socio demographic profile of the respondents, followed by a brief history of the disease for which the patient attended the AYUSH clinic and allopathic treatment, if taken. Questions on the satisfaction levels of patients regarding services available at the clinic, attitude of doctors, infrastructure and maintenance of the clinic were also included in the schedule in addition to their willingness to recommend these services to their friends and relatives etc. Likert scale in the form of money scale was used to record the satisfaction levels of the patients. Each item statement was read to the patient, who would indicate his/her level of agreement with the statement by mentioning a money
amount. The interview schedule contained 12 items. Record review was done to get information regarding utilization of the AYUSH scheme. All the information sought from the officials was marked in the checklists. Morbidity data of AYUSH as well as Allopathic clinics was compiled from the records.

The study was approved by Institutional ethics committee and informed consent was obtained from all participants.

Results

Infrastructure and operationalisation

Set up of the AYUSH clinics was as per Indian Public Health Standards (IPHS) except for posting of AYUSH specialist and space for pharmacy cum store in one CHC (Table 1).

AYUSH professionals were involved in various health societies, but their involvement in the national health programmes was very minimal. They had received some trainings on some programmes and they were involved in conducting health educational activities such as organisation of AYUSH Melas, health camps etc. There were not involved in planning, and managerial functions at the CHC levels.

Socio-economic and morbidity profile of the patients

Socioeconomic profile of the patients is given in Table 2. Ayurvedic clinics catered to 30.6 per cent of the total patient load whereas, 68.6 per cent of the patients were registered at homeopathic clinics of these centres.

One year (2009-10) morbidity analysis revealed that more patients attended these clinics for chronic problems (Figure 1). Homeopathy was more popular for allergic/skin disorders, whereas, Ayurvedic stream was preferred for joint problems and digestive system/diarrhoeal diseases (17.9%). Ranking of morbidity, body system wise was done and top 10 ailments in both allopathic and AYUSH streams are given in Table 3. Whereas, allopathic systems were preferred for surgical and orthopaedic problems requiring acute care or for infections, AYUSH systems were preferred for chronic skin problems and joint problems.

Record review for the period of three months (January-March, 2010) revealed only 17 per cent of patients used AYUSH services. Over the period of three months, old registration was more in Ayurvedic and Homeopathic streams as compared to Allopathy and; more females attended AYUSH as well as Allopathic clinics than males. (Figure 2)

77 per cent (95% CI 67.5-84.8) of the patients had taken allopathic treatment for the same health problem and 29 per cent of them had taken the allopathic medicines for

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Table 1: IPHS Standards Compliance Chart of AYUSH clinics

<table>
<thead>
<tr>
<th>Services to be delivered at CHC:</th>
<th>IPHS Standards:</th>
<th>Status at selected health centres:</th>
</tr>
</thead>
<tbody>
<tr>
<td>AYUSH Specialist (Post-graduate in AYUSH)</td>
<td>One</td>
<td>None</td>
</tr>
<tr>
<td>AYUSH General duty Medical Officer (Graduate in AYUSH)</td>
<td>One</td>
<td>One each clinic.</td>
</tr>
<tr>
<td>AYUSH Pharmacist</td>
<td>One</td>
<td>One each clinic.</td>
</tr>
<tr>
<td>Pharmacy cum store</td>
<td>6.4 x 3.2 m</td>
<td>CHC 1: 2.5 x 3 m CHC 2: 6 x 3 m</td>
</tr>
<tr>
<td>Space for each AYUSH doctor’s room</td>
<td>3.2 x 3.2 m</td>
<td>CHC 1: 4 x 3 m CHC 2: 5 x 4 m</td>
</tr>
<tr>
<td>AYUSH drugs</td>
<td>As per the list in annexure</td>
<td>Available as per the list.</td>
</tr>
</tbody>
</table>

Table 2: Socioeconomic characteristics of the patients attending AYUSH clinics (n= 100)

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Percentage of population</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
</tr>
<tr>
<td>0-10 years</td>
<td>8</td>
</tr>
<tr>
<td>11- 20 years</td>
<td>14</td>
</tr>
<tr>
<td>21-30 years</td>
<td>21</td>
</tr>
<tr>
<td>31-40 years</td>
<td>28</td>
</tr>
<tr>
<td>41- 50 years</td>
<td>15</td>
</tr>
<tr>
<td>51- 60 years</td>
<td>7</td>
</tr>
<tr>
<td>61-70 years</td>
<td>6</td>
</tr>
<tr>
<td>71-80 years</td>
<td>1</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>27</td>
</tr>
<tr>
<td>Female</td>
<td>73</td>
</tr>
<tr>
<td>Occupation</td>
<td></td>
</tr>
<tr>
<td>House wife</td>
<td>51</td>
</tr>
<tr>
<td>Business</td>
<td>4</td>
</tr>
<tr>
<td>Government job</td>
<td>11</td>
</tr>
<tr>
<td>Laborer</td>
<td>1</td>
</tr>
<tr>
<td>Non worker</td>
<td>14</td>
</tr>
<tr>
<td>Private job</td>
<td>19</td>
</tr>
<tr>
<td>Level of education</td>
<td></td>
</tr>
<tr>
<td>Illiterate</td>
<td>10</td>
</tr>
<tr>
<td>Primary education</td>
<td>23</td>
</tr>
<tr>
<td>Higher secondary</td>
<td>12</td>
</tr>
<tr>
<td>Matriculation</td>
<td>20</td>
</tr>
<tr>
<td>Graduation</td>
<td>24</td>
</tr>
<tr>
<td>Post graduation</td>
<td>10</td>
</tr>
</tbody>
</table>

Figure 1: Bar diagram representing comparative morbidity profile of patients of AYUSH clinics
years altogether, before switching over to AYUSH clinic. Out of the patients who had availed allopathic treatment before, 73 per cent (95% CI: 63.2-81.4, n=77) of them did not tell their allopathic practitioner about the switchover to AYUSH. Sixty five per cent (95% CI: 53.8-73.4) of total patients were referred to these clinics. Of these 49% (95% CI: 53.8-73.4) of the referrals were made by friends/relatives. Only 4% of the referrals were made by allopathic doctors. Among patients, who were not referred by anyone, source of information was variable, with 24% coming on their own, and receptionist sending 11% (95% CI: 5.6-18.8) of the patients due to non-availability of an allopathic doctor. Advertisements (newspapers and televisions) contributed 8% of the patients who had come to the clinic without any referral. Only 1% of the patients came by reading the information board displayed in the dispensary.

**Why a switch over?**

Side effects and non-effectiveness of the allopathic medicines were quoted as the major reasons for switching over from allopathic to AYUSH clinic, by 40% and 54% of the patients respectively. Only 3% of the patients cited non-satisfaction with allopathic doctor’s behaviour as the reason for switching over to AYUSH clinic. 51% (95% CI: 40.8-61.1) of the patients attending clinic had no prior experience of AYUSH system of medicine.

**Satisfaction level of patients with different aspects of services being provided**

High satisfaction to the level of above 80% was expressed by patients regarding doctor’s communication with respect to diagnosis, treatment plan, follow up plan, and precautions need to be taken; however regarding medicines being provided, less than half of the patients were satisfied to this much extent (Table 4).

**Discussion**

This study conducted to assess the status of operationalization of AYUSH scheme in Chandigarh, its utilisation by patients and perceptions of AYUSH and Allopathic practitioners about the scheme has given some important insights.

First, Chandigarh is on track for mainstreaming AYUSH, by putting necessary infrastructure and manpower in place. Involvement of AYUSH practitioners in various national health programmes and health educational activities is an encouraging sign. Major shortfalls in the operationalization of the AYUSH scheme are that; specialists in the AYUSH stream have not been engaged as yet, and medicine supplies are not sufficient. According to the IPHS standards there should be at least one specialist in each AYUSH dispensary. Consequently, speciality services like Panchkarma, Ksharasutra, etc. are not being provided by these facilities and the AYUSH doctors are merely functioning as general practitioners. Unlike an unpublished report on mainstreaming of AYUSH in four states of India in 2009, there are no major bottlenecks with respect to infrastructure, manpower and materials except for having more medicines and specialists.

Secondly, patient satisfaction with AYUSH services was found to be very high. There seems to be no major issue with respect to sitting place, cleanliness and examination facilities. However, not many were satisfied with the medicines and with the ultimate relief from the problem they were suffering with.

Most of these patients shifted to AYUSH after trying...
allopathic treatment. This may have happened due to greater and reliable access to this alternate treatment. However, inability to provide relief to these patients may initiate reverse referral. Non relief may be due to non-availability of specialist doctors and specialised treatment therapies.

Thirdly, study has further shown poor referral by allopathic doctors. It may be due to their misperceptions and non-orientation to AYUSH system. In our study also one of the allopathic doctors expressed his distrust for AYUSH system. Such perceptions can be changed through sensitization workshops and repeated dialogues.

There are some noteworthy observations regarding extent of AYUSH service utilisation and the factors influencing the utilisation. Annual AYUSH clinics utilisation of 17 per cent was less than the utilisation of CAM use in Singapore, Israel, Japan and India. In our study, there was higher usage of Homeopathy as compared to the Ayurvedic system, contrary to other available evidences where Ayurveda was found to be more popular.

Increased utilisation of AYUSH services over time may reflect their dissatisfaction in some way with conventional treatment. More than three-fourth of the patients in the present study had taken allopathic medicines and were dissatisfied, prior to shifting over to the AYUSH systems. Higher cost of treatment was not the major reason for switching over to other systems. In the present study, more than half of the patients were found to have switched over to AYUSH systems of medication due to the non-effectiveness of the allopathic medicine, whereas 40% did so due to their side effects. Modern medicine is considered to bring only symptomatic improvement and is not curative of the underlying problem. Patients while shifting, preferred not to inform their allopathic doctors, as was observed in other studies.

Patients with chronic diseases were found to be more likely to use AYUSH services, as was found in other studies. For pregnancy related issues and deliveries, women had to resort to the co-located allopathic system. AYUSH practitioners were neither providing any emergency services nor any specialized services to patients.

Education has inconsistent association with use of CAM. Our findings of higher AYUSH use among lower income groups, was also inconsistent with the other available evidence. It may be due to overall lack of utilization of public services by high income group population.

In conclusion, AYUSH systems of medicines have been implemented successfully in Chandigarh and people are using these services. There is need to generate more awareness among allopathic practitioners, and appoint specialist AYUSH practitioners.

Conflict of interest: None declared

Funding: None

References


<table>
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<tr>
<th>S. No.</th>
<th>Parameters measured</th>
<th>1-20 paisa</th>
<th>21-40 paisa</th>
<th>41-60 paisa</th>
<th>61-80 paisa</th>
<th>81 paisa-1 Re</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>With the doctor’s communication with respect to</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a.</td>
<td>Diagnosis</td>
<td>0%</td>
<td>1%</td>
<td>5%</td>
<td>17%</td>
<td>77%</td>
</tr>
<tr>
<td>b.</td>
<td>The treatment plan</td>
<td>1%</td>
<td>2%</td>
<td>10%</td>
<td>19%</td>
<td>68%</td>
</tr>
<tr>
<td>c.</td>
<td>The follow up plan</td>
<td>0%</td>
<td>2%</td>
<td>5%</td>
<td>31%</td>
<td>62%</td>
</tr>
<tr>
<td>d.</td>
<td>The precautions that need to be taken</td>
<td>2%</td>
<td>3%</td>
<td>8%</td>
<td>26%</td>
<td>50%</td>
</tr>
<tr>
<td>2.</td>
<td>With the medicines given*</td>
<td>1%</td>
<td>1%</td>
<td>17%</td>
<td>25%</td>
<td>44%</td>
</tr>
<tr>
<td>3.</td>
<td>With sitting place in clinic</td>
<td>0%</td>
<td>5%</td>
<td>10%</td>
<td>36%</td>
<td>49%</td>
</tr>
<tr>
<td>4.</td>
<td>With the cleanliness</td>
<td>0%</td>
<td>2%</td>
<td>11%</td>
<td>38%</td>
<td>49%</td>
</tr>
<tr>
<td>5.</td>
<td>With the examination facility</td>
<td>0%</td>
<td>0%</td>
<td>10%</td>
<td>29%</td>
<td>61%</td>
</tr>
<tr>
<td>6.</td>
<td>Relief felt from the problem**</td>
<td>4%</td>
<td>17%</td>
<td>19%</td>
<td>26%</td>
<td>11%</td>
</tr>
<tr>
<td>7.</td>
<td>Will come again to avail services here</td>
<td>1%</td>
<td>2%</td>
<td>34%</td>
<td>55%</td>
<td>8%</td>
</tr>
<tr>
<td>8.</td>
<td>Will recommend this treatment to others</td>
<td>1%</td>
<td>3%</td>
<td>31%</td>
<td>57%</td>
<td>8%</td>
</tr>
<tr>
<td>9.</td>
<td>Overall satisfaction with AYUSH doctor’s services at the health centre</td>
<td>0%</td>
<td>0%</td>
<td>16%</td>
<td>44%</td>
<td>40%</td>
</tr>
</tbody>
</table>

*12% did not respond ** 33% gave no response (or had just started treatment)
tertiary care hospital for management of osteoarthritis (knee)-


ABO blood group and secretor status in IHD patients

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Abstract
Various studies have been done trying to relate ABO blood groups with diseases e.g. Peptic ulcer, duodenal ulcer, pernicious anaemia, gallstones, carcinoma of stomach and IHD. The same blood group antigens are secreted in the body fluids. The blood group antigens have protective action on its secreting mucosa. Some studies have related secretor status with various diseases. Very few studies have been done in India, to relate IHD with ABO blood groups and secretor status. The present study was undertaken to determine, whether the correlation exists between IHD and ABO blood group and secretor status.

Two hundred IHD patients admitted in SJIC, Bangalore were studied for blood group and secretor status and compared with age and sex matched controls. Among the control group ‘O’ was most common while in the IHD patients, an overall excess of blood group ‘A’ was found. In the gender difference serum HDL values were significantly higher in females than males. Serum triglyceride values were significantly higher in blood group ‘A’ compared to other groups. IHD cases have been divided into MI and non MI group and its relation has been observed with blood group. There was significant excess of Myocardial infarction in blood group ‘O’ and ‘B’.

In this study we have not observed any significant relation of IHD with secretor status.

Key Words
Ischemic Heart Disease; ABO Blood group; secretor status.

Introduction
ABO agglutinogens are not strictly confined to the red cells alone but they could also be demonstrated in other body tissues and are secreted in body fluids. The presence of A and B antigens in the saliva of A and B group individuals, was first discovered in 1930 by Lehr and Putkomen who classified the world population into two groups, secretors and non-secretors.

Obesity, dyslipidemia, hypertension, diabetes mellitus, smoking, stress, sedentary lifestyle and genetic factors are some well known risk factors for IHD. Considering ABO blood group phenotype and secretor status as a part of the genetic factors, this study was undertaken to observe the influence of genetic factors on IHD.

Knowledge of secretor status can be applied for tissue transplantation among other applications. Numerous studies have been reported, correlating the frequency of diseases with the ABO blood groups and secretor status. Several studies have suggested with the ABO blood group phenotype and secretor status of individuals are associated with an increased risk of Ischaemic Heart Disease. Ischaemic Heart Disease has emerged as the number one killer disease, the world over, including India. The association between ABO blood group and secretor status and IHD in India, was recognized as an important lacuna to be filled. Hence, the need for such a study.

Material and Methods
Subjects recruited for this study were Ischaemic Heart Disease (IHD) patients, in Sri Jayadeva Institute of Cardiology, Bangalore, Karnataka. Patients were diagnosed by cardiologist on the basis of clinical history, ECG, blood investigations, enzyme markers and angiography. Two hundred patients were studied, one hundred and sixty were males and forty were females, in the age group of 35 to 65 years. IHD patients with other systemic disorders (eg. CVA, COPD, endocrine disorders) except Diabetes Mellitus and Hypertension were excluded.

Blood group of volunteer blood donors in the St. Martha’s hospital and Sri Jayadeva Institute of Cardiology blood bank, Bangalore, Karnataka were included as controls. Two hundred blood donors (age and sex matched) with no evidence of any disease were included in the control group.

ABO blood group: ABO blood group was determined, using slide agglutination technique¹. Secretor status was examined using the agglutination inhibition technique¹. Saliva was collected in between 9 AM to 11AM. before breakfast after rinsing the mouth. The sample was examined within one hour.

If the blood group antigens are present in saliva, when an appropriate antiseraum is added to it, an antigen – antibody reaction occurs. The antibodies in the serum neutralise the antigens in saliva. When a red cell suspension of the same blood group is now added to this mixture, there is no agglutination due to previous inhibition of the antisemur. Thus in the case of secretors
there will be no agglutination seen. In the case of non-secretors, there will be an agglutination reaction.

Lipid profile: Total serum cholesterol, serum triglycerides, low density lipoproteins (LDL) and high density lipoproteins (HDL), were estimated in milligram percent, using the autoanalyser, (Hitachi LTD – Tokyo – Japan 704-1703 Sr. No. 0391 – 1) and expressed in milligram percent. The hemoglobin percent was done by laboratory technician and reports were collected from laboratory.

Statistical analysis done using SPSS version-10.6. One-way analysis of variance (ANOVA) and the Chi-square tests were used to analyze the data.

Results

Distribution of ABO blood group in the case group was; ‘O’ group 88(44%), ‘B’ group 66 (33%), ‘A’ group 37 (18.5%) and ‘AB’ group 9 (4.5%). Among the control group blood group ‘O’ is the commonest followed by B, A and AB. In IHD group, overall the number of patients with blood group ‘O’ were highest. Compared to controls, there was an overall excess of blood group ‘A’ and the difference was statistically significant. There was a significant deficit in blood group ‘O’.

Lipid profile values obtained were higher in females, as compared to males but this difference was statistically significant only for serum HDL.

The mean serum cholesterol level was greater in patients belonging to blood group-A, compared to the other group but this was not statistically significant.

<table>
<thead>
<tr>
<th>Blood Group</th>
<th>Lipid profile</th>
<th>Serum cholesterol (mg%)</th>
<th>Serum triglycerides (mg%)</th>
<th>LDL (mg%)</th>
<th>HDL (mg%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>O n=77</td>
<td></td>
<td>191.9 + 44.1</td>
<td>191.9 + 9.5</td>
<td>119.3 + 38.5</td>
<td>32.9 + 5.9</td>
</tr>
<tr>
<td>A n=62</td>
<td></td>
<td>201.1 + 30.7</td>
<td>242.2 + 105.2*</td>
<td>121.2 + 32.5</td>
<td>32.6 + 6.3</td>
</tr>
<tr>
<td>B n=46</td>
<td></td>
<td>191.8 + 41.2</td>
<td>165.5 + 65.9</td>
<td>128.9 + 33.9</td>
<td>30.2 + 8.2</td>
</tr>
<tr>
<td>AB n=15</td>
<td></td>
<td>181.6 + 31.6</td>
<td>180.9 + 87.7</td>
<td>140.9 + 45.1</td>
<td>32.9 + 6.3</td>
</tr>
</tbody>
</table>

The mean serum triglycerides level was greater in patients belonging to blood group-A, compared to the other groups and this was statistically significant.

The mean hemoglobin level was 13.19+1.84 gm%.

The chi-square test shows, that the difference between patients presenting with myocardial infarction, versus those without infarction was statistically significant. There was a significant excess of ‘O’ in patients with myocardial infarction.

Lipid profile showed higher values for serum cholesterol, serum triglycerides and LDL in secretors, compared to nonsecretors but this was not statistically significant.

Discussion

The controls in our study showed a proportionate frequency of the ABO blood groups as follows: the maximum number belonged to the ‘O’ group (44%), followed by ‘B’ group (33%), ‘A’ group (18.5%) and ‘AB’ group (4.5%). This is similar to the findings of KRC Naidu in Bellary, Karnataka2.

In the IHD group, the overall frequency of ‘O’ group was higher (38.5%), followed by ‘A’ (31%), ‘B’ (23%) and ‘AB’ (12.5%) groups. As compared to the controls, an overall excess was found in ‘A’ group and an overall deficit in the ‘O’ group.

The frequency of secretors noted was 73.29% and of nonsecretors 26.71%, in the control population. In our study, in the IHD group 77.5% were secretors and 22.5% non-secretors.

Table I: Distribution of ABO blood group among controls and IHD cases

<table>
<thead>
<tr>
<th>Blood Group</th>
<th>Observed number (O)</th>
<th>Expected number (E)</th>
<th>O – E</th>
</tr>
</thead>
<tbody>
<tr>
<td>O</td>
<td>Cases 77</td>
<td>82.5</td>
<td>-5.5</td>
</tr>
<tr>
<td></td>
<td>Controls 88</td>
<td>82.5</td>
<td>+5.5</td>
</tr>
<tr>
<td>A</td>
<td>Cases 62</td>
<td>49.5</td>
<td>+12.5*</td>
</tr>
<tr>
<td></td>
<td>Controls 37</td>
<td>49.5</td>
<td>-12.5</td>
</tr>
<tr>
<td>B</td>
<td>Cases 46</td>
<td>56</td>
<td>-10</td>
</tr>
<tr>
<td></td>
<td>Controls 66</td>
<td>56</td>
<td>+10</td>
</tr>
<tr>
<td>AB</td>
<td>Cases 15</td>
<td>12</td>
<td>+3</td>
</tr>
<tr>
<td></td>
<td>Controls 9</td>
<td>12</td>
<td>-3</td>
</tr>
</tbody>
</table>

Table II: Lipid profile in different blood groups in IHD patients (One-way ANOVA)

<table>
<thead>
<tr>
<th>Blood Group</th>
<th>IHD Group</th>
<th>Observed number (O)</th>
<th>Expected number (E)</th>
<th>O – E</th>
</tr>
</thead>
<tbody>
<tr>
<td>O n=77</td>
<td>MI</td>
<td>50</td>
<td>38.5</td>
<td>+11.5*</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>27</td>
<td>38.5</td>
<td>-11.5</td>
</tr>
<tr>
<td>A n=62</td>
<td>MI</td>
<td>37</td>
<td>31</td>
<td>+6</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>25</td>
<td>31</td>
<td>-6</td>
</tr>
<tr>
<td>B n=46</td>
<td>MI</td>
<td>30</td>
<td>23</td>
<td>+7**</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>16</td>
<td>23</td>
<td>-7</td>
</tr>
<tr>
<td>AB n=15</td>
<td>MI</td>
<td>9</td>
<td>7.5</td>
<td>+1.5</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>6</td>
<td>7.5</td>
<td>-1.5</td>
</tr>
</tbody>
</table>

Table III: Distribution of IHD patients in MI and Non-MI groups

<table>
<thead>
<tr>
<th>Blood Group</th>
<th>IHD Group</th>
<th>Observed number (O)</th>
<th>Expected number (E)</th>
<th>O – E</th>
</tr>
</thead>
<tbody>
<tr>
<td>O n=77</td>
<td>MI</td>
<td>41</td>
<td>31.5</td>
<td>+9.5*</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>22</td>
<td>31.5</td>
<td>-9.5</td>
</tr>
<tr>
<td>A n=48</td>
<td>MI</td>
<td>31</td>
<td>24</td>
<td>+7**</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>71</td>
<td>24</td>
<td>-7</td>
</tr>
<tr>
<td>B n=38</td>
<td>MI</td>
<td>27</td>
<td>19</td>
<td>+8***</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>11</td>
<td>19</td>
<td>-8</td>
</tr>
<tr>
<td>AB n=11</td>
<td>MI</td>
<td>7</td>
<td>5.5</td>
<td>+1.5</td>
</tr>
<tr>
<td></td>
<td>Non-MI</td>
<td>4</td>
<td>5.5</td>
<td>-1.5</td>
</tr>
</tbody>
</table>
were nonsecretors. This was similar to the findings in other studies in India and western countries. Age and gender are considered to be risk factors for atherosclerosis. Males are at a higher risk than females, but the gap between sexes narrows as age increases.

The human male in his early and mid-adult years is several times more likely to develop atherosclerosis than the female, suggesting that the male sex hormones might be atherogenic or conversely, that female sex hormones might be protective. In our study it was observed that 80% were males and 20% were females.

Experimental studies suggest that excess blood levels of iron can lead to atherosclerosis, perhaps by forming free radicals in the blood that damage the vessel wall. In our study none of the IHD patients were anaemic.

Blood groups are one of the most prominent inherited characters. Therefore, if genetic factors influence diseases, the susceptibility to IHD should also be reflected in the pattern of blood group distribution. This has been found to be the case by several workers.

Among the control group the blood group ‘O’ was the commonest, while in the IHD series of patients, an excess of blood group ‘A’ as compared to controls, was found. This agrees with the findings of other workers. In our study the serum cholesterol was on the higher side in blood group ‘A’ patients, than in other groups but the difference was not statistically significant. The analysis of variance showed that the difference for lipid profile values, between blood groups was statistically significant for serum triglycerides. Serum triglyceride values were higher in blood ‘A’ patients. However there is a less certain association between triglycerides and coronary heart disease (as compared to LDL cholesterol). O’Brein and Patton proposed that the reason for the association between ABO blood groups and serum cholesterol levels, may probably be related to the site of cholesterol in the erythrocyte membrane, of which the ABO antigen forms a part and this supposes the ABO locus itself, as the source of association.

Mehta et al., found that blood group ‘A’ neonates have a high serum cholesterol and low serum phospholipid levels. It can be suggested that an increase in cholesterol, which is hydrophobic and a decrease in phospholipid, obstruct the transport of lipids. Thus high serum cholesterol and a low serum phospholipid level should make an individual more prone for development of atherosclerosis in the future. Though our study does not include serum phospholipid estimation, cholesterol levels were found to be higher in group ‘A’ subjects, which supports the above hypothesis.

In our study, there was an excess of blood group ‘A’ and a deficit of blood group ‘O’ among IHD patients and this difference in the two groups was found to be statistically significant. This finding is similar to those reported by Denborough, Beg et al., Srivastava et al. and Bronte Stewart et al. A comparative table of the ABO blood group distribution reported by different authors and in our study is shown below.

Table V: Secretor status in male and female patients

<table>
<thead>
<tr>
<th>Sex</th>
<th>Secretor status</th>
<th>Secretors</th>
<th>Non-secretors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>Secretors</td>
<td>127 (79.4)</td>
<td>33 (20.6)</td>
</tr>
<tr>
<td></td>
<td>Non-secretors</td>
<td>28 (70)</td>
<td>12 (30)</td>
</tr>
<tr>
<td>Female</td>
<td>Secretors</td>
<td>28 (70)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Non-secretors</td>
<td>12 (30)</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>Secretors</td>
<td>155 (77.5)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Non-secretors</td>
<td>45 (22.5)</td>
<td></td>
</tr>
</tbody>
</table>

Table VI: Lipid profile and secretors status in IHD patients (One-way ANOVA)

<table>
<thead>
<tr>
<th>Secretor Status</th>
<th>Lipid profile</th>
<th>Serum Cholesterol (mg%)</th>
<th>Serum Triglycerides (mg%)</th>
<th>LDL (mg%)</th>
<th>HDL (mg%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Secretors</td>
<td>196 + 40.2</td>
<td>207 + 98.3</td>
<td>125.2 + 38.4</td>
<td>33.3 + 6.2</td>
<td>33.29 + 7.9</td>
</tr>
<tr>
<td>Non-Secretors</td>
<td>184.8 + 33.2</td>
<td>178.08 + 76.5</td>
<td>118.5 + 29.8</td>
<td>33.29 + 7.9</td>
<td>33.29 + 7.9</td>
</tr>
</tbody>
</table>

Table VII: Blood Group Distribution in Various Studies of IHD

<table>
<thead>
<tr>
<th>Study</th>
<th>Blood Group</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Group</td>
</tr>
<tr>
<td>M. Beg et al 1987</td>
<td>Cases</td>
</tr>
<tr>
<td></td>
<td>Controls</td>
</tr>
<tr>
<td>D.K. Srivastava et al 1966</td>
<td>Cases</td>
</tr>
<tr>
<td></td>
<td>Controls</td>
</tr>
<tr>
<td>M A Denborough 1962</td>
<td>Cases n=205</td>
</tr>
<tr>
<td></td>
<td>Controls n=17,700</td>
</tr>
<tr>
<td>B Bronte Stewart et al 11 1962</td>
<td>Cases</td>
</tr>
<tr>
<td></td>
<td>Controls</td>
</tr>
<tr>
<td>Present study 2005</td>
<td>Cases</td>
</tr>
<tr>
<td></td>
<td>Controls</td>
</tr>
</tbody>
</table>
On the other hand as these studies were confined mainly to survivors of IHD, the finding could have another interpretation. Group A and B individual are more likely to survive, than group ‘O’. A large prospective study is required to elucidate this association. The following possibilities may be considered (i) the genetic influence may be operating, by determining the anatomy of coronary vessels, or their liability to the atherosclerotic process or (ii) Group ‘O’ in some unknown way offers a protective substance, which reduces the incidence of coronary heart disease.

There was an excess of group ‘O’ patients presenting with myocardial infarction. When compared with group ‘A’. Our results differ from those of Allan15. They found an excess of ‘A’ group patients presenting with MI compared to ‘O’ group patients with IHD. Sweeney et al also found lower level of factor VIII: and vWF: Ag in group ‘O’ which increases bleeding tendency15. The presence of blood group ‘A’ determinants on the vWF locus is known and ‘A’ antigen is also expressed on platelets. The amount of blood group ‘A’ substance on platelets is proportional to the amount present in plasma. The presence of ‘A’ antigen interferes with the interaction between platelets and vWF. In group ‘O’ individuals lacking the A determinant, platelets would aggregate more effectively and occlude the vessels.

It has been observed, that blood group ‘O’ subjects have lower levels of antihemophilic globulin (factor VIII) and are more likely to bleed. Hence it can be concluded, that atherosclerotic patients with blood group ‘O’, might not tend to thrombose but rather hemorrhage into the arterial walls and sustain more tissue damage, leading to myocardial infarction.

A recent study reported that ‘non-O’ individuals have higher levels of vWF and fVIII than ‘O’ individuals due to prolonged clearance rates of vWF17.

‘Non-O’ individuals could have a high risk of venous thromboembolism via having greater level of fVIII or vWF. AsfVIII levels are determined mainly by vWF, higher levels of vWF might explain the higher incidence of venous thromboembolism among non-O individuals.

Small intestinal alkaline phosphatase is known to play an important role in handling of fat. The occurrence of small intestinal alkaline phosphatase activity in serum has been found to depend on the ABO blood group and ABH secretor status of individuals.

Thus group ‘A’ non-secretors might be more prone to atherosclerosis because serum alkaline phosphatase activity is altered and hence fat metabolism will also be altered in these individuals.

In our study we have not found any significant difference in the secretor status, between the IHD and control group. However the IHD patients show a preponderance of secretors. This result is similar to the finding of Hall et al in peripheral arteriosclerosis17 and contrasts with Langman’s results.

**Conclusion**

The group ‘A’ individuals appear more prone to IHD. Since the study was conducted in survivors of IHD, the progress of the disease might be slower in group ‘A’ than in ‘O’. The frequency of secretor status was similar in case and control series, indicates no influence of secretor status in IHD in our study.

**References**

12. Srivastava DK, Thakur CP and Das M. ABO blood groups in relation to Ischaemic Heart Disease. Indian Heart J. 1966; 18:140-149.
Central Adiposity and General Adiposity as Major Risk Predictors of Various Life Styles Related Diseases especially Hypertension and Type 2 DM

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Abstract

The functional roles of Waist circumference (WC), Waist and hip ratio (W/H ratio) and Body mass index (BMI) as risk factors for hypertension and type 2 diabetes mellitus (DM) in the Nomads Tribal population of Rajasthan State are evaluated in the paper. Four hundred and forty-five participants, aged between 18 to 100 years old are taken as subjects for scientific study. The prevalence of pre-hypertension (34.4%), hypertension (22.5%), prediabetes 5.2% (n=23) and diabetes (approximately 3.4 %) (n =15) is noted in the select population. Body mass index, waist circumference, and waist/hip ratio in the study are shown to be associated with type 2 DM and hypertension. Women with W/H ratio > 85+ and men with W/H ratio > 1.0+ are shown to be prone to increased risk of diabetes. And subjects having body mass index (BMI) particularly in the range of (obese) >25 appear to be vulnerable to hypertension. Similarly subjects with greater waist circumferences look to be exposed to hypertension. The study in the paper also supports the observation that both BMI and WC yield almost the same results. The findings suggests that by keeping a check on modifying risk factors like BMI, WC and W/H ratio we can control the prevalence of life styles associated diseases like hypertension and type 2 DM.

Key Words

Nomads, adiposities, incidence, SBP, DBP and FBG levels.

Introduction

The prevalence of type 2 DM is rising throughout the world, most rapidly in populations like India which are undergoing the epidemiologic transition from a mainly rural subsistence economy to an increasingly urban industrial economy1,2. The associations of increased weight and body fat composition with non-communicable diseases, particularly cardiovascular diseases and diabetes, are well established3. Obesity has become a major worldwide epidemic affecting more than 300 million people. It is an important risk factor for type 2 DM, a chronic disorder of carbohydrate, fat, and protein metabolism. From the clinical perspective, visceral adipose tissue is known to generate diabetogenic substance4 and, as such, may be more informative than total fat for diagnostic evaluation.

Body mass index, waist circumference, and waist/hip ratio have been shown to be associated with type 2 DM and hypertension. As expected, epidemiologic studies have demonstrated that these three obesity indicators are strong and consistent predictors of lifestyles related diseases like type 2 DM and hypertension. However, despite the clear, clinical difference between visceral and other forms of fat, little epidemiologic difference would be expected in the relation of diabetes with body mass index versus waist hip ratio. From a statistical perspective, the two measures yield similar information, with the correlation coefficient typically about 0.85. Several studies have shown that WC is a better predictor of type 2 DM, than is BMI, but these findings are inconclusive,6-8 while other studies provide evidence that W/H ratio has a positive effect independent of BMI9-11.

In addition, the ability of these obesity indicators to predict diabetes may differ by ethnicity, age, and sex12-14. For example, among Asian populations, central obesity has been shown to be a more consistent predictor of diabetes than is total obesity11,15, while general obesity has been shown to be a better predictor among white US populations and Europeans16,17. To compare associations of hypertension and type 2 DM incidence with general and central obesity indicators, we conducted a pilot study based on Tribal population of Rajasthan in India. Not a single study has been done so far on the population chosen so ours is the first attempt to study the role of central adiposity and general adiposity as indicators for the lifestyle related diseases especially diabetes and hypertension.

Study design and Data collection

Present search was limited to tribal population of Jhunjhunu district of Rajasthan state in India. Data was collected through snow and ball method of sampling. We had gone to twelve to sixteen camps and collected data of nearly four hundred and forty-five participants. The study was approved by the institutional human ethics committee at BITS, Pilani and performed according to the Declaration of Helsinki. All study members received detailed explanation of the study in their regional language before giving consent. A survey questionnaire was designed and finalized after a field trial. Men and women ≥18 years of age were considered eligible except pregnant women, seriously ill subjects, and those
who were on herbal medication or on drugs such as corticosteroids and oral contraceptive pills. Each subject was asked to report at a selected investigation site after an overnight fast.

Each participant was interviewed to know about occupation, education, housing, sanitation, family income and number of members in the family. Their status of physical activities, family history of diabetes, hypertension, and other kinds of diseases was also taken into account for this study. Other investigations included anthropometry, systolic and diastolic blood pressure, fasting plasma glucose and random plasma glucose test. Measurements of height, weight, and waist and hip circumference were taken with light clothes and without shoes. The weighing tools were calibrated daily by known standard weights. For taking standing height, the measurement has been taken with the help of stadiometer. Waist circumference was measured at the belly button or just above it. Similarly, the hip circumference was measured at its widest part. Blood pressure was taken after a 10-min rest with digital blood pressure machine. Classification of hypertension was based on ADA guidelines (a) Healthy blood pressure: <120.80 (b) Pre-hypertension: between 120/80 and 140/90 (c) Hypertension: 140/90 or higher. For diabetes the diagnostic criteria of the American Diabetes Association were used. Statistical analysis: Linear regression analysis and scatter plots was drawn to find out the correlation between associated risk factors and the prevalence of type 2 DM and hypertension.

**Results**

Overall, mean and standard deviation values for each variables i.e. FBG levels, systolic blood pressure(SBP), diastolic blood pressure(DBP), and general, central adiposity and dietetic pattern were available for all 445 participants given in Table 1.

As can be seen from Table 2 a higher number of females were overweight (12.4%) and obese (17.6 %) as compared to males. It was observed that there was (11.3%) obese female among the participants. In overweight category 13.1% of participants were having early high blood pressure and 17 % obese subjects were having early high blood pressure (pre-hypertension). Similarly in overweight category 11% of participants were having high blood pressure and 23 % obese respondents were having high blood pressure (hypertension). The subjects found positive for diabetes had BMI in the range of 23-25 (33.3%) and obese persons had BMI more than 25 or greater (20 %). Those subjects that found at risk zone had BMI in the range of 23-25 (13.3%) and obese persons had BMI more than 25 or greater (17.4%). Females were more prone to increased Body mass index (BMI) related diseases as compared to males. Similarly it was noticed that females had more W/H ratio i.e. 67.6% as compared to males (10.3%). Waist to hip ratio has been divided into three categories i.e. low risk, moderate risk and high risk. The participants come under high risk category having cut off w/h ratio (85+) cm for females and (1.0+) for males. In high risk group, 31.7% of subjects had early high blood pressure and 28.6 % had high blood pressure, 33.3% of them had positive blood glucose test and 56.5% were at risk zone of pre-diabetes. The waist circumference was also found to be high in case of females (30%) as compared to males i.e. (12.8%). Thirty three (32%) subjects having circumferences higher than 80 cm for females and 90 cm for men had early high blood pressure and 33% of

**Table 1:** Descriptive Statistics of study participants

<table>
<thead>
<tr>
<th>Parameters (n=445)</th>
<th>Data</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yrs) – Mean+ SD</td>
<td>45.39 +18.39</td>
</tr>
<tr>
<td>Sex- n (%)</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>250 (50.2)</td>
</tr>
<tr>
<td>Female</td>
<td>195 (43.8)</td>
</tr>
<tr>
<td>Height (cm) – Mean + SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>160.21 + 9.36</td>
</tr>
<tr>
<td>Female</td>
<td>154.37 + 11.18</td>
</tr>
<tr>
<td>Weight (cm) – Mean + SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>53.73 +3.96</td>
</tr>
<tr>
<td>Female</td>
<td>20.94 +3.96</td>
</tr>
<tr>
<td>BMI(kg/m2) – Mean + SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>23.0 +2.4</td>
</tr>
<tr>
<td>Female</td>
<td>20.94 +3.96</td>
</tr>
<tr>
<td>Waist circumference(cm) – Mean+SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>74.74 + 12.35</td>
</tr>
<tr>
<td>Female</td>
<td>79.45 + 13.4</td>
</tr>
<tr>
<td>Hip circumference (cm) –Mean+SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>84.19 + 11.88</td>
</tr>
<tr>
<td>Female</td>
<td>85.17 + 12.35</td>
</tr>
<tr>
<td>Waist to hip ratio(cm) – Mean+SD</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>0.8899 + .08335</td>
</tr>
<tr>
<td>Female</td>
<td>0.8074 + 0.0775</td>
</tr>
<tr>
<td>Blood Pressure</td>
<td></td>
</tr>
<tr>
<td>Systolic(mm/Hg) - Mean+SD</td>
<td>125.44 +18.89</td>
</tr>
<tr>
<td>Diastolic(mm/Hg) - Mean + SD</td>
<td>80.36 +13.12</td>
</tr>
<tr>
<td>FBG levels (mg/dl) –Mean+ SD</td>
<td>108.32 +27.64</td>
</tr>
</tbody>
</table>

**Table 2:** Prevalence of associated risk factors by gender, prehypertension hypertension, diabetes and prediabetes

<table>
<thead>
<tr>
<th>Variables</th>
<th>Male</th>
<th>Female</th>
<th>Pre-hypertension</th>
<th>Hypertension</th>
<th>Diabetes</th>
<th>Prediabetes</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N (%)</td>
<td>N (%)</td>
<td>N (%)</td>
<td>N (%)</td>
<td>N (%)</td>
<td>N (%)</td>
</tr>
<tr>
<td>Body mass index:Kg/m²</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal(18-22.9)</td>
<td>250 (50.2)</td>
<td>195 (43.8)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Overweight(23-25)</td>
<td>22 (11.3%)</td>
<td>11 (18.0%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Obese (&gt; 25)</td>
<td>56 (28.7%)</td>
<td>32 (53.3%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Under weight (below 18</td>
<td>97 (49.7%)</td>
<td>109 (43.6%)</td>
<td>64 (41.8%)</td>
<td>49 (49.0%)</td>
<td>5 (33.3%)</td>
<td>10(4.5%)</td>
</tr>
<tr>
<td>Waist to hip ratio: cm</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Low risk *</td>
<td>16 (8.2%)</td>
<td>26 (13.3%)</td>
<td>8 (34.8%)</td>
<td>7 (30.4%)</td>
<td>3 (20%)</td>
<td>1(4.3%)</td>
</tr>
<tr>
<td>Moderate risk **</td>
<td>20 (10.3%)</td>
<td>35 (14.0%)</td>
<td>16 (62.6%)</td>
<td>11 (18.0%)</td>
<td>5 (33.3%)</td>
<td>1(4.3%)</td>
</tr>
<tr>
<td>High risk ***</td>
<td>133(68.2%)</td>
<td>169(76.1%)</td>
<td>60 (31.7%)</td>
<td>54 (28.6%)</td>
<td>5 (33.3%)</td>
<td>5 (21.7%)</td>
</tr>
<tr>
<td>Normal****</td>
<td></td>
<td></td>
<td>69 (40.1%)</td>
<td>28 (16.3%)</td>
<td></td>
<td>13 (56.5%)</td>
</tr>
<tr>
<td>Abdominal Overweight: (&gt;80cm) &amp; (&gt;90 cm)</td>
<td>25 (12.8%)</td>
<td>75 (30%)</td>
<td>32 (32%)</td>
<td>33 (33%)</td>
<td>6 (40.0%)</td>
<td>6 (26.1%)</td>
</tr>
</tbody>
</table>

*0.80 & 0.95 or below, ** 0.81 to 0.85 &*** 0.96 to 1.0+
participants had high blood pressure whereas 26.1% of them were at risk zone for prediabetes and 40.0% of them had positive blood glucose test.

Linear regression analysis was done to find out the correlation between general adiposity i.e. BMI and SBP and BMI and DBP, a positive relationship ($r=0.226$) was observed between SBP and Body mass index and based on the t-value (4.872) and p-value (0.001). Similarly it was found positively correlated between DBP and BMI ($r=0.183$) based on t value (3.923) and p-value (0.001) as shown in Figure 1.

Binary Linear regression analysis was observed between abdominal circumference and SBP and DBP. There was found a strong positive correlation for SBP ($r = .215, p=0.001$) and DBP ($r=.163, p = 0.001$) Figure 2.

Although correlation between increasing body mass index($r=0.09$, $p=0.05$), Waist circumference($r=0.08$ $p=0.07$) and the prevalence of type 2 DM disease was observed positive but it was not found as strongly correlated as with the incidence of hypertension.

Logistic regression analysis was done to correlate the relationship between waist to hip ratio and with the incidence of type 2 DM among Tribal population. It was found to be highly correlated ($r=0.109$) based on t-value (2.305) $p= 0.022$ as shown in Figure 3. Whereas the value of regression was not giving significant correlation with Systolic i.e. ($r= .084, t=1.778, p=.076$) and Diastolic blood pressure ($r= -.034, t= -.719, p=.473$)

**Discussion**

The association of BMI, WC, and W/H ratio with incident type 2 DM and hypertension were confirmed in our study by significant logistic regression analysis. When comparing the associations with these indicators - BMI and WC and W/H ratio, the body mass index (57.3%) was found to be strongly associated with hypertension (including both overweight and obese category) and waist to hip ratio ($r=.109, p=0.022$) was correlated strongly with positive blood glucose test. Waist circumference was also found to be correlated strongly with hypertension (chi-square= 8.559, $p=.014$, df =2) as compared to diabetes.

Ford et al. support the use of WC as a measure of obesity to predict health risk. Among their arguments are that WC has been shown to be a good or better predictor than BMI of the metabolic syndrome, diabetes, cardiovascular disease, and all-cause mortality. It provides information about health risk in addition to body mass index; and it is conceptually easy to measure, although it does require
some training and standardization. However, others have noted that substitution of BMI by WC as an indicator of risk for cardiovascular disease and diabetes may be an oversimplification. Some counter arguments nonetheless prove that waist circumference is strongly correlated to body mass index ($r = 0.08$) (19, 23, 24). Our study also supported that BMI and WC ($r = 0.05$) (Fig 4) were strongly correlated with each other and both predicts the same results for hypertension. Other indicators have been suggested to describe fat distribution associated with abdominal obesity i.e. W/H ratio. In our analysis, we had also included W/H ratio as it was the most common obesity-related predictor of diabetes and it has a weaker correlation with body mass index than waist circumference. However, some have argued against the use of W/H ratio as a measure of obesity because of its ambiguous biologic interpretation, its lesser sensitivity to weight gain, its greater variability across age, sex, and ethnic groups, and its greater computational complexity and interpretation in a public health context.

**Conclusion**

The present study demonstrated consistently strong associations between body mass index versus waist circumference and waist circumference versus waist to hip ratio. It also showed positive association with type 2 diabetes and hypertension. Finally we can conclude our study by saying that all the indicators were strongly associated with each other and Body mass index and waist circumference had strong association with hypertension than diabetes. Waist to hip ratio had also positive association with hypertension but had strongly correlated with diabetes.

**References**

Comparative assessment of efficacy of a single application of Potassium Oxalate and N-Butyl 2-Cyanoacrylate in treatment of hypersensitive teeth-A clinical study

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¹Prof., Department of Periodontics, S.J.M. Dental College and Hospital, Chitradurga, Karnataka, India, ²Lecturer, Department of Oral Pathology, S.J.M. Dental College and Hospital, Chitradurga, Karnataka, India

Abstract
Pain has been so closely associated with dentistry that the word ‘pain’ and ‘dentistry’ have almost become inseparable, and one of the most important objective of dentistry has been the control/elimination of pain.

Dentin hypersensitivity is a common painful condition of the teeth, which has caught the attention of people since our ancient times. In today’s busy world, it is difficult for the suffering patient to keep up the repeated visits to get treated for hypersensitivity. This has led to evolving of treatment modalities utilizing single application medicaments like the resins, oxalates and cyanoacrylates.

Keywords
Hypersensitivity, cyanoacrylates, oxalate.

Introduction
Pain has been so closely associated with dentistry that the word ‘pain’ and ‘dentistry’ have almost become inseparable, and one of the most important objective of dentistry has been the control/elimination of pain.

Dentinal hypersensitivity can be defined as pain arising from exposed dentin, typically in response to chemical, thermal, tactile or osmotic stimuli that cannot be explained as arising from any other form of dental defect or pathology⁶.

In the earlier days dentinal hypersensitivity was treated randomly using traditional cures like materials of natural origin (roots, plant extracts, animal parts, and naturally occurring salts). Most of the therapies proposed, rely on one of the two major suppressive mechanisms i.e. sealing off the dentinal tubules or dampening neural impulses.

Almost all of the agents used in the effective treatments of dentin hypersensitivity have major drawbacks such as requiring multiple applications, application by professionals. In today’s busy world, it is difficult for the suffering patient to keep up the repeated visits to get treated for hypersensitivity. This has led to evolving of treatment modalities utilizing single application medicaments like the resins, oxalates and cyanoacrylates.

Various salts of oxalates, like the ferric oxalate, potassium oxalate are used. These salts precipitate to occlude the dentinal tubule orifice and thus reduce sensitivity.

Cyanoacrylates have made its way into dentistry as a tissue adhesive, based on which property the use of cyanoacrylates to block the dentinal tubule orifice is studied.

The present short term clinical trial aims at an in vivo assessment of 30% dipotassium oxalate and n-butyl 2 cyanoacrylates as desensitizing agents.

Aims & Objectives
To compare the clinical efficacy of a single application of 30% dipotassium oxalate and n-butyl 2-cyanoacrylate in treating hypersensitive teeth.

Materials & Method
This short term clinical study was carried out on 45 teeth from 15 patients from the out patient section of Department of Periodontics, for a comparative evaluation of the efficacy of single application of 30% dipotassium oxalate and n-butyl 2-cyanoacrylate in the management of hypersensitive teeth.

The criteria for selection of patient were:

Inclusion criteria:
• Patients with a history of dentinal hypersensitivity due to exposure of root surface by periodontal disease.
• The labial surfaces of the anteriors were selected.

Exclusion criteria:
• Presence of caries.
• Recent treatment of any type for hypersensitivity by the patients or professionals.
• Hypersensitivity due to abrasion, erosion and attrition.
• Patients with any systemic disease, cardiac pacemakers.

Pretreatment assessments: The presence of hypersensitivity was assessed prior and during treatment in two ways:

1. Subjective assessment.
   a. Visual analogue scale (VAS)
2. Objective assessment
   a. Tactile stimulation
   b. Electrical stimulation

The subjective and objective assessments were recorded prior to the treatment; immediately after treatment and at weekly intervals thereafter for 12 weeks.
Treatment modalities
A total of 15 patients with hypersensitive teeth due to denudation of root because of periodontal disease, but not due to wasting disease (attrition, abrasion, erosion) were selected. Three anterior teeth were selected per patient. The selected teeth were divided into 3 groups:

**Group I** - Application of distilled water for 1 min (control group) with a cotton pallet.

**Group II** - Application of 30% dipotassium oxalate for 1 min with a cotton pallet.

**Group III** - Application of n-butyl 2-cyanoacrylate for 1 min with a cotton pallet.

Freshly prepared 30% dipotassium oxalate solution was used for every application. About 30 grams of dipotassium oxalate crystals were dissolved in 100 ml of distilled water at room temperature, to get 30% dipotassium oxalate solution.

Scaling was done and patients recalled after one week. At that time patients who continued having subjective and objective symptoms of hypersensitivity were treated. As previously mentioned each patient was divided into 3 groups and treated with respective solutions. After application for 1 min the patients were asked to rinse with water, following which subjective and objective assessments were redone and recorded immediately. Assessment was done at weekly intervals thereafter for 12 weeks.

**Observations & Results**
A total of 45 teeth were treated (in vivo), which were divided into 3 groups of 15 each, assigned to as group I, II and III. Group I specimens (control group) were treated with distilled water; Group II were treated with 30% dipotassium oxalate and Group III were treated with N-butyl 2-cyanoacrylate.

Severity of hypersensitivity was assessed before application, immediately after application, later on at weekly intervals for 12 weeks.

Table 1:

<table>
<thead>
<tr>
<th>Time</th>
<th>Control (distilled water) Mean± SD</th>
<th>30% Dipotassium oxalate Mean± SD</th>
<th>N-butyl 2-cyanoacrylate Mean± SD</th>
<th>F* Value</th>
<th>Pair wise comparison</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>I - II</td>
</tr>
<tr>
<td>BA</td>
<td>7.3 ± 1.1</td>
<td>7.7 ± 0.9</td>
<td>8.0 ± 0.8</td>
<td>1.85</td>
<td>NS</td>
</tr>
<tr>
<td>IA</td>
<td>7.2 ± 1.1</td>
<td>3.1 ± 0.5</td>
<td>2.7 ± 0.8</td>
<td>128.9</td>
<td>&lt; .01</td>
</tr>
<tr>
<td>VI Week</td>
<td>7.2 ± 1.1</td>
<td>3.1 ± 0.5</td>
<td>3.1 ± 0.7</td>
<td>12.1</td>
<td>&lt; .01</td>
</tr>
<tr>
<td>X I I Week</td>
<td>6.4 ± 0.6</td>
<td>3.1 ± 0.5</td>
<td>3.3 ± 1.0</td>
<td>93.2</td>
<td>&lt; .01</td>
</tr>
</tbody>
</table>

*ANOVA F – Test [If F > 8.2, P < 0.001] = Highly Significant
Newman-Keul’s range test [If Difference > minimum mean value = Significant range, P < 0.01 = Significant]
BA - Before Application, IA - Immediately after application
weekly intervals for 12 weeks. The statistical analysis of the data collected was done using one factor ANOVA, Newman Keul’s range test.

One factor ANOVA was used for comparing different groups. For pair wise group comparison, Newman Keul’s range test was used. The values are expressed as Mean ± S.D. A ‘P’ value for less than .05 was considered significant.

Visual analogue scale (Table - 1):
It was observed that the mean value during the initial period i.e. before treatment showed a definite evidence of hypersensitivity, mean: Group I - 7.3; Group II - 7.7; Group III - 8.0.

After treatment there was a gradual decline in the readings right after the immediate application to 12th week, except for the control group (i.e. group treated with distilled water).

Mean value of visual analogue scale with regards to Group I showed a reduction of sensitivity at 7th week, (Mean - 6.6). Group II showed a reduction (Mean - 3.1) immediately after application and was the same for 12 weeks, Group III also showed reduction (Mean - 2.7) immediately after application, but the mean score increased at 6th week (Mean - 3.1).

Analysis variance (ANOVA) test revealed that the mean values for visual analogue scale method at different periods decreased significantly (F - 93.2).

Pair wise (Newman Keul’s Range test) showed that comparison between Group I & II, I & III were statistically significant throughout the 12 weeks (P<.01), whereas group II & III were not statistically significant.

Tactile stimulus (Table - 2):
It was also observed that the mean value during the initial period I before treatment, mean: Group 1-2.7; Group II - 2.7; Group III -.7 showed a definite evidence of hypersensitivity.

In Group I the reduction of mean score was noted after seventh week (Mean - 1.7), in group II there was an immediate reduction in mean score after immediate application (Mean - 0) and was the same for the following12 weeks. Group III showed reduction (Mean - 0) immediately after application, but the mean score increased after 7th week (Mean - 0.9). Analysis variance (ANOVA) test revealed that the mean value for tactile stimulus score method at different periods decreased significantly (F - 79.4).

Pair wise (Newman – Keul’s Range test) showed that comparison of group I & II, I & III were statistically significant throughout the 12 weeks (P<.001). Group II & III were statistically significant after 7th week (P<.01).

Electric stimulus (Table - 3):
The mean value during the initial period i.e. before treatment also showed a definite evidence of hypersensitivity, mean: Group I -13.5; Group II - 13.5; Group III - 14.6.

In Group I the increase in mean score was noted after 7th week (Mean - 14.3), in Group II there was an increase in the mean score immediately after application (Mean - 43.3) and was the same for 12 weeks. In Group III, there was a similar increase in the mean score immediately after application (Mean - 54.5), which gradually reduced by 6th week (Mean - 48.9).

Analysis variance (ANOVA) test revealed that the mean value for electric stimulus score method at different periods decreased significantly (F -203).

Pair wise (Newman Keul’s range test) showed that comparison between Group I & II, I & III was statistically significant through out the 12 weeks (P<.001) but Group II & III were statistically significant upto 7th week (P<0.01).

Discussion
Many therapeutic means and agents have been tried to treat dentin hypersensitivity. Patient’s testimony concerning every day experiences provides a useful guide to the state of the problem and the need for treatment.

To prevent and to treat hypersensitivity more effectively, attention must be given to etiology, mechanism and methods for the assessment of hypersensitivity.

In the present study, only the hypersensitive teeth with recession due to periodontal diseases but not due to abrasion or attrition were selected. In patient with vigorous home care procedures by brushing (abrasion), electron micro radiographic findings demonstrated that dentinal tubules on and near the surface had been obstructed by mineral deposition and these areas were never sensitive (William H. Hiatt 1972)8. The patients after scaling were recalled one week later for the continuation of the study. David H. Pasley (1990)3 stated that the smear layer created during manipulation of root surface may last for 5 to 7 days.

Table 2:

<table>
<thead>
<tr>
<th>Time</th>
<th>Control (distilled water) Mean± SD</th>
<th>30% Dipotassium oxalate Mean±SD</th>
<th>N-butyl 2-cyanoacrylate Mean±SD</th>
<th>F* Value</th>
<th>Pair wise comparison</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>I - II</td>
</tr>
<tr>
<td>BA</td>
<td>2.7 ± 0.5</td>
<td>2.7 ± 0.5</td>
<td>2.7 ± 0.5</td>
<td>0.0</td>
<td>NS</td>
</tr>
<tr>
<td>IA</td>
<td>2.7 ± 0.5</td>
<td>2.7 ± 0.5</td>
<td>0</td>
<td>44.80</td>
<td>&lt; .01 NS</td>
</tr>
<tr>
<td>VI Week</td>
<td>2.6 ± 0.5</td>
<td>2.6 ± 0.5</td>
<td>0</td>
<td>44.80</td>
<td>&lt; .01 NS</td>
</tr>
<tr>
<td>X 11 Week</td>
<td>1.6 ± 0.5</td>
<td>1.6 ± 0.5</td>
<td>1.1 ± 0.3</td>
<td>79.9</td>
<td>&lt; .01 NS</td>
</tr>
</tbody>
</table>

Newman-Keul’s range test (If Difference > minimum mean value = Significant range, P < 0.05, P < 0.01 = Significant)
This study focuses on clinical efficacy of 30% dipotassium oxalate and n-butyl 2-cyanoacrylate by single application. The present study showed that the mean values of visual analogue scale with 30% dipotassium oxalate decreased significantly immediately after application (Table - 1). The mean values of tactile stimulus & electric stimulus with 30% dipotassium oxalate decreased immediately after application and were the same for the 12 weeks (Table - 2, 3).

It can be concluded that there was an immediate decrease in hypersensitivity after 30% dipotassium oxalate single application which was in agreement of the results presented by M.S Seo. et al., (1991)7, whereas in the study by Kathleen Brough Muzzin (1989)5 no significant difference was noted for teeth treated with distilled water followed by 30% dipotassium oxalate. This could be attributed to the fact that, they used distilled water prior to the application of 30% dipotassium oxalate.

The present study revealed the decrease in the mean values of visual analogue scale significantly immediately after application of n-butyl 2-cyanoacrylate and gradual increase in the mean values was noted at VII weeks (Table - 1). The mean value of tactile stimulus with n-butyl 2-cyanoacrylate decreased immediately after application, but a gradual increase in the mean values was noted at around VII week (Table - 2).

The current which is required to elicit hypersensitivity, increases as tooth sensitivity decreases, the less sensitive the tooth, the greater the voltage required to deliver the current flow. The mean value of electric stimulus with n-butyl 2-cyanoacrylate increased immediately after application, but a gradual decrease in the mean value was noted at around V week in our study (Table - 3).

Finally, the present study with the use of n-butyl 2-cyanoacrylate concluded that there was immediate reduction in hypersensitivity after single application but there was a gradual increase in hypersensitivity noted around 6th week which is in accordance with the study of Bahram Javid et al. (1987)2.

### Summary & Conclusion

Comparison of 30% dipotassium oxalate and n-butyl 2-cyanoacrylate exhibited no significant difference with VAS. There was a statistical significant difference in tactile stimulus after 7 weeks for both experimental groups, (as compared to control) wherein the 30% dipotassium oxalate group showed consistent absence of hypersensitivity till the end of the study. With electric stimulus, the difference was statistically significant up to 6 week for both experimental groups, (as compared to control) wherein the n-butyl 2-cyanoacrylate group showed more reduction in hypersensitivity than 30% dipotassium oxalate.

With respect to cost effectiveness and long term efficacy 30% dipotassium oxalate could have an edge over n-butyl 2-cyanoacrylate as a desensitizing agent.

In conclusion of the study, it was seen that a single application of 30% dipotassium oxalate and n-butyl 2-cyanoacrylate were almost equally effective as

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**Table 3:**

<table>
<thead>
<tr>
<th>Time</th>
<th>Control (distilled water) Mean±  SD</th>
<th>30% Dipotassium oxalate Mean±SD</th>
<th>N-butyl 2-cyanoacrylate Mean±SD</th>
<th>F* Value</th>
<th>Pair wise comparison</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>BA</td>
<td>1.35 ± 1.8</td>
<td>1.35 ± 2.4</td>
<td>1.81</td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td>IA</td>
<td>1.37 ± 2.1</td>
<td>4.33 ± 4.8</td>
<td>57.22</td>
<td>&lt; .01</td>
</tr>
<tr>
<td></td>
<td>VI Week</td>
<td>1.37 ± 1.8</td>
<td>4.34 ± 4.8</td>
<td>&lt; .01</td>
<td>&lt; .01</td>
</tr>
<tr>
<td></td>
<td>X I Week</td>
<td>1.51 ± 3.5</td>
<td>4.37 ± 4.4</td>
<td>&lt; .01</td>
<td>&lt; .01</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>&lt; .01</td>
<td>N S</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>&lt; .01</td>
<td>&lt; .01</td>
</tr>
</tbody>
</table>

Newman-Keul’s range test (If Difference > minimum mean value = Significant range, P < 0.01 = Significant)
desensitizing agents upto 6-7 weeks after application - Beyond this, the 30% dipotassium oxalate group showed a consistent absence of hypersensitivity till the end of the study.

References
Role of human bite mark identification and DNA technology in forensic odontology: A review

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Abstract

Out of the numerous fields of study and analysis in forensic odontology an important principle is the recognition and interpretation of marks and injuries produced for human bites in foods founded in crime scenes or in the human skin. Thus, the knowledge of the anatomical detail and particularities of human dentition and distribution in the dental arcs can be of real value to forensic odontologist. These bite marks are usually found over the skin, as a result of rapes, fights, assaults, abuses and child violence, this evidence can mean the crime resolution, assuming a decisive role in the criminal identification. In this direction, the DNA technology can be used in the recovery of the genetic material, through the saliva deposited in the skin, after the production of a bite mark, due to the fact that dental impression can be adulterated easily and showing problems on expert analysis. This review emphasizes the importance of the recognition and identification of these injuries combining them with the use of the DNA technology for individual identification.

Key Words
Bite marks, DNA, saliva.

Introduction

The forensic dentist field of activity is not restricted just to the examination of dental remains but extending to several areas such as anthropology, genetics, biochemistry, forensic traumatology, radiology, computing and images mixing, and is regulated by a pertinent federal law.

Today, oral conditions analysis has been presented as a crucial tool to the forensic investigation team members when solving identification problems and it has been helpful to Criminal and Civil Justice1.

One of the possibilities in Forensic Dentistry field is related to the human identification process, such as in cases which involve the study of bite marks. Bite can be defined as the mark made by human or animal teeth in the skin of living people, cadavers or unanimated objects with relatively softened consistence³.

Besides the agent identification, bite mark analysis, in a forensic investigation, can elucidate the kind of violence and the elapsed time between its production and the examination. It can show if the bite was produced ante mortem or post-mortem and, in case of several bite marks, identify the sequence of them².

Therefore, by observing, analyzing and interpreting, the bite marks constitute a important medical-judiciary proof in some cases of offense and help in suspects' exclusion or point out the culpability elements⁵. However, bite marks do not embody all the requisites of an ideal identification method, but it can represent, in some cases, the unique signs of real value to criminal investigation.

Nowadays, there are three kinds of human identification using oral characters. Two of them have been used since long time. The first one is denominated comparative dental identification and involves a comparison between ante mortem and post-mortem records. The second one is composed by the reconstruction of post-mortem dental profile and is used in the cases in which there is no suspicion of the person or his/her descendents. The third kind works on the application of the modern techniques of DNA, in order to establish the identity⁴.

The present work intends to display some applications of DNA technology in human identification by studying the bite marks in Forensic Dentistry.

Production of Bite marks

Bite marks are produced as a result of teeth pressure on skin, described as force combination. In general, when a person bites an object, the superior teeth hold the object while the inferior ones cut it. The mark left by the superior teeth, however, is extremely relevant in order to provide information such as: dental alignment, size and shape of dental archs5. Human adult dentition consists of 32 teeth and each one of them has its own size, shape and features. The action of the dental arch on the skin may produce many kinds of lesions, as the dental elements act as incisive instruments or even incisive-cut⁶.

The greatest challenge is Forensic Dentistry are bite marks found in human skin, because of the distortion presented and the time elapsed between the production and the analysis⁷. Moreover, many factors can affect the structure of the lesions produced by bite marks, which include: applied force, bite duration and movement between tissues and teeth.

Use of DNA in bite mark identification

Techniques involving DNA in Forensic Dentistry offers a new tool when traditional identification methods fail due to the effects of heat, traumatism or autolytic processes⁹,
as well as in distortions and difficulties in analysis. DNA analysis in forensic samples has been increasingly used in human identification processes.

Due to this abundance of material, the use of the technique based on PCR (Polymerase Chain Reaction) has acquired great importance in DNA post-mortem analysis in forensic cases. Polymerase Chain Reaction is an enzymatic amplification of a specific DNA sequence, aiming millions of copies production from this sequence in a test tube, which was first described by Kary Mullis, in the late 1980's, and enabling a new strategy of gene analysis though a simple and fast method, excusing all the laborious stages of genic cloning.

When treating of forensic samples, the DNA study is usually done through the analysis of regions of short tandem repeats (STR), which can be defined as DNA hypervariable regions that present tandem repeats of fragment that have from two until nine pairs of bases (pb). The most valuable STR's to human identification present a higher polymorphism (larger alleles amount), smaller size, greater heterozygosity frequency (higher than 90%) and low mutation frequency.

In order to perform human identification, it is more interesting to use the molecular markers that have great variability within the population. In other words, high level of polymorphism, enabling that the probability of two people that present the same alleles gets smaller. And, when we wish to identify a subject that comes from a certain population, the study of different markers in that population is necessary, in order to know what the present alleles are and how often they appear, with the purpose of defining the best markers to be used.

Besides the genomic DNA, inside the cell nucleus, it is possible to use mitochondrial DNA. This organela has a number that ranges from 100 to 10,000 copies per cell, enabling the material analysis with limited amounts and also DNA samples partially degraded.

Saliva is a very useful DNA source due to the fact of being collected by painless and non-evasive way, able to be used even when it is stored in the most different conditions. The amount of saliva deposited on the skin is generally very little in bite mark cases, making it necessary to use methods for collecting, whose result in the recovery is the maximum possible amount of saliva and minimizes any contamination through the victim's skin cells. When checking the DNA analysis reproducibility of collected saliva on the skin, simulating cases that involve bite marks in 20 samples, the double swab technique showed to be sensitive and efficient in criminal cases when there is presence of saliva in bite marks. Saliva, in contact with intact skin, maintains itself in stable conditions and can be recovered, at least, 60 hours after its deposit.

However, it is not always possible to recover DNA from a bite mark, due to the fact that it will be subject to a series of modifications, such as contamination, degradation and putrefaction, depending on the circumstances the body and/or object were submitted.

Bite Mark identification: DNA analysis and genotypical composition of oral bacteria.

The human oral cavity has a large and varied bacterial community, many of which are unique for this habitat. There is wide evidence that oral bacteria are transferred during the human bite act and, in some cases, survive and multiply, creating infections. Besides, there are evidences that individuals shelter unique bacterial species stocks in the oral cavity and that those stocks can be identified by techniques such bacterial typing and protein profiles.

It is important to note that the oral Streptococcus recovery from the skin or objects seems to imply the contact with oral surfaces or deposit saliva which can lead to evidences of buccal-envolvement in the injury.

In a study where 10il fresh saliva sample was collected without stimulation and applied to areas of the upper left quadrant of the thorax, so that the loss rate of units that make up the colony and its recovery ranged from 45 to 50% per hour. They have also noticed that 6, 25 hours after the saliva deposition, oral viable streptococcus could be recovered. In another research, volunteers bit their own arms firmly and the bite marks were sampled in time intervals to recover isolated viable streptococcus, in order to make a genotypical comparison with bacteria from the oral cavity. It was concluded that it is possible to recover bacteria up to 24 hours after the production of the bite, but identification assertiveness is only possible when compared to samples acquired from the subjects’ teeth responsible for the bite. Isolated Streptococcus from recent bite marks can be listed by PCR and compared to the teeth that were responsible for the bite. Moreover, they claim it is preferable to recover the subject's DNA, but such strategy is not always possible, the recovery of the bacteria derived from the subject’s tooth may enable the link with the suspect of a crime.

In conclusions, the knowledge proceeding from Forensic Dentistry and Molecular Biology has great importance to the expert practice when we think of a dentist inserted in forensic investigation team in a bite mark case. It's necessary to broaden the pertinent studies of the theme, in order to establish protocols to allow additional tools in criminal investigation.

It is stated that, in judicial proceedings involving Dentistry, being Civil or Criminal, it is extremely necessary the presence of a professional that militates in Forensic Dentistry as a judicial expert.

References
Congenital malformations detected at birth – A prospective study in Bangalore

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Abstract
In a prospective study of 4280 births in Vanivilas Hospital, Bangalore Medical College, Bangalore 61 cases of congenital malformations were observed with an Incidence of 14.25/1000 live births. A total of 117 malformations were observed. Among these, 95 were major and 22 were of minor type. The central nervous system, gastro-intestinal and multiple system involvement were the commonest with 12 cases each (19.67%), followed by musculoskeletal 11 (18.03%), cardio-vascular 4 (6.56%), genito urinary 02 (3.30%) cases, cutaneous 1 (1.64%) and 07 (11.49%) cases were miscellaneous. Malformations in the new born males were 52.46%, in females it was 45.90% and 1.64% are of ambiguous genitalia thereby showing male preponderance. Maximum of 42 cases (68.85%) were born to non-consanguineous group and 19 cases (31.15%) were to consanguineous couples.

Keywords
Congenital malformations, Neonate, Consanguinity.

Introduction
A Congenital malformation is a functional or structural anomaly which result from chromosomal aberrance, deficiency or damage to the organism in the process of growth and development occurring prior to birth. These congenital malformations occur all over the world and constitute an impenetrable core of perinatal mortality in developed countries. The awareness of incidence, prevalence and patterns of malformations can help medical and paramedical personnel to identify ‘at-risk’ cases early and plan appropriate and effective intervention in treatment and prevention if possible. Hence the present study was undertaken with a view to assess the frequency of congenital malformations in newborns and to analyze the magnitude of the problem.

Material and Methods
The study was conducted at Vanivilas Hospital, attached to Bangalore Medical College, Bangalore for a period of 1 year. All the live newborns delivered at Vanivilas Hospital were screened for congenital anomalies within 24–48 hours after birth with a written consent from parents/relatives for examination.

The details of each of the congenital anomalous newborns regarding sex, major and minor malformations and parental consanguinity were recorded. The information was sought from the mother or father regarding consanguinity and previous births.

Major malformations were included, the anomalies which caused serious structural, cosmetic and functional disability requiring surgical or medical management. The minor anomalies included those anomalies which were unlikely to cause serious hindrance to life like preauricular tag, polydactyly etc.

Results
Out of total 4280 births, 61 cases of congenital malformations were observed. In these 61 cases 117 malformations were noted in which, thirty two (52.46%) were male and twenty eight (45.90%) were females with one case of ambiguous genitalia. The distribution of congenital malformation is high among male children with a male female ratio of 1.4:1.

Discussion
The incidence of congenital malformations in our study is 14.25 per 1000 live births carried out at Vanivilas Hospital, Bangalore which includes both major and minor anomalies. Several studies reported to have higher incidence of 34/1000 by Kulshresta et al. WHO has reported 18.87/1000, 22.54/1000 at Melbourne and Johannesburg respectively. These reported incidence rates may not reflect the true as McIntosh et al reported that only 43% of anomalies will be detected at birth and

Table 1: System Wise Distribution of Abnormalities

<table>
<thead>
<tr>
<th>SL NO.</th>
<th>Abnormalities</th>
<th>Total number</th>
<th>%</th>
<th>Incidence /100 live births</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>CNS</td>
<td>12</td>
<td>19.67</td>
<td>2.80</td>
</tr>
<tr>
<td>2</td>
<td>GIT</td>
<td>12</td>
<td>19.67</td>
<td>2.80</td>
</tr>
<tr>
<td>3</td>
<td>Multiple system</td>
<td>12</td>
<td>19.67</td>
<td>2.80</td>
</tr>
<tr>
<td>4</td>
<td>Musculoskeletal Sys.</td>
<td>11</td>
<td>18.03</td>
<td>2.57</td>
</tr>
<tr>
<td>5</td>
<td>CVS</td>
<td>4</td>
<td>6.56</td>
<td>0.93</td>
</tr>
<tr>
<td>6</td>
<td>Genito urinary Sys.</td>
<td>2</td>
<td>3.30</td>
<td>0.47</td>
</tr>
<tr>
<td>7</td>
<td>Skin</td>
<td>1</td>
<td>1.64</td>
<td>0.23</td>
</tr>
<tr>
<td>8</td>
<td>Miscellaneous</td>
<td>7</td>
<td>11.40</td>
<td>1.64</td>
</tr>
</tbody>
</table>
remaining by first year of life.

Out of 61 cases of total congenital malformations, 55 infants 1.29% (12.85/1000) have major type of malformation either single or multiple, 05 infants 0.14% (1.40/1000 live births) have minor type. Among the 95 major malformations, central nervous system anomalies (19/95) were dominating in the present study, followed by multiple system involvements having 12 cases each (19.67%), musculoskeletal systems (18,03%), Cardiovascular (6.56%) and Gastrointestinal system (3.30%).

Goravalingappa8 have reported highest incidence of Central Nervous System, We observed 12/61 cases (19.67%) of Central Nervous System malformations, with an incidence rate 2.80/1000 LB which consisted of 8 cases (1.87/1000) of meningocoele, myelomeningocoele and one associated with spina bifida, 2 cases (0.47/1000 LB) of hydrocephalus, one case (0.23/1000 LB) of microcephaly and a case of (0.23/1000) Choroid plexus cyst. Among these, Neural tube defects are commonest. Isolated myelomeningocoele accounted 0.69/1000LB for highest incidence among central nervous system anomalies in our series, which is consistently high when compared to Choudhury A et al9, Datta V et al10, WHO study1 (0.05/1000). Whereas Mathur BC et. al13 reported 0.9/1000, Swain S et al9 0.76/1000, which is high when compared with our study.

The overall incidence of congenital malformations in relation to sex of the child is more in males than in females. Total 32 males (52.46%), 28 female (45.90%) presented with abnormality and one case (1.64%) with Ambiguous genitalia. Later karyotyping revealed the latter as being the male. The sex ratio in our study for male to female is 1.4 : 1, which is well comparable to the previous observations by Ajay Kalra et al17, Vikram Datta et al16, Goravalingappa J P et al16 and Thirumalaikolundusubramanian P et al18 and Arjun Singh et al1. Whereas, other studies reported high incidence in female child than males in Zilfalil B A et al19, but authors like Swain et al9 have noted no difference in the incidence of congenital malformation in relation to the sex of the baby. Why males are prone for the congenital malformations is not yet proved. Possibly, it requires a genetic background to explain the reason. Hence it requires more extensive study to prove, whether really the male sex or both sexes are equally susceptible to congenital malformations.

A total of 19/61 cases (31.15%) of congenital malformations, in our study belonged to consanguineous couples as compared to 42/61 (68.85%) births of malformations, were to non-consanguineous group. Uncle niece was the most common form of marriage 14/61 (22.95%) and followed by first cousin marriages 05/61 ((8.19%). Radha Rama Devi A et.al20 reported that incidence of uncle niece union in Bangalore as 12.66% by surveying seventeen hospitals. But our study shows an incidence of 31.15% among consanguineous couples who gave birth to congenital malformed infants with a higher rate of uncle–niece union of 22.95%. These reports are consistent with the extensive study of Kulkarni ML21. WHO study in Bombay and Alexandria showed the frequency of congenital malformations was significantly higher in offspring's of first cousins and closer relationships than in those related less closely than first cousin1. Thus it seems that closer the family relationships of the parents, the greater the chances of congenital abnormalities.

The study frequency of malformations affecting major systems is high, particularly affecting multiple system(7/12). Cardiovascular system involvement in parental consanguineous group shows 50% involvement (02/04 cases), whereas 1/3 of CNS and 1 / 4 of GIT cases are involved in parental consanguinity. Whereas minor
malformations and skin anomalies shows no association with parental consanguinity. Consanguinity contributes to intra uterine growth retardation. There are recessive genes with a slightly retarding effect on fetal growth present in general population\textsuperscript{22}.

**Summary and Conclusion**

In present study out of 4280 new born delivered 61 cases of congenital malformations were studied with an Incidence of 14.25/1000 live births. In all the 61 cases, a total of 117 malformations were observed. Among these 95 were major & 22 were of minor type. Malformations in the new born males were 52.46%, in females it was 45.90% and 1.64% are of ambiguous genitalia thereby showing male preponderance. The central nervous system, gastro-intestinal and multiple system involvement were the commonest with 12 cases each (19.67%), followed by musculoskeletal 11 (18.03%), cardio-vascular 4 (6.56%), genito urinary 02 (3.30%) cases, cutaneous 1 (1.64%) and 07 (11.49%) cases were miscellaneous. The commonest anomaly was neural tube defects (NTD) particularly myelomeningocoele in central nervous system, ano-rectal anomaly in gastro-intestinal system, talipes equinovarus in musculoskeletal and congenital heart disease in cardiovascular and multiple

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### Table 3: Major Malformations (22.20/1000 Live births)

<table>
<thead>
<tr>
<th>SL. NO.</th>
<th>ANOMALIES</th>
<th>TOTAL</th>
<th>Incidence /1000LB</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>CTEV</td>
<td>6</td>
<td>1.40</td>
</tr>
<tr>
<td>2</td>
<td>Rocker bottom foot</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>3</td>
<td>Rhizomelic limb</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>4</td>
<td>Torticollis</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>5</td>
<td>Cong. Diaphragmatic hernia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>6</td>
<td>Gum hypertrophy</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>7</td>
<td>Natal tooth buds</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>8</td>
<td>Tibial bone deformity</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>9</td>
<td>Omphalocele</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>10</td>
<td>Multiple limb anomalies</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>11</td>
<td>Anorectal anomaly</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>12</td>
<td>Imperforate anus with fistula</td>
<td>9</td>
<td>2.10</td>
</tr>
<tr>
<td>13</td>
<td>Cleft lip/Cleft palate</td>
<td>4</td>
<td>0.93</td>
</tr>
<tr>
<td>14</td>
<td>Oesophageal atresia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>15</td>
<td>Duodenal atresia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>16</td>
<td>Tracheoesophageal fistula</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>17</td>
<td>CHD-VSD</td>
<td>3</td>
<td>0.70</td>
</tr>
<tr>
<td>18</td>
<td>CHD-PDA</td>
<td>5</td>
<td>1.16</td>
</tr>
<tr>
<td>19</td>
<td>CHD-ASD</td>
<td>4</td>
<td>0.93</td>
</tr>
<tr>
<td>20</td>
<td>CCHD</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>21</td>
<td>Single umbilical artery</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>22</td>
<td>Harlequin syndrome</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>23</td>
<td>Hydrocephalus</td>
<td>5</td>
<td>1.16</td>
</tr>
<tr>
<td>24</td>
<td>Myelomeningocoele</td>
<td>9</td>
<td>2.10</td>
</tr>
<tr>
<td>25</td>
<td>Spina bifida</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>26</td>
<td>Dandy walker cyst malformation</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>27</td>
<td>Microcephaly</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>28</td>
<td>Cong. Choroids plexus cyst</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>29</td>
<td>Cong. Hydronephrosis</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>30</td>
<td>Scrotum like fused phallus</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>31</td>
<td>Ambiguous genitalia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>32</td>
<td>Ectopia vesicae</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>33</td>
<td>Micrognathia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>34</td>
<td>Sacrococcygeal teratoma</td>
<td>3</td>
<td>0.70</td>
</tr>
<tr>
<td>35</td>
<td>Low set ears</td>
<td>6</td>
<td>1.40</td>
</tr>
<tr>
<td>36</td>
<td>Corneal opacity</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>37</td>
<td>Hypertelorism</td>
<td>5</td>
<td>1.16</td>
</tr>
<tr>
<td>38</td>
<td>Clouding of cornea</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>39</td>
<td>Multiplecraniofacial anomaly</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>40</td>
<td>Proptosis</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>41</td>
<td>Fused eye balls</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>42</td>
<td>Inguinal hernia</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td><strong>TOTAL</strong></td>
<td></td>
<td><strong>95</strong></td>
<td></td>
</tr>
</tbody>
</table>

### Table 4: Minor Malformations (5.14/1000 LB)

<table>
<thead>
<tr>
<th>SL. NO.</th>
<th>MALFORMATIONS</th>
<th>TOTAL NUMBER</th>
<th>Incidence /1000 live births</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Polydactyly</td>
<td>4</td>
<td>0.93</td>
</tr>
<tr>
<td>2</td>
<td>Syndactyly</td>
<td>4</td>
<td>0.93</td>
</tr>
<tr>
<td>3</td>
<td>Genu recurvatum</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>4</td>
<td>Preauricular tag</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>5</td>
<td>Prominent occiput</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>6</td>
<td>Short neck</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>7</td>
<td>Depressed nasal bridge</td>
<td>2</td>
<td>0.47</td>
</tr>
<tr>
<td>8</td>
<td>Fish like mouth</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>9</td>
<td>High arched palate</td>
<td>3</td>
<td>0.70</td>
</tr>
<tr>
<td>10</td>
<td>Incurving of fingers</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>11</td>
<td>Long toes</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td>12</td>
<td>Congenital hydrocele</td>
<td>1</td>
<td>0.23</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td><strong>22</strong></td>
<td></td>
</tr>
</tbody>
</table>

### Table 5: Relation of Consanguinity to Congenital Malformations

<table>
<thead>
<tr>
<th>Consanguinity</th>
<th>Cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>Consanguineous</td>
<td>19(31.15%)</td>
</tr>
<tr>
<td>Non-consanguineous</td>
<td>42(68.85%)</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td>61</td>
</tr>
</tbody>
</table>

### Table 6: Incidence of Congenital Malformation at Various Places Compared With Present Study

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>AUTHOR</th>
<th>REGION</th>
<th>INCIDENCE (PER 1000 BIRTHS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ghosh &amp; Bali L\textsuperscript{a}</td>
<td>Delhi</td>
<td>34.0</td>
</tr>
<tr>
<td>2</td>
<td>Stevenson AC et al- WHO\textsuperscript{1}</td>
<td>Bombay</td>
<td>8.6</td>
</tr>
<tr>
<td>3</td>
<td>Stevenson AC et al- WHO\textsuperscript{1}</td>
<td>Calcutta</td>
<td>3.1</td>
</tr>
<tr>
<td>4</td>
<td>Mathur BC\textsuperscript{3}</td>
<td>Hyderabad</td>
<td>31.0</td>
</tr>
<tr>
<td>5</td>
<td>Choudhury et.al\textsuperscript{2}</td>
<td>Calcutta</td>
<td>2.9</td>
</tr>
<tr>
<td>6</td>
<td>Mishra PC et al\textsuperscript{7}</td>
<td>Allahabad</td>
<td>14.64</td>
</tr>
<tr>
<td>7</td>
<td>Goravalingappa\textsuperscript{8}</td>
<td>Hubli</td>
<td>31.27</td>
</tr>
<tr>
<td>8</td>
<td>Swain S et al\textsuperscript{9}</td>
<td>Banaras</td>
<td>12.0</td>
</tr>
<tr>
<td>9</td>
<td>Datta V et.al\textsuperscript{10}</td>
<td>Sevagram (Wardha)</td>
<td>12.4</td>
</tr>
<tr>
<td>10</td>
<td>Presents Study</td>
<td>Bangalore</td>
<td>14.25</td>
</tr>
</tbody>
</table>
systems involvement. Maximum of 42 cases (68.85%) were born to non-consanguineous group and 19 cases (31.15%) were to consanguineous couples.

In conclusion, the incidence of congenital malformations in this study is comparatively low to other parts of India and abroad. Mostly etiological factors remain obscure, but require detailed history taking and thorough investigations for the early diagnosis and treatment. Genetic study should be made mandatory for all the investigations for the early diagnosis and treatment. But require detailed history taking and thorough investigations for the early diagnosis and treatment. Genetic study should be made mandatory for all the investigations for the early diagnosis and treatment.

Table 11: Comparative study showing system wise distribution of cases

<table>
<thead>
<tr>
<th>S. NO.</th>
<th>AUTHOR</th>
<th>CN S</th>
<th>GIT</th>
<th>MS</th>
<th>CVS</th>
<th>RS</th>
<th>GUS</th>
<th>SKIN</th>
<th>MULTIPLE</th>
<th>MISC</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>McIntosh Ret et al7</td>
<td>60</td>
<td>41</td>
<td>154</td>
<td>33</td>
<td>7</td>
<td>39</td>
<td>88</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>2</td>
<td>Ghosh S et al8</td>
<td>38</td>
<td>3</td>
<td>26</td>
<td>64</td>
<td>14.7</td>
<td>6</td>
<td>1</td>
<td>0.2</td>
<td>0.15/1000</td>
</tr>
<tr>
<td>3</td>
<td>WHO BOMBAY Stevenson AC et al11</td>
<td>142</td>
<td>38</td>
<td>1.95/1000</td>
<td>9</td>
<td>0.22/1000</td>
<td>0</td>
<td>6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Stevenson AC et al – Calcutta1</td>
<td>11</td>
<td>1.86/1000</td>
<td>3</td>
<td>0.15/1000</td>
<td>22</td>
<td>1.14/1000</td>
<td>0.10/1000</td>
<td>0.20/1000</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Tibrewala NS et al10</td>
<td>19</td>
<td>1.54/1000</td>
<td>56</td>
<td>1.00/1000</td>
<td>0</td>
<td>41</td>
<td>4.53/1000</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>6</td>
<td>Sugunabai NS et al13</td>
<td>21</td>
<td>2.9/1000</td>
<td>31</td>
<td>1.00/1000</td>
<td>0</td>
<td>4</td>
<td>0.6/1000</td>
<td>2.2/1000</td>
<td>30</td>
</tr>
<tr>
<td>7</td>
<td>Choudhury A et al</td>
<td>13</td>
<td>0.69/1000</td>
<td>15</td>
<td>0.74/1000</td>
<td>9</td>
<td>0.44/1000</td>
<td>0</td>
<td>0</td>
<td>0.09/1000</td>
</tr>
<tr>
<td>8</td>
<td>Datta V et al10</td>
<td>5</td>
<td>1.71/1000</td>
<td>8</td>
<td>2.7/1000</td>
<td>9</td>
<td>1.11/1000</td>
<td>0.69/1000</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>9</td>
<td>PRESENT STUDY</td>
<td>12</td>
<td>2.80/1000</td>
<td>12</td>
<td>2.80/1000</td>
<td>11</td>
<td>2.57/1000</td>
<td>0.93/1000</td>
<td>0</td>
<td>2</td>
</tr>
</tbody>
</table>

(Figures underneath the number indicate incidence)

References

A rare form of aggressive pregnancy gingival enlargement - A case report

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Abstract
Gingival enlargement related to pregnancy is sometimes seen in the oral cavity. Pregnancy is a physiological state that brings full of changes in a woman’s life. The metabolism and immunology of the body are modified by progesterone and estrogen as well as other local factors, these sex hormones may modify the oral mucosa and may lead to various periodontal diseases. A case of female patient 23 yrs of age reported during 8th month of pregnancy with a localised gingival enlargement affecting the buccal aspect of left maxillary central incisor upto canine. The hormonal changes occurring during pregnancy may be associated with generalized or localised gingival enlargement and the presence of local factors may accentuate the gingival response.

Introduction
Pregnancy has far reaching systemic effects extending beyond the reproductive system involving complex physical and psychological changes that have an impact on even the healthy woman. These effects occur mainly due to hormones on almost every organ system¹². Progesterone is the main sex pregnancy hormone. The level of it rises until the 8th month of pregnancy and after that it becomes stable until giving birth. The estrogen level rises slowly until the end of the pregnancy³. The high level of hormones in blood and saliva may cause some periodontal reaction and may increase or cause periodontal disorders. The gingival enlargement has been associated with a variety of local and systemic factors so differential diagnosis becomes an important aspect for complete management of the lesion. Most of the causative factors lead to an unusual hyperplastic tissue response to chronic inflammation associated with local irritants such as plaque, calculus or bacteria and their products⁴. Hormonal changes occurring during pregnancy and puberty, have been known to be associated with varying types of gingival enlargement. Hormonal changes can significantly potentiate the effects of local irritants on gingival connective tissue⁵.

In all forms of enlargements, good oral hygiene is necessary to minimize the effects of systemic factors, Gingivoplasty or Gingivectomy may be required after pregnancy but should be done in combination with prophylaxis and oral hygiene instructions⁶. The pregnancy gingival tumour is a disorder found in .2 to .5% of the pregnant females. It is a benign rapidly growing lesion occurring usually in 1st trimester of pregnancy and extends upto 3rd trimester. Pregnancy tumour is a smooth or lobulated exophytic lesion with a pedunculated or sessile base. This paper presents a case report of a pregnancy associated aggressive gingival enlargement of enormous size.

Case Report
A female patient of 23 years of age reported to the Department of Periodontics, with a localized massive gingival overgrowth on the upper left front region in relation to maxillary left central incisor, lateral incisor and canine with protruding lips (fig:1,2,3 and 4). There was no history of drug intake and hereditary reasons. Patient was eight months pregnant and revealed that her gums used to bleed on brushing since three months of pregnancy, but the enlargement came to the present size at this time, the patient reported that she felt difficulty in chewing, speech and closing of the lips, her esthetics was also compromised because of the enlargement.

Fig. 1: Gingival enlargement i.r.t 21, 22 & 23 during 8TH month of pregnancy

Fig. 2: Gingival enlargement extraoral view
The gingival swelling was so massive that it was evident extraorally. On Intraoral examination the size of the lesion was about 3.5 x 2 cms in dimension and pedunculated. The tumour was bright red in colour, soft that bled on slightest provocation. Subgingival calculus and plaque was present. Patient was unable to maintain oral hygiene in this area, because of gingival enlargement. Oral prophylaxis was performed after routine haematological investigation. Instructions regarding maintenance of oral hygiene were given and advised to return post partum. She reported to the department one month after the delivery of the baby. The dimension of the tumour was same as before delivery. The gingival overgrowth was excised with the help of 15 no. b.p blade and electrocautery, the bleeding was controlled by the help of the ball tip of the electrocautery (fig:5,6). Periodontal dressing was done in the involved region.(fig:9) The excised lesion (fig:7) was sent for histopathological examination, which revealed epithelial proliferation and underlying capillary proliferation along with marked inflammatory cell infiltration seen (fig:8). The healing was uneventful and no relapse was seen on follow up. (fig:10)

**Discussion**

The gingival changes in pregnancy were described as early as 1898 by Pinard. Gingivitis in pregnancy is caused by bacterial plaque, like in non-pregnant individuals. Pregnancy accentuates the gingival response to plaque. The incidence of gingivitis in pregnancy varies from around 50 to 100%. Pregnancy does not alter the healthy gingiva it effects the severity of previously inflamed area. In some cases the inflamed gingiva forms a discrete mass referred to as pregnancy tumour7. Kornman & Loesch (1980) reported that the subgingival flora changes to a more anaerobic flora as pregnancy progresses Prevotella intermedia is the microorganism that increases significantly during pregnancy6. They also stated that the increase is due to elevations of levels of systemic estradiol and progesterone. O’Neil (1979) suggested that the altered tissue response to plaque is due to depression of the maternal T lymphocyte9. Formicola & Associates (1970) have shown that gingiva is a target organ for female sex hormones10.Therefore, the maintenance of oral hygiene before and during pregnancy is very important in order to reduce the incidence and the severity of gingival inflammation. It is generally accepted that increase in gingival inflammation typically begins in the second month and reaches the maximal level during the eighth month of pregnancy. These inflammatory changes may lead to gingiva that appears edematous, hyperplastic and erythematous; the changes may be localized or generalized, and are usually noted on the marginal gingiva and interdental papilla11,12.

**Summary & Conclusion**

The local factors i.e. plaque and calculus are known to be responsible for gingival enlargement during pregnancy. The hormonal factors also play a role in aggravating the hyperplasia. Therefore, the importance of regular dental check up and oral prophylaxis(scaling) cannot be overlooked. In all forms of gingival enlargements, good oral hygiene is necessary to minimize the effects of
systemic factors. In the present case, the massive size of the hyperplastic tissue was excised by surgical excision and the contouring was done by electrocautery thus relieving the patient from difficulty in speech, chewing, closing of lips and esthetics.

References
1. Sangeeta gajendra oral health and pregnancy NYSDJ. 2004, 40-4
A two year study of polycystic ovary syndrome in Davangere, Karnataka


1Associate Professor, 2Assistant Professor, Department of OBG, 3Associate Professor, Department of Surgical Oncology, 4Associate Professor, Department of Forensic Medicine & Toxicology, 5Assistant Professor, Dept. of Pharmacology, 6Associate Professor, Dept. of Microbiology, S.S Institute of Medical Sciences and Research Centre, Davangere- 577005, Karnataka

Abstract

Introduction
Polycystic ovarian syndrome (PCOS) is one of the common gynaecological condition of diverse etiology 30% of the infertile women have anovulation due to polycystic ovarian disease, 20-30% of them have hyperandrogenemia and hyperprolactenemia.

Methods
An explorative type of hospital based study of 102 patient, during a 2 year period was done with 75 (73.52%) married women and 27 (26.47%) unmarried women with varied symptomatology.

Results
67.6% were in the age group 21-30 years. Ovulation induction was done in all the infertile group, an ovulation induction rate of 88.40% was achieved and of these 21 (34.42%) conceived, others on follow up.

Of the unmarried women, all had menstrual irregularities along with obesity (51.85%) and hirsutism (48.14%), after treatment 66.66% had an improvement in their symptoms, other still on follow up.

Conclusion
This study demonstrates the various presentations either single or in combination and their response to the various medical and surgical management.

Introduction
PCOS is an incompletely understood enigmatic disorder of heterogeneous nature and is the commonest endocrinopathy of the reproductive age. PCOS starts appearing from 15-25 years of age and it may take years, for its clinical presentation to appear. It is diagnosed in 5-10% of women between adolescence and menopause. It has an autosomal dominant inheritance with great phenotypic variability presenting with infertility (mean incidence 74%), menstrual irregularity (mean incidence of DUB 29%, and ammenorrhoea 51%) and androgen excess (mean incidence of hirsutism 69% while of virilization being 21%), along with serious long term consequences like endometrial carcinoma, obesity, hyperinsulinsim, cardiovascular risks, bone disease etc., leaving their impact on quality of life, morbidity and mortality.

The diagnosis of PCOS depends on confirming the presence of hyperandrogenism with elevated serum, testosterone and an elevated LH:FSH ratio. Either or both of these may be found in 60-70% of patients, with greatest sensitivity and specificity.

Ultrasound is now considered as the gold standard in the evaluation of infertility. Diagnostic laparoscopy with or without ovarian drilling is used as a second line of management in cases of failed induction or in unexplained infertility and also to assure tubal patency.

The purpose of this study is to know about the clinical course and outcome of polycystic ovarian disease.

Study Aims and Objectives
Aim is to find out the various presentations either single or in combination and their response to the various medical and surgical management.

Objectives
1) To study the clinical presentations and hormonal imbalances in these women with polycystic ovarian disease.
2) Sonological evaluation of women with polycystic ovarian disease and its correlation with hormonal values.
3) Laparoscopic evaluation and/or ovarian drilling of infertile women with polycystic ovarian disease, after failed ovulation induction.
4) To study the outcome of all the modalities of treatment particularly in hirsutism and infertility.

Study Design and Study Population
Type of study
Explorative type of hospital based study.
**Place of study**
All the women attending the gynaecology out patient department at S.S. Institute of Medical Science and Research Centre with complaints of hirsutism, infertility and menstrual irregularities were studied.

**Study Eligibility Criteria**

**Sample size of duration of study**
The criteria was that a total of about 100 patients or the number of patients with polycystic ovarian disease in a period of 2 years from Oct. 2008 to Oct. 2010 were recruited for the purpose of the study. The related data were collected by detailed history and clinical examination.

**Inclusion Criteria**
All patients with polycystic ovarian disease irrespective of age and body weight.

**Exclusion Criteria**
Patients were later excluded if the cause of hirsutism, infertility and obesity was of non-ovarian origin.

**Methodology**

**Study Procedure**
In the out patient clinic, an informed consent was taken, and patients were recruited for the study. Then a detailed history and general examination was done.

A diagnosis of PCOS was made based on the relevant history, clinical findings and confirmed by relevant investigations which included basic investigations like hemoglobin levels, urine routine and random blood sugar estimations, hormonal profile testosterone, follicle stimulating hormone, leutinising hormone, prolactin, thyroid stimulating hormone levels were assessed within 3 days of menstruation and an ultrasound examination for prediction of ovulation was done around the time of ovulation.

The semen analysis of the husband, if the patient presented with infertility was done to rule out an associated male factor as a cause of infertility.

For those who desired to conceive, ovulation induction with clomiphene citrate was the initial choice, with 50 mg/day for 5 days, starting on D2 of cycle. Follicular study was done on 10th, 12th and 14th day of the cycle.

To improve the mid cycle LH surge, Inj HCG (5000 – 10000 u) Im was given around the time at which the follicle size was 18-20 mm and HCG administered patients required intense follow-up for any evidence of ovarian hyperstimulation symptoms.

Patients with hirsutism were more resistant to CC alone and so dexamethasone was added at a low dose of 0.5 mg, orally and those with high prolactin levels were given bromocriptine strating at an usual dose of 2.5 mg/day, orally for 2 months along with clomiphene. A repeat hormonal assay was done after 2 months of treatment to see for the response.

If all the above combinations failed, then preparations of gonadotropins (HMG) was given for follicle stimulation beginning by 7-14 days of the cycle with continuous daily intramuscular injections beginning with 75 u/day or with purified FSH preparations with a dose of 1 amp/day (75 lu FSH) intramuscularly was given and monitored.

Metformin was added to obese or those with insulin resistance as 500 mg tid for 3-4 months. If when the medical line of treatment was unsuccessful to achieve conception, diagnostic hystero laparoscopy with chromopertubation was planned, with or without ovarian drilling under general anaesthesia. Following laparoscopy, clomiphene was restarted. Conception rates improved, conceived women were followed up to know the outcome of pregnancy.

In the other group of unmarried women with polycystic ovarian disease, apart from a general advise of weight reduction, they were started on either oral contraceptives with or without Finasteride (Once a day) and Aldactone (Once a day) for a period of 3 to 6 months and then evaluated for the improvement of symptoms being regularization of cycles, decreasing hirsutism scores and reduction in body weight. A repeat hormonal assay was done 3 months later for the response to the treatment.

**Results**
Of the 102 patients with PCOS, 67.6% group of 21-30 years.
75 women (73.52%) were married and 27 (26.47%) were unmarried women.

Chief complaints in the married woman group.

<table>
<thead>
<tr>
<th>Table 1: Chief complaints</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Presenting complaints</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Infertility</td>
<td>75</td>
<td>73.50</td>
</tr>
<tr>
<td>Primary infertility</td>
<td>57</td>
<td>78.66</td>
</tr>
<tr>
<td>Secondary infertility</td>
<td>18</td>
<td>21.33</td>
</tr>
<tr>
<td>Hirsutism</td>
<td>26</td>
<td>25.49</td>
</tr>
<tr>
<td>Menstrual irregularities</td>
<td>56</td>
<td>74.66</td>
</tr>
<tr>
<td>Obesity</td>
<td>47</td>
<td>46.07</td>
</tr>
<tr>
<td>Early virilisation</td>
<td>02</td>
<td>01.96</td>
</tr>
<tr>
<td>Total</td>
<td>75</td>
<td></td>
</tr>
</tbody>
</table>

Infertility (73.5%) was their chief complaint along with menstrual irregularities (74.66%) and obesity (46.07%) and Virilisation in 1.96%.
Table 2: Symptomatology of the 27 unmarried women

<table>
<thead>
<tr>
<th>Presenting complaints</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hirsutism</td>
<td>13</td>
<td>48.14</td>
</tr>
<tr>
<td>Menstrual irregularities</td>
<td>27</td>
<td>100</td>
</tr>
<tr>
<td>Obesity</td>
<td>14</td>
<td>51.85</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>27</strong></td>
<td><strong>100</strong></td>
</tr>
</tbody>
</table>

All had menstrual irregularities, (100%) followed by obesity (51.85%) and hirsutism (48.14%).

Table 3: Investigations

<table>
<thead>
<tr>
<th>Investigation</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random blood sugar levels (&gt; 120mg/dl)</td>
<td>20</td>
<td>19.60</td>
</tr>
<tr>
<td>Serum luteinising hormone level (&gt; 6.7 IU/L)</td>
<td>57</td>
<td>55.88</td>
</tr>
<tr>
<td>Serum Follicle stimulating hormone level (&gt; 10.5 IU/L)</td>
<td>16</td>
<td>15.68</td>
</tr>
<tr>
<td>Serum LH : FSH ration (μ 2:1)</td>
<td>32</td>
<td>31.37</td>
</tr>
<tr>
<td>Serum testosterone level (&gt; 10 ng/ml)</td>
<td>88</td>
<td>86.27</td>
</tr>
<tr>
<td>Serum prolactin level (&gt; 20 pg/ml)</td>
<td>23</td>
<td>18.62</td>
</tr>
<tr>
<td>Serum TSH level (&gt; 6.5 μu/ml)</td>
<td>07</td>
<td>6.86</td>
</tr>
<tr>
<td>21 days progesterone level (&lt; 10 ng/ml)</td>
<td>02</td>
<td>2.66</td>
</tr>
</tbody>
</table>

20 of them (19.60%) had glucose intolerance, serum LH was raised in 57 (55.88%) of patients, serum FSH was significant in 16 (15.68%) of patients, but serum LH:FSH ratio of 2:1 was seen in 32 (31.37%) of the study group.

Hyperandrogenemia was seen in 88 (86.27%) patients, 23 (18.62%) of cases had hyperprolactenemic, serum TSH was raised in 07 (6.86%), 2 (2.66%) had a luteal phase defect as detected by the 21 day progesterone levels.

Table 4: Ultrasound examination

<table>
<thead>
<tr>
<th>Scan findings</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Done in</td>
<td>91</td>
<td>89.21</td>
</tr>
<tr>
<td>Not done in</td>
<td>11</td>
<td>10.78</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>102</strong></td>
<td><strong>100</strong></td>
</tr>
<tr>
<td>Typical PCO picture seen in</td>
<td>79</td>
<td>86.81</td>
</tr>
<tr>
<td>Normal ovaries seen in</td>
<td>12</td>
<td>13.18</td>
</tr>
</tbody>
</table>

Typical picture of polycystic ovaries was seen in 79 cases (86.81%) and the findings of normal ovaries was seen in 12 cases (13.18%).

Table 5: Findings on diagnostic laparoscopy

<table>
<thead>
<tr>
<th>Findings</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Polycystic ovaries</td>
<td>26</td>
<td>86.66</td>
</tr>
<tr>
<td>Normal ovaries</td>
<td>03</td>
<td>10.00</td>
</tr>
<tr>
<td>Hyperthecosis</td>
<td>01</td>
<td>3.33</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>30</strong></td>
<td><strong>100</strong></td>
</tr>
<tr>
<td>Ovarian drilling done in</td>
<td>27</td>
<td>90</td>
</tr>
<tr>
<td>Ovarian drilling not done in</td>
<td>03</td>
<td>10</td>
</tr>
<tr>
<td>Tubal patency</td>
<td>30</td>
<td>100</td>
</tr>
</tbody>
</table>

On diagnostic laparoscopy, 26 (86.66%) had polycystic ovaries and 3 (10%) had normal ovaries, 1 (3.33%) had the severe form of polycystic ovaries.

Table 6: Outcome of pregnancy

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Full term normal delivery</td>
<td>9</td>
<td>42.85</td>
</tr>
<tr>
<td>Emergency LSCS (Obstetric indications)</td>
<td>2</td>
<td>9.52</td>
</tr>
<tr>
<td>Missed abortion</td>
<td>2</td>
<td>9.52</td>
</tr>
<tr>
<td>Inevitable abortion</td>
<td>1</td>
<td>4.76</td>
</tr>
<tr>
<td>Ongoing pregnancy (&gt; 28 weeks)</td>
<td>2</td>
<td>9.52</td>
</tr>
<tr>
<td>Ongoing pregnancy (&lt; 28 weeks)</td>
<td>5</td>
<td>23.80</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>21</strong></td>
<td><strong>100</strong></td>
</tr>
</tbody>
</table>

Of the 21 patients who conceived, 9 of them (42.85%) had full term deliveries, 2 (9.52%) underwent an emergency LSCS for fetal distress, while the abortion rate was (14.28%) and 7 (33.32%) with ongoing pregnancy, of whom 2 (9.52%) have already crossed the period of viability, with no ectopic / multiple gestation.

Table 7: Other modalities of treatment

<table>
<thead>
<tr>
<th>Other modalities of treatment</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>CC + B – Crip + HCG</td>
<td>04</td>
</tr>
<tr>
<td>CC + Dexa + B-crip</td>
<td>04</td>
</tr>
<tr>
<td>CC + HCG + Eltroxin</td>
<td>0</td>
</tr>
<tr>
<td>CC + HCG</td>
<td>17</td>
</tr>
<tr>
<td>CC + Metformin</td>
<td>06</td>
</tr>
<tr>
<td>CC + FSH + HMG</td>
<td>01</td>
</tr>
<tr>
<td>Combination therapy</td>
<td>01</td>
</tr>
<tr>
<td>CC + Progesterone supplements</td>
<td>2</td>
</tr>
<tr>
<td>OC alone</td>
<td>1</td>
</tr>
<tr>
<td>Tab Finast + Tab Aldactone</td>
<td>8</td>
</tr>
<tr>
<td>OC + Finast + Aldactone</td>
<td>5</td>
</tr>
<tr>
<td>OC + Finast</td>
<td>2</td>
</tr>
<tr>
<td>OC + Aldactone</td>
<td>4</td>
</tr>
<tr>
<td>Tab Finast</td>
<td>4</td>
</tr>
<tr>
<td>Tab Aldactone</td>
<td>3</td>
</tr>
<tr>
<td>CC + Dexamethasone</td>
<td>12</td>
</tr>
<tr>
<td>CC + B-crip</td>
<td>23</td>
</tr>
<tr>
<td>CC + HMG</td>
<td>01</td>
</tr>
</tbody>
</table>

The above table that there is no single mode of approach and treatment of polycystic ovarian disease but it is simply a multi-modality approach.

Table 8: Outcome of treatment in unmarried women

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Regularization of M. cycles</td>
<td>9</td>
<td>33.33</td>
</tr>
<tr>
<td>Weight reduction</td>
<td>5</td>
<td>18.51</td>
</tr>
<tr>
<td>Improvement in Hirsutism</td>
<td>4</td>
<td>14.81</td>
</tr>
<tr>
<td>Others on followup</td>
<td>9</td>
<td>33.33</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>27</strong></td>
<td><strong>100</strong></td>
</tr>
</tbody>
</table>
With oral contraceptives, 9 (33.33%) had cycle regularization, hirsutism improved with aldactone and finasteride in 14.8% cases.

Table 9: Final outcome of the infertility group

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total patients treated</td>
<td>75</td>
<td>100</td>
</tr>
<tr>
<td>Ovulation induction done in</td>
<td>69</td>
<td>92.0</td>
</tr>
<tr>
<td>Patients who ovulated</td>
<td>61</td>
<td>88.40</td>
</tr>
<tr>
<td>No. of women who conceived</td>
<td>21</td>
<td>34.42</td>
</tr>
<tr>
<td>Unexplained infertility</td>
<td>04</td>
<td>5.33</td>
</tr>
<tr>
<td>Mild OHSS</td>
<td>02</td>
<td>2.89</td>
</tr>
<tr>
<td>On followup</td>
<td>48</td>
<td>69.56</td>
</tr>
</tbody>
</table>

With 88.40% of ovulation induction rates, (34.42%) conceived, with a total unexplained infertility 4 (5.33%) and incidence of mild OHSS occurred in 2 (2.89%) while the others 48 (69.56%) are still on follow up.

Table 10: Correlation of clinical presentation and hormonal imbalances in the infertility group

<table>
<thead>
<tr>
<th>Correlations</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Menstrual irregularities</td>
<td>56</td>
<td>74.66</td>
</tr>
<tr>
<td>Obesity</td>
<td>47</td>
<td>62.66</td>
</tr>
<tr>
<td>Hirsutism</td>
<td>26</td>
<td>34.66</td>
</tr>
<tr>
<td>Virilisation</td>
<td>02</td>
<td>2.66</td>
</tr>
<tr>
<td>Serum LH (&gt; 6.7 Iu/L)</td>
<td>44</td>
<td>58.66</td>
</tr>
<tr>
<td>Serum FSH (&gt; 10.5 Iu/L)</td>
<td>13</td>
<td>17.33</td>
</tr>
<tr>
<td>Serum LH/FSH (≥ 2:1)</td>
<td>26</td>
<td>34.66</td>
</tr>
<tr>
<td>Serum testosterone (&gt; 10 mg/dl)</td>
<td>46</td>
<td>61.33</td>
</tr>
<tr>
<td>Serum prolactin (&gt;10 pg/ml)</td>
<td>16</td>
<td>21.33</td>
</tr>
<tr>
<td>Serum TSH (&gt; 6.5 μu/ml)</td>
<td>07</td>
<td>9.33</td>
</tr>
<tr>
<td>Random blood sugars (&gt; 120 mg/dl)</td>
<td>20</td>
<td>19.60</td>
</tr>
<tr>
<td>Total</td>
<td>75</td>
<td></td>
</tr>
</tbody>
</table>

A significant LH:FSH ratio of ≥ 2:1 was seen in only 34.66% cases but a raised serum testosterone value definitely correlated with the symptoms in 61.33% of these cases. And so serum testosterone can be used as a definitive hormonal test for diagnosing PCOS.

Carbohydrate intolerance, hyperprolactinaemia and hypothyroidism were present in 19%, 21% and 9% respectively.

Table 11: Correlation of scan findings with hormonal values in the infertility group

<table>
<thead>
<tr>
<th>Correlations</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>On transabdominal scan</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Typical PCO picture seen in</td>
<td>63</td>
<td>84.00</td>
</tr>
<tr>
<td>Normal ovaries seen in</td>
<td>12</td>
<td>16.00</td>
</tr>
<tr>
<td>Serum FSH (&gt; 10.5 Iu/L)</td>
<td>13</td>
<td>17.33</td>
</tr>
<tr>
<td>Serum LH (&gt; 6.5 Iu/L)</td>
<td>44</td>
<td>58.66</td>
</tr>
<tr>
<td>Serum LH/FSH ratio (&gt; 2:1)</td>
<td>26</td>
<td>34.66</td>
</tr>
<tr>
<td>Serum testosterone (&gt; 10 mg/ml)</td>
<td>46</td>
<td>61.33</td>
</tr>
<tr>
<td>Total</td>
<td>75</td>
<td></td>
</tr>
</tbody>
</table>

It could be concluded that scan picture correlated with the serum testosterone value and is more definitive in the approach towards the disease.

Statistical Analysis

Analysis shows that a high LH:FSH ratio (p < 0.01) cases fared better with surgical therapy and high testosterone (p < 0.05) cases fared better with medical therapy.

Discussion

Key findings of the results are as follows:

- 81.37% presented with menstrual irregularities, with polycystic ovaries on scan is 86% and 92% of them had atleast one hormonal abnormality to support the diagnosis, correlated with the incidence of 75%, 56.75% and 93% respectively, reported by Adams et al.1
- Results reveal that 26 (34.56%) of infertile patients with abnormal LH/FSH ratio, only 21 (80.76%) had typical polycystic ovarian morphology on ultrasound and 5 (19.23%) had normal ovaries on scan. Ardaens et al in their study have showed that transvaginal scan is more sensitive in diagnosing polycystic ovarian when compared to a transabdominal scan.6
- Elevated LH levels was seen in 58.66% FSH in 17.35% and testosterone in 61.33% and LH/FSH ratio > 2:1 with p value < 0.01 are similar to results of the study conducted by Robinson et al.7,8
- Ovulation induction rates of 88.4%, with average follicular size 19 x 21 mm and 41.93% conception rates correlate with Rajan et al9 and abortion rates of 13% correlated with Compo et al.10

Conclusion

In polycystic ovarian syndrome there is no single test as the gold standard to prove the disease, as it is a cohort of abnormalities requiring a multi modality approach towards the disease.

In this study, there was an overall improvement of symptoms in 66.66% of unmarried women with PCOD, after appropriate treatment.

In the infertility group of patients, the ovulation rate of 88.40%, pregnancy rate of 34.43% and the pregnancy rate after laparoscopy / drilling of 26.7% was acceptable statistically. With a significant difference between the medical and surgical modalities of management with respect to a high LH:FSH ratio and testosterone values.

Proper selection of the patients, intense monitoring and early detection of complications go a long way towards the success of conception.

References


A pilot study of relation of polycystic ovary syndrome and metabolic disorder in Davangere, Karnataka

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1Assistant Professor, 2Associate Professor, 3Professor and Head, Department of Obstetrics and Gynaecology, 4Assistant Professor, Department of Anatomy, 5Associate Professor, Department of Forensic Medicine & Toxicology, 6Associate Professor, Department of Surgical Oncology, S.S.Institute of Medical Sciences & Research Centre, Davangere-577005, Karnataka, India

Abstract

Introduction
Polycystic ovarian syndrome (PCOS) is seen in 5 – 10% of women of the reproductive age, is the most common form of anovulatory infertility1,2,3.
Evidence prove that these women exhibit a characteristic dyslipidaemia non insulin dependent diabetes and cardiovascular disease in later life, which are the hallmarks of the metabolic syndrome4.
Presence of 3 of 5 common cardiovascular risk factors as a criteria, the prevalence of metabolic syndrome in PCOS is very high (43 – 46%)5.

Aim of the Study
Aim is to establish the relation between PCOS and metabolic syndrome.

Material and Methods

Study Design
A prospective, noninterventional case control study was conducted with 60 primary infertility patients, attending the gynaecological department, S.S. Institute of Medical Sciences and Research Centre, Davangere, Karnataka.
Study population: All these patients had clinical features suggestive of PCOS, of these 30 patients with ultrasound findings of polycystic ovaries were taken as cases and the others were age matched contemporary controls.
Age group: 20 – 30 years.
Duration of study: 8 months. (Dec. 2009 – July 2010)

Methodology
After a brief history, waist circumference and blood pressure was recorded, then patients were asked to come on overnight fasting for a fasting blood sugar level and lipid profile assessment.

Results
In the present study all the five parameters needed for diagnosing metabolic syndrome were raised in PCOS cases and compared to controls. Values were statistically significant by student’s unpaired ‘t’ test.
An incidence of 38% of metabolic syndrome was noted in the PCOS cases studied.

Conclusion
Women with PCOS were strongly associated with metabolic syndrome when compared with the controls. Obesity, high triglycerides and low high density lipoprotein levels are closely linked to insulin resistance and they are independent predictors of myocardial infarction and cardiovascular disease.
Results strongly indicate the need for comprehensive screening and education program for women of all ages with PCOS. Modification of lifestyle factors such as diet and exercise along with insulin sensitizers and lipid lowering agents can prevent long term health risks.

Introduction
Polycystic ovarian syndrome (PCOS) is the most commonly encountered endocrinopathy in women of reproductive age (5-10%)3 with significant reproductive and non reproductive consequences.6
The existence of two of the following criteria like oligoovulation, anovulation, clinical or biochemical signs of hyperandrogenism and polycystic ovaries on ultrasound is required to make the diagnosis of PCOS, as per the American society of reproductive medicine6. PCOS patients are at a higher risk for the metabolic syndrome, which is a group of cardiovascular risk factors that include dyslipidaemia, type II diabetes mellitus, hypertension and obesity5. Hyperinsulinemia is noted in 50-70% of these patients while the rates of obesity ranges from 38-87%5. Thus, polycystic ovarian syndrome seems to have many of the hallmarks of the metabolic syndrome and should no longer be considered a purely gynaecological disorders.
Affected women seem to have subclinical insulin resistance and a form of the metabolic syndrome that
manifests itself in early adult life with gynaecological symptoms. They may therefore gain particular benefit from early screening for metabolic syndrome.7

Criteria used to define metabolic syndrome

Most widely adapted are the criteria proposed by the American National Cholesterol Education Program (NCEP). Adult Treatment Panel III (ATP-III)). These criteria require the presence of 3 of 5 common CV risk factors (increased waist circumference, blood pressure, elevated fasting blood glucose, low serum high-density lipoprotein (HDL)-cholesterol, and increased triglycerides). Using these criteria, the prevalence in PCOS has been reported to be extremely high, 43-46%, primarily on the basis of abnormal lipids and increased waist circumference.5

Incidence

Metabolic syndrome (MBS) which is a common disorder related to visceral obesity and insulin resistance (IR) is associated with atherosclerosis and cardiovascular (CV) disease. The prevalence of MBS is high, occurring in 23.7% of the USA population over 20 years. The prevalence also increases with age, from 6.7% in the third decade to 43.5% in the 7th decade.5

Aim of the Study

To establish the relation between PCOS and metabolic syndrome.

Study Objectives

Primary objective

1) Correlating fasting blood glucose levels with systolic and diastolic blood pressure of cases to control groups.
2) To compare waist circumference with total cholesterol values in both cases and control population.
3) To correlate total triglyceride (TTG), high density lipoprotein – C (HDL-C) values and TTG/HDL-C ratio in PCOS cases and controls.

Secondary objective

1) To compare the obese and the nonobese of the PCOS group with respect to insulin resistance (FBS), total triglycerides and HDL-C levels as a predictor for cardiovascular disease.
2) To know the prevalence of raised total triglycerides to HDL-C ratio in PCOS women as a marker for atherosclerosis in later life.

Study design and study population

Study design: A prospective Case control hospital based non interventional study.

Study population: 50 primary infertility patients attending the gynaecological outpatient department at S.S. Institute of Medical Science and Research Institute, Davangere, Karnataka.

Cases : 25 primary infertility patients with an ultrasound diagnosis of polycystic ovaries.
Controls: 25 primary infertility patients with a normal pelvic study on ultrasound examination.

Study start date: November 2009
Completion date: April 2010

Subject eligibility criteria

Sample size: A total of 50 primary infertility patients in the age group 18-35 years will be randomized into the study.

The study will be conducted at the gynaecological, radiological and biochemistry departments at S.S. Institute of Medical Sciences and Research Institute Davangere, Karnataka 05.

Inclusion criteria

• Age: 18 – 35 years.
• Marital status: Married
• Primary infertility patients at gynaecology outpatient department with clinical features of PCOS like abdominal obesity ie., increased waist circumference (> 35 inches), hirsutism, menstrual irregularities and/or ultrasound findings are taken as physically identifiable factors for detecting PCOS.
• Informed consent.

Exclusion criteria

• Patients who refuse to participate in the study.
• Secondary infertility patients excluded.
• Cases of primary infertility of non PCOS group will be excluded.

Study procedure

At the outpatient department the investigator will complete the history protocol, informed consent taken and their height, weight and waist circumference is measured.

Body weight being recorded to the nearest 0.1 kg using a standard balance scale with subjects bare foot. Body height being measured to 0.1 cm with free standing magnimeter stadiometer. Body mass index (BMI) will then be assessed waist circumference measured in inches.

Patient is asked to empty her bladder and lie down in supine position, privacy maintained. A blood pressure is checked in supine position, right upper arm, Korotkov V sound used for diastolic pressure measurement.

A general physical examination and a pelvic examination is done as a routine examination.

Then the patients will be sent to the radiology department for a pelvic ultrasound and if the scan shows a polycystic picture, they will be included in the study as cases and those with normal pelvic study on ultrasound examination will be considered and controls.

Later these patients will be asked to come fasting the next
day morning at 9 am to the biochemistry department and blood samples will be taken for the measurement of following parameters.

1) Fasting glucose.
2) Lipid profile
   a) Total triglycerides
   b) Total cholesterol
   c) HDL

**Normal reference values**

\[
\text{BMI} = \frac{\text{wt (kg)}}{\text{ht sqm}^2}
\]

- Normal 18.5 – 24.9
- Underweight Low < 18.5
- Obese high > 30
- Overweight 25 – 29.9

**Waist circumference**

> 35 inches is significant, the WHO clinical criteria for metabolic syndrome uses BMI > 30 and a waist circumference > 35 inches.

Blood pressure range (supine position) (>130 / >85 mmHg)

- Systolic 120 mmHg
- Diastolic 80 mmHg

1) Fasting blood glucose will be done by glucose Drytech (DT method) and a reference normal range will be 60-100 mg/dl.

2) Lipid profile will be done using cholesterol Drytech DT method for total cholesterol levels.

**Reference interval**

<table>
<thead>
<tr>
<th></th>
<th>Conventional units mg/dl</th>
<th>SI units mmol/L</th>
</tr>
</thead>
<tbody>
<tr>
<td>Desirable</td>
<td>&lt; 200</td>
<td>&lt; 5.2</td>
</tr>
<tr>
<td>Borderline (high)</td>
<td>200 – 239</td>
<td>5.2 – 6.2</td>
</tr>
<tr>
<td>High</td>
<td>&gt; 240</td>
<td>&gt; 6.2</td>
</tr>
</tbody>
</table>

3) HDL cholesterol levels using dry tech (DT) method, using end point caloximetric model.

**Reference interval for HDL conventional units**

<table>
<thead>
<tr>
<th></th>
<th>&lt; 40</th>
<th>&gt; 60</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low</td>
<td></td>
<td></td>
</tr>
<tr>
<td>High</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

4) Triglycerides are analyzed using the Drytech (DT) method, calorimetric model, with reference intervals for Trig DT.

**Results**

**Tabulation**

1) Correlation between fasting blood glucose levels with systolic and diastolic blood pressure of cases to control groups.

<table>
<thead>
<tr>
<th></th>
<th>Controls FBG (mg%)</th>
<th>Systolic mmHg</th>
<th>Diastolic mmHg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average</td>
<td>84.84</td>
<td>109.2</td>
<td>77.6</td>
</tr>
<tr>
<td>S.D. ±</td>
<td>5.17</td>
<td>7.44</td>
<td>4.35</td>
</tr>
<tr>
<td>PCOS cases</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average</td>
<td>101.36</td>
<td>126</td>
<td>84.4</td>
</tr>
<tr>
<td>S.D. ±</td>
<td>10.53</td>
<td>13.54</td>
<td>5.06</td>
</tr>
<tr>
<td>p value</td>
<td>p &lt; 0.001</td>
<td>p &lt; 0.001</td>
<td>p &lt; 0.001</td>
</tr>
<tr>
<td>Significance</td>
<td>Significant</td>
<td>Significant</td>
<td>Significant</td>
</tr>
</tbody>
</table>

A raised fasting blood glucose levels, rise in systolic and diastolic blood pressure was noted in the cases and the values are statistically significant.

2) To compare waist circumference with total cholesterol levels in both cases and control population.

<table>
<thead>
<tr>
<th></th>
<th>Controls Waist circumference (inches)</th>
<th>Total cholesterol (mg%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average</td>
<td>33.36</td>
<td>197.12</td>
</tr>
<tr>
<td>S.D. ±</td>
<td>1.25</td>
<td>7.41</td>
</tr>
<tr>
<td>PCOS cases</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Weist circumference (inches)</td>
<td>36.96</td>
<td>200.12</td>
</tr>
<tr>
<td>Total cholesterol (mg%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>S.D. ±</td>
<td>2.95</td>
<td></td>
</tr>
<tr>
<td>p value</td>
<td>p &lt; 0.001</td>
<td>p &lt; 0.001</td>
</tr>
<tr>
<td>Significance</td>
<td>Significant</td>
<td>Less significant</td>
</tr>
</tbody>
</table>

Waist circumference and total cholesterol levels too were higher in PCOS cases.

3) To correlate total triglyceride (TTG), high density lipoprotein C (HDL-C) values and TTG/HDL-C ratio in PCOS cases and controls.

<table>
<thead>
<tr>
<th></th>
<th>Controls TTG (mg%)</th>
<th>HDL-C (mg%)</th>
<th>TTG/HDL-C ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average</td>
<td>134.48</td>
<td>49.68</td>
<td>2.71</td>
</tr>
<tr>
<td>S.D. ±</td>
<td>9.00</td>
<td>3.11</td>
<td>0.17</td>
</tr>
<tr>
<td>PCOS cases</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average</td>
<td>157.68</td>
<td>43.38</td>
<td>3.65</td>
</tr>
<tr>
<td>S.D. ±</td>
<td>6.33</td>
<td>3.29</td>
<td>0.31</td>
</tr>
<tr>
<td>p value</td>
<td>p &lt; 0.001</td>
<td>p &lt; 0.001</td>
<td>p &lt; 0.0005</td>
</tr>
<tr>
<td>Significance</td>
<td>Significant</td>
<td>Significant</td>
<td>Highly significant</td>
</tr>
</tbody>
</table>

The patients are instructed to take a non fatty meal the previous night.

Health education will then be given to patients regarding modification of life style factors such as diet and exercise to modify risk factors such as diet and exercise to prevent long term effects of polycystic ovarian syndrome.

The infertility management will then continue as per the standard protocol of the respective unit chiefs.

Qualitative data were analyzed by using students unpaired t-test odds ratio.
Total triglycerides levels and high density lipoprotein cholesterol levels were higher in PCOS cases when compared to controls, with p value being significant (p < 0.001).

Ratio of total triglycerides to high density lipoprotein cholesterol levels in PCOS cases is highly significant (p < 0.0005).

4) Analyzing the PCOS group with respect to BMI, FBS, total triglycerides and HDL-C levels.

<table>
<thead>
<tr>
<th></th>
<th>Obese PCOS</th>
<th>FBS mg/dl</th>
<th>TTG (mg/d)</th>
<th>HDL-C (mg%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average</td>
<td>36.96</td>
<td>101.36</td>
<td>157.68</td>
<td>43.38</td>
</tr>
<tr>
<td>S.D ±</td>
<td>2.95</td>
<td>10.53</td>
<td>6.33</td>
<td>3.29</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Nonobese PCOS</th>
<th>FBS mg/dl</th>
<th>TTG (mg/d)</th>
<th>HDL-C (mg%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average</td>
<td>33.26</td>
<td>84.84</td>
<td>134.48</td>
<td>49.68</td>
</tr>
<tr>
<td>S.D ±</td>
<td>1.25</td>
<td>5.17</td>
<td>9.00</td>
<td>3.11</td>
</tr>
<tr>
<td>p value</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

Significance: Significant Significant Significant Significant Significant

Above findings, depict that the FBS, TTG and HDL-C values were significantly higher in obese PCOS cases as compared to nonobese PCOS cases.

Discussion

In our study women with abdominal obesity i.e. increased waist circumference (> 35 inches) and / or ultra-sound findings are taken as physically identifiable factors for detecting PCOS cases. Ultra-sound findings are similar in women with abdominal obesity and in a smaller group of lean patients with PCOS. Although categorical abdominal obesity (waist circumference > 35 inches) is most strongly associated with metabolic syndrome, the world health organization clinical criteria for metabolic syndrome used BMI more than 30 or waist/hip ratio interchangeably. Our findings are similar to the prevalence of metabolic syndrome in women with PCOS reported using abdominal waist circumference.

The results of our study indicate that women with PCOS are strongly associated with metabolic syndrome compare with age-matched, increase of metabolic syndrome in women with PCOS compared with age matched contemporary controls as per NHANES III (3rd National Health and Nutrition Examination Survey Study). Abdominal obesity, abnormalities in HDL-C, and triglycerides were most frequently detected in women with PCOS.

Obesity and insulin resistance have been implicated in the pathophysiology of metabolic syndrome due to its association with hypertension, hyperglycemia, and dyslipidemia. In our study group, some nonobese women exhibited metabolic syndrome. High triglycerides and low HDL-C levels are closely linked to insulin resistance and are independent predictors of myocardial infarction and cardiovascular disease.

Increase triglycerides/HDL – C ratio is also a marker for atherogenesis with small, dense LDL particles. In a study the findings revealed that triglycerides/ HDL – C more than 3.2 had high sensitivity and specificity, suggestion the possible use of this cutoff to screen for metabolic syndrome.

Conclusion

Polycystic ovarian syndrome presents with overt symptoms of infertility, hirsutism and acne and is strongly related to obesity and chronic hyperinsulinemia this is well established after National Cholesterol Education Program and Adult Treatment Panel III. In the present study most of the women showing more than three risk factors out of five risk factors of metabolic syndrome. Risk factors may not always progress to disease. Treatment of the individual components of the syndrome, including dyslipidemia, obesity and hypertension clearly decrease the incidence of cardiovascular disease. Therefore, the approach requires emphasis on the modification of lifestyle factors such as diet and exercise to modify risk factors in preventing clinical disease.

Young women diagnosed with PCOS should be informed of the possible long term risks to health. Healthcare providers should educate these women regarding tranquil life style with an appropriate diet and exercise program. Early pharmacotherapy with insulin sensitizers and lipid lowering agents may be considered if therapeutic lifestyle changes are unsuccessful.

References

Clinical use and safety of medical method of first trimester Abortion

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Abstract

Introduction
Women have a right to undergo an abortion if she wishes. Medical abortion is legally accepted under the Indian law. Mifepristone and misoprostol combination offers a safe, nonsurgical alternative for pregnancy termination with good success rates.

Aims and Objectives
The purpose of this study was to evaluation of the clinical efficacy of the combination of mifepristone and misoprostol orally for early termination of pregnancy.

Methods
A prospective clinical study conducted during a one year period included 100 healthy pregnant women consenting for a legal abortion with an ammenorrhoea of no more than 63 days.

Results
This method had a success rate of 94%, with an average induction abortion interval between 40-50 hours, one women needed surgical curettage. No serious side effects expect for nausea was noted.

Conclusion
The combination of mifepristone and misoprostol is effective for the termination of early pregnancy in terms of success, tolerance, safety and practicality.

Introduction
Medical termination of pregnancy today has become a way of life. Women have come a long way since the days, when abortion was illegal, secret and socially unacceptable procedure, hidden from family members. The Indian parliament liberalized the abortion laws of the country as a measure of the socioeconomic need of the country. The aim of this social legislation was to remove the social stigma attached to this procedure and to minimize maternal morbidity and mortality arising out of the need to go to quacks for procuring an abortion.

Public education and awareness have led to a healthy trend where women now seek for abortion earlier in the pregnancy. The medical termination act in India was implemented from April 1972 and specifies the place, duration of pregnancy and the person licenced to perform the procedure1.

Though early first trimester surgical termination of pregnancy is safe and simple, but needs a well trained doctor, operation theatre and anaesthesia facilities to perform. About 40-50 million abortions are estimated to take place annually in the world, more than half are illegal and performed by unskilled persons in unhygienic conditions2.

Today, we have the best option of using an antiprogestational drug Ru 486 (Mifepristone) along with PGE, (Misoprostol) for termination of early pregnancy. This combination offers a safe, non surgical alternative for pregnancy termination, which is readily acceptable by women Ru 486 when administered alone, succeeds in termination of pregnancy in about 75% of patients, but when combined with a prostaglandin analogue the success rate goes upto 95-98%1.

Background and Rationale
The “abortion pill” is mifepristone (Ru 486), a progesterone antagonist leads to decidual necrosis and detachment of the products of conception3 and endogenous prostaglandin activity,4 was approved by the FDA in 2000. Prostalandin E, is a prostaglandin analogue, both drugs are active on oral administration.

Aim of the Study
To know the clinical efficacy of the combination of Ru 486 and prostaglandin E1 misoprostol, orally for early termination of pregnancy.

Objectives

• To evaluate the effectiveness of medical methods for early termination of pregnancy with 9 weeks of gestation.
• To evaluate the morbidity and mortality associated with medical method of early termination of pregnancy.

Study design and study population
Type of study: Prospective interventional hospital based study.

Place of study: Women attending the gynaecology outpatient department at S.S. Institute of Medical Sciences and Research Centre requesting for early pregnancy termination.

Study eligibility criteria
The criteria was that a total of 100 consecutive patients requesting for early pregnancy termination in a period of 1 year from September 2009 to September 2010 were recruited for the study.

Inclusion criteria
• Patient requesting for abortion, with pregnancy less than 9 weeks of gestation
• Pregnancy confirmed by clinical examination, and ultrasound examination.
• Singleton, intrauterine gestation
• Willing to terminate pregnancy by surgical method, in case of failure of treatment.
• Informed consent taken.

Exclusion criteria
• Cardiovascular disorder and other serious medical disorder.
• Gestational age more than 9 weeks.
• Threatened abortion cases.
• Patient age more than 35years
• Smoking
• Asthma
• Hemoglobin less than 8g%.

Methodology
Study procedure
Women fulfilling the criteria underwent a thorough general physical examination, systemic examination including cardiovascular respiratory, per abdomen, per speculum and per vaginal examination was done.

Investigations
• Urine for albumin, sugar, microscopy
• Blood for hemoglobin, blood group and Rh typing, HIV and HbsAg status
• Urine pregnancy test.
• Pelvic ultrasound

Informed consent was taken. Then on
Day one: 200mg Ru 486 was given orally to the women on outpatient basis.

Day three: after 36 hours, cases were reviewed and received two tablets of 200 mcg of misoprostol and 6 hours later another single tablet of 200mg of misoprostol was given. So the women received 600mcg of misoprostol in total. Patients being informed about possible side effects like vomiting, diarrhea, abdominal pain, expulsion of product of conception.

On the tenth day cases were reviewed for number of days of bleeding, pain abdomen and any other side effects. Pelvic ultrasound confirmed the completeness of the abortion.

Assessing outcome of the treatment
Successful outcome
Success was defined as complete expulsion of the conceptus, without the need for a surgical procedure.

The outcome of treatment was considered as successful when the following criteria were fulfilled,
• History of onset of bleeding per vagina.
• History of expulsion of products of conception
• Decrease in uterine size
• Pelvic ultrasound showing empty uterine cavity with no retained products.
• No intervention required.

Unsuccessful outcome
This comprised of incomplete abortion or continuing of pregnancy or trial interruption. In these conditions pregnancy is terminated or abortion process is completed by appropriate surgical procedures. The tissue obtained should be sent for histopathological examination.

1) Incomplete abortion: The outcome of treatment was considered as incomplete abortion when the following conditions were fulfilled.
• History of bleeding per vagina
• Uterine size remaining at pretreatment size or bulky
• Ultrasound report showing retained products of conception.
• Some intervention was required to complete the abortion.

2) Continuation of pregnancy
The outcome of treatment was considered as continuation of pregnancy when the
• Size of uterus increased from pretreatment assessment.
• Ultrasound report showing ongoing pregnancy.
• Bleeding may or may not have started.

3) Trial interruption: The outcome of treatment was also considered as unsuccessful when the pregnancy had to be interrupted before day 10 due to,
• Heavy bleeding
• Other medical reasons.

Results
1) Age distribution

<table>
<thead>
<tr>
<th>Age in years</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>20-25 years</td>
<td>34</td>
<td>34%</td>
</tr>
<tr>
<td>26-30 years</td>
<td>62</td>
<td>62%</td>
</tr>
<tr>
<td>31-35 years</td>
<td>4</td>
<td>4%</td>
</tr>
</tbody>
</table>

Majority of them belonged to the 26-30 years age group.

2) Parity Index

<table>
<thead>
<tr>
<th>Gravida</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0</td>
<td>0%</td>
</tr>
<tr>
<td>2</td>
<td>42</td>
<td>42%</td>
</tr>
<tr>
<td>3</td>
<td>52</td>
<td>52%</td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4%</td>
</tr>
<tr>
<td>5</td>
<td>2</td>
<td>2%</td>
</tr>
</tbody>
</table>

None of them were primigravidas and majority of them were second and third gravidas.

3) Period of gestation

<table>
<thead>
<tr>
<th>Weeks in gestation</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 6 weeks</td>
<td>24</td>
<td>24%</td>
</tr>
<tr>
<td>&gt; 6 weeks with in 9 weeks</td>
<td>76</td>
<td>76%</td>
</tr>
</tbody>
</table>

 Majority of the women were beyond 6 weeks of gestation.

4) Induction abortion interval

<table>
<thead>
<tr>
<th>Time in hours</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>40 – 50 hours</td>
<td>90</td>
<td>90%</td>
</tr>
<tr>
<td>51 – 60 hours</td>
<td>6</td>
<td>6%</td>
</tr>
<tr>
<td>61 – 70 hours</td>
<td>4</td>
<td>4%</td>
</tr>
</tbody>
</table>

90% of women had an induction abortion interval of around 40-50 hours.

5) Comparison with gestational age and induction abortion time

<table>
<thead>
<tr>
<th>Gestation in weeks</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 6 weeks</td>
<td>20</td>
<td>40-50 hours</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>46-50 hours</td>
</tr>
<tr>
<td>&gt; 6 weeks with in 9 weeks</td>
<td>66</td>
<td>40-50 hours</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>51-60 hours</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>61-70 hours</td>
</tr>
</tbody>
</table>

24 women with gestational age less than 6 weeks, had an induction abortion interval within 40-50 hours, all of them had a complete abortion, while in the remaining who had a gestational age of more than 6 weeks but less than 9 weeks, 66 women had an abortion in 40-50 hours time, 6 of them required 51 – 60 hours and another 4 of them took almost 61-70 hours for an abortion.

6) Side effects

<table>
<thead>
<tr>
<th>Side effects</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nausea</td>
<td>52</td>
<td>52%</td>
</tr>
<tr>
<td>Vomiting</td>
<td>6</td>
<td>6%</td>
</tr>
<tr>
<td>Severe abdominal pain</td>
<td>8</td>
<td>8%</td>
</tr>
<tr>
<td>Chills</td>
<td>2</td>
<td>2%</td>
</tr>
<tr>
<td>Diarrhoea</td>
<td>2</td>
<td>2%</td>
</tr>
</tbody>
</table>

Nausea was the commonest (52%) side effect noted, severe abdominal pain during the abortion process was seen in 8% of women in the study.

7) Success rate

<table>
<thead>
<tr>
<th>Results</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complete abortion</td>
<td>94</td>
<td>94%</td>
</tr>
<tr>
<td>Incomplete abortion</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Continuation of pregnancy</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lost to follow – up</td>
<td>4</td>
<td>4%</td>
</tr>
<tr>
<td>Trial of interruption</td>
<td>2</td>
<td>2%</td>
</tr>
</tbody>
</table>

94% of them had a complete abortion, 2 women needed trial of interruption as they started to have heavy bleeding after 11 hours of expulsion of the products of conception, requiring emergency curettage. 4 cases were lost followup.

Discussion

The results of our study reveal a success rate of 94%, with less morbidity and without any mortality. This can be compared with similar success rates of 94% by Mc Kinley et al and 95% (Sang et al)5.

Ru 486 when administered alone succeeds in 75% of early trimester pregnancies but when followed by a prostaglandin analogue achieves complete abortion in 95-98%.6

The age and the parity index of the women did not alter the outcome in the present study.

Side effects due to the drugs used were of minor concern, more so nausea (52%), pain abdomen (8%) and vomiting (6%) were observed. An antispasmodic was prescribed in a few cases (4%). Similar side effects noted by studies conducted by Peyron et al7 and Refaey et al. These side effects were mainly due to prostaglandin administration.

Efficacy of this method of medical abortion seems to be clearly linked to gestational age, the failure rate is increased in pregnancies with higher gestational age, as shown by Norman et al3 who included women in 8 weeks of gestation and showed low failure rates.

The induction abortion interval was between 40-50 hours in 90% of cases and it is within 51-60 hours in 6% of cases and within 61-70 hours in 4% of cases, can be compared with Peyron et al7, who had an average of 12-36 hours
interval. Trial of interruption with an emergency check curettage was done in one woman (1%) and 2 cases did not turn up for followup on the 10th day. Clinical studies shows failure rate of 2-3%

In spite of the satisfactory results reported here, we wish to emphasize that the abortion procedure should be done under medical supervision. Appropriate counseling of the patients, detailed history to rule out contraindications for usage and proper followup of patients is of importance.

Conclusion
Pregnancy termination is an important maternal health issue. In developing countries, legal abortions carry a mortality rate of 5 per 100000, but the mortality rate due to illegal abortions is 100 per 100000. In India, though abortion was legalized in 1972, illegally induced abortion is a major killer of women. It accounts for one of ten maternal deaths due to lack of information and because facilities cannot meet the demand.

Unmarried teenagers and most of the married women requesting abortion are reluctant to go to a registered clinic or hospital because of want of adequate privacy and fear about surgical procedure. Medical abortion should be made available to abortion seekers through a supervised national programme, it may revolutionise abortion management and decrease maternal deaths, especially in the rural areas.

Attractive features of medical abortion include the ease of swallowing pills simpler, privacy, autonomy and avoidance of psychological effect of physically invasive operation. It is more comfortable and less dangerous. The combination of mifepristone and misoprostol is effective for the termination of early pregnancy in terms of success, tolerance, safety and practicality. It is a desirable option which should be made available to women through the providers of family planning services.

References
5. Wel SG et al. Termination of early pregnancy by two regimens of mifepristone with misoprostol and PGOS – a multicentre randomized clinical trial in china; Contraception; 1994;50:501-510.
Abstract
Breast cancer is the first common cancer among women in Meerut city and western India. All breast cancer cases were collected during 2008-2009, of these 60 (21%) death cases out of 285 were found in many hospitals/home. The survival rates by Kaplan Meier and Estimated survival rate by Logistic Regression for 1, 2 and more than 3 years were found 88.30%, 76.27% & 63.63% and 77.38%, 63.98% & 49.98% respectively. Cancer stage have <5cm displayed poorer survival rate (51.2%) at more than 3 year than stage <2cm with survival rate (72.73%), and the survival rate in religion, Hindu (55.47%) has better than Muslim with survival rate (54.51%), both were statistically significant. More research and interventions are suggested to formulate and implement strategies to educate and increase the knowledge of communities for healthy life style and disease prevention strategies including early detection of breast cancer to control in India.

Key Words
Breast cancer; survival rate by Kaplan Meier and logistic regression; developing countries.

Introduction
Cancer is not a communicable disease. In India, the registration cases require an active tracing for records for cancer patients. Breast cancer is the second most common cancer among women in Meerut city, Delhi, Mumbai and Southern India after cervix cancer. This study depends upon the data of survival rate of female breast cancer patients, who were registered in hospitals and clinical nursing homes in Meerut city. The Meerut city, belong to the Uttar Pradesh with a population of about 1.6 million in 2009. The morbidity data are collected by personal interviews of patients themselves and medical record review. The follow-up data are obtained from obstructing mortality information from Statistical Division of Municipality of Meerut city.

The data was based upon the different parameters and date of diagnose of cancer, method of confirmation of diagnosis (histology, radiography, FNACs, etc) and clinical extent of disease, which was equivalent to tumor stages. Data of different types of tumor stages was based on clinical assessment before the treatment. The criteria use for coding of diseases of clinical extant are define for population based cancer registries.

Material and Methods
This data was taken between the first diagnosis date of cancer (from 1st January, 2008) and closing date of study (31st December, 2009) at Meerut city in Utter Pradesh. Only 60 (21%) breast cancer cases were registered as death, and had been excluded from the study. The remaining 225 (79%) breast cancer cases were found a part of analysis for the present study. In this study used some method as below.

Observed survival was computing by the Kaplan Meier Method and cumulative survival rate at the end of each year of follow-up, the proportion of patients still surviving can be calculated at intervals as short as the accuracy of recording date of death. Log rank test was used to evaluate the prognostic factors in an univariate analysis. The relationship between the prognostic factors and survival in Meerut city was studied using a proportional hazards regression model. Cox Regression Method is useful in situations where we have censured observations, where some of the patients do not die during the observation period. Another important method is used as below.

This study, data was divide by two categories such as complete to follow up belong to 155 cases and incomplete follow up belong to 130 cases out of 285 total cases. In this situation, we find 36 more death probabilities in incomplete data with the base of complete to follow up by using the logistic regression. So that estimated deaths were calculated 96 out of the total cases and check the goodness of this data by using the Cross Validation Method.

Results
The entire variable in table 1 has been observed survival rate by using given methods and table-2 showed the hazard rate of above parameters respectively.

Table1. Showed that survival rate with 95% confidence interval by using the Kaplan Meier Method were found 60 deaths and estimated survival rate by using the logistic regression method were found 96 deaths out of 285 cancer cases.
By age group

The table (1) showed that the survival rate for age group <40, 40-49, 50-59 and 60+ years were found to be 67.93% ,54.23% ,59.19% and 51.28% and found no significant in all the age groups respectively. All age group, the cumulative survival rate was found to be 50.4% with median 51. The cumulative survival rates had been observed 88.30%, 76.27% and 63.63% at 1, 2, and 3+ year respectively. A clearly decreasing trend was found in cumulative survival rate with increasing age. From the table (2) showed that the hazard ratio for the age group 40-49 years was 33% reduced, 50-59 years and ≥ 60-years were found, 16% and 38 % increased respectively for the corresponding age group.

Similarly another method showed that the survival rate for age group 40-49 years was 33% reduced, 50-59 years and ≥ 60-years were found, 16% and 38 % increased respectively for the corresponding age group.

By Menopausal Stage

In this study 127 cases belonging to pre-menopausal Stage were included, which had cumulative survival rate 57.51% and 55.34% and 51 and 37 median survival respectively and 158 cases of post-menopausal Stage had cumulative survival rate of 50.52% with median 50 and 51.86% with 28 median for respectively tables. The result was statistically significant (p < 0.05) by logistic regression methods and the hazard ratio of post menopausal stage was found 46% increased and post menopausal stage was found 66% increased corresponding pre menopausal stage.

By Size of Cancer i.e. stages

There were 65 (23%) breast cancer patients, which diagnosed with a lump size of < 2 cm. and had cumulative survival rates 72.73% and 67.41% from table 1. While 65 (23%) breast cancer cases was obtained from the data with a lump size of 2-5 cm., having cumulative survival rates 73.95% and 55.67%. Where 155 (54%) breast cancer patients had a lump size more than 5 cm. gave the cumulative survival rates 51.2% and 53.06. The elevation in the lump size shows statistically highly significant results (p < 0.001). The risk of dying among those patients, which had lump size more than 5 cm., was approximately 3 times greater than those patients with the lump size of < 2 cm.

By Religion

This factor reveals that 233(82%) breast cancer cases were belong to Hindu religion in Meerut city and had cumulative survival rates 55.47% and 30.52% respectively methods from table 1. While 52 (18%) breast cancer cases were belong to Muslim religion and had

---

Table 1: Depicts the estimate of survival rates by using Kaplan Meier

<table>
<thead>
<tr>
<th>Factor</th>
<th>Total Cases</th>
<th>% Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age &lt;40</td>
<td>59</td>
<td>21</td>
<td>11</td>
<td>30-42</td>
<td>67.93</td>
<td>0.117</td>
<td>17</td>
<td>27-47</td>
<td>50.49</td>
</tr>
<tr>
<td>40-49</td>
<td>92</td>
<td>32</td>
<td>15</td>
<td>31-75</td>
<td>54.23</td>
<td>0.0005</td>
<td>8</td>
<td>34-47</td>
<td>67.41</td>
</tr>
<tr>
<td>50-59</td>
<td>72</td>
<td>25</td>
<td>16</td>
<td>23-61</td>
<td>59.19</td>
<td>0.0079</td>
<td>17</td>
<td>24-76</td>
<td>55.67</td>
</tr>
<tr>
<td>60+</td>
<td>62</td>
<td>22</td>
<td>18</td>
<td>6-72</td>
<td>51.28</td>
<td>0.9197</td>
<td>24</td>
<td>15-43</td>
<td>51.94</td>
</tr>
<tr>
<td>All Ages</td>
<td>285</td>
<td>100</td>
<td>60</td>
<td>37-65</td>
<td>50.40</td>
<td>0.0327</td>
<td>76</td>
<td>21-39</td>
<td>50.40</td>
</tr>
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<table>
<thead>
<tr>
<th>Menopausal Stage</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre</td>
<td>127</td>
<td>45</td>
<td>22</td>
<td>34-68</td>
<td>57.51</td>
<td>0.148</td>
<td>32</td>
<td>23-51</td>
<td>55.34</td>
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<tr>
<td>Post</td>
<td>158</td>
<td>55</td>
<td>38</td>
<td>41-59</td>
<td>50.52</td>
<td>0.0005</td>
<td>64</td>
<td>14-42</td>
<td>51.86</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Stages</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 2 cm</td>
<td>65</td>
<td>23</td>
<td>5</td>
<td>37-50</td>
<td>72.73</td>
<td>0.0005</td>
<td>8</td>
<td>34-47</td>
<td>67.41</td>
</tr>
<tr>
<td>2 to &lt;5 cm</td>
<td>65</td>
<td>23</td>
<td>7</td>
<td>44-56</td>
<td>73.95</td>
<td>0.0079</td>
<td>17</td>
<td>24-76</td>
<td>55.67</td>
</tr>
<tr>
<td>more than 5cm</td>
<td>155</td>
<td>54</td>
<td>48</td>
<td>24-48</td>
<td>51.20</td>
<td>0.0000</td>
<td>71</td>
<td>17-35</td>
<td>53.06</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Religion</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hindu</td>
<td>233</td>
<td>82</td>
<td>43</td>
<td>33-67</td>
<td>55.47</td>
<td>0.0379</td>
<td>72</td>
<td>29-45</td>
<td>50.62</td>
</tr>
<tr>
<td>Muslim</td>
<td>52</td>
<td>18</td>
<td>17</td>
<td>12-62</td>
<td>54.51</td>
<td>0.0917</td>
<td>24</td>
<td>15-43</td>
<td>51.86</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Laterality</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Right</td>
<td>117</td>
<td>41</td>
<td>24</td>
<td>52-66</td>
<td>55.36</td>
<td>0.327</td>
<td>76</td>
<td>21-39</td>
<td>50.40</td>
</tr>
<tr>
<td>Left</td>
<td>168</td>
<td>59</td>
<td>36</td>
<td>31-53</td>
<td>50.96</td>
<td>0.0000</td>
<td>59</td>
<td>23-37</td>
<td>50.80</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Treatment</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th>Estimated Death</th>
<th>C.I</th>
<th>Survival Rate</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Surgery</td>
<td>218</td>
<td>76</td>
<td>46</td>
<td>25-59</td>
<td>52.29</td>
<td>0.0005</td>
<td>76</td>
<td>21-39</td>
<td>50.40</td>
</tr>
<tr>
<td>Radiation</td>
<td>13</td>
<td>4.6</td>
<td>5</td>
<td>32-60</td>
<td>65.45</td>
<td>0.0000</td>
<td>5</td>
<td>32-60</td>
<td>65.45</td>
</tr>
<tr>
<td>Chemotherapy</td>
<td>54</td>
<td>19</td>
<td>9</td>
<td>36-54</td>
<td>65.77</td>
<td>0.0000</td>
<td>15</td>
<td>21-79</td>
<td>55.67</td>
</tr>
</tbody>
</table>

---
cumulative survival rates 54.51% and 51.49%. This result was statistically significant results (p < .01) and the risk of dying of Hindu community is 78% and 49% increase compared to Muslim community breast cancer patients.

By Laterality

Regarding to this parameter, there were 117 (41%) breast cancer patients affected to right breast and they had cumulative survival rates 55.36% and 51.87%. In the case of left breast cancer cases 168(59%) breast cancer patients was found, which had cumulative survival rates 50.96% and 50.80% from the above table and hazard ratio of left breast cancer patients was approximately equal to the right breast cancer patients.

By Treatment

In the present study, 218 (76%) breast cancer cases were treated by surgery, which had cumulative survival rates 51.29% and 50.40% using both methods. Whereas 13(5%) breast cancer cases were observed to treated by radiation and have cumulative survival rates 65.45% and 65.45 % with same median. There were 54 (19%) breast cancer patients also treated by chemotherapy, which had cumulative survival rates 65.77% and the risk of dying of women breast cancer patients which were treated with chemotherapy and radiation had approximately 30% and 50% reduced compared to other treatment such as surgery.

Discussion

The present study provides substantial information about the survival rates of women diagnosed with breast cancer. The survival data is not readily available in many developing and under developed countries because of lack of proper information systems and the difficulties faced for follow up in due to some unavoidable reason of cancer cases until death, and another important factor completeness of cancer registration which give the affecting results of survival analysis. While in complete registration, the complete information of cancer cases is not available in most of the developed and under

Table 2: Hazard Ratio was found of all cases with 95% confidence interval for overall data and Estimated data

<table>
<thead>
<tr>
<th>Factor</th>
<th>Original data with 60 deaths</th>
<th>Estimated data with 96 deaths</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Hazard Ratio</td>
<td>C.I</td>
</tr>
<tr>
<td>Age-group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;40</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>40-49</td>
<td>0.672</td>
<td>(.370-1.471)</td>
</tr>
<tr>
<td>50-59</td>
<td>1.165</td>
<td>(.538-2.524)</td>
</tr>
<tr>
<td>60+</td>
<td>1.389</td>
<td>(.645-2.990)</td>
</tr>
<tr>
<td>Menopausal Stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pre</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>Post</td>
<td>1.463</td>
<td>(.863-2.480)</td>
</tr>
<tr>
<td>Stages</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt; 2 cm</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>2 to &lt;5 cm</td>
<td>0.972</td>
<td>(.307-3.076)</td>
</tr>
<tr>
<td>More than 5cm</td>
<td>3.134</td>
<td>(1.242-7.912)</td>
</tr>
<tr>
<td>Religion</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hindu</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>Muslim</td>
<td>1.786</td>
<td>(1.013-3.149)</td>
</tr>
<tr>
<td>Laterality</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Right</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>Left</td>
<td>0.974</td>
<td>(.580-1.636)</td>
</tr>
<tr>
<td>Treatment</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Surgery</td>
<td>1.000</td>
<td>-</td>
</tr>
<tr>
<td>Radiation</td>
<td>0.915</td>
<td>(.357-2.350)</td>
</tr>
<tr>
<td>Chemotherapy</td>
<td>0.702</td>
<td>(.343-1.438)</td>
</tr>
</tbody>
</table>

Analysis of Survival rates on the base of original and estimated data by Kaplan Meier Method
developed country because of skill problem or financial constraints; the time estimate of the survival data is not available, if we delete this incomplete data. We may have under estimate of survival rate. While in complete registration of cases and major exclusion are likely to bias the survival estimates, inadequate information on several variables would limit interpretation of the results.

An another research showed that the cumulative survival rate was better among older than younger patient and it was challenged by a few studies. Several studies have been shown that the age has a significant prognostic effect even after stratification for stage of disease (6-7) and average annual age-adjusted incidence rate of 22.0 in during (1982-93) in Madras8.

Several studies have shown that breast cancer in younger women especially those less than 40 years old. Several studies have shown that breast cancer in younger and average annual age-adjusted incidence rate of 22.0 and it was challenged by a few studies. Several studies have been shown that the age has a significant prognostic effect even after stratification for stage of disease and average annual age-adjusted incidence rate of 22.0 in during (1982-93) in Madras8.

Several studies have shown that breast cancer in younger women especially those less than 40 years old has a poor prognosis9,10. In present study, it was found that the large proportion of the cases (32%) occurring in the age group 40-49 years, which over half (53%) < 50 years of age, and in the age group < 40 years, the survival rate was found (67.93%) compare with the age >60 years survival rate (51.28%). The overall survival rate was 50.4%, which was poorer compare the above age group by Kaplan Meier Method but Logistic Regression showed that less than 40 years age group had survival rate 50.49% compare with the age group was greater than 60 year age group having survival rate 51.16%. The overall survival rate have 51.30%, was not a poorer the respective age group.

In menopausal stage and survival rate was statistically insignificant and significant due to corresponding methods. The post-menopausal Stage has hazard rate corresponding to pre-menopausal Stage, which are 40% and 60% increased from respective methods. Therefore increase the age of patient give the more risk of breast cancer.

The present work reveals that 23% of cases have tumor size 2-5 cm and 23% have tumor size < 2 cm, while 54% cases have the size of tumor more than 5 cm, which were statistically significant stages and cumulative survival rates. However Fakhro11 observed that 51.3 % cases belong to 2-5 cm cancer size and 70% cases belong to the tumor size < 2 cm.

According to the religion parameter observed that 82% breast cancer cases belong to the Hindu religion and 18% breast cancer cases belong to the Muslim religion. The present findings indicate that the breast cancer risk compare of Hindu and Muslim religion, it was observed that higher elevated breast cancer risk was found in Muslim religion by using Cox regression Method.

There was statistically insignificant association found between cumulative survival rates and laterality during this study, which include 41% breast cancer patient are affected in the right breast and 59% in the right breast. The survival rates to regard this parameter 55.3% and 50.96 % by Kaplan Meir Method and 51.87% and 50.80% with logistic regression. This is due to the effect explaining left side the more aggressive disease present in of breast cancer. The similar study has been done by Arkoob12 and concluded that survival rate 61.1% belong to the right breast and 59.1 % survival rate belong to the left breast in Jordon.

In reviewing the line of treatment, the present study reveals that 52.29% and 50.40% of cases are managed by surgery alone, 65.45% cases are treated by radiation and 65.77% & 55.67% cases are also treated by chemotherapy with both methods respectively.

This type of studies requires identifying the incidence and probability of developing breast cancer in relation to specific demographic variables in Meerut. On the other hand some weaknesses occur in present study such as medical record of many patients was not complete, they were lost to follow up that's why some important information was missing in L.L.R.M. College and in private nursing homes.

References
Urban-rural differences in prevalence of CHD: A population based Study

Mahajan H1, Chaturvedi M2, Lal P3, Saha R4, Grover VL5
1Assistant Professor, Community Medicine, SMS & R, Greater Noida, UP, 2Associate Professor, Community Medicine, SMS&R, Greater Noida, UP, 3Professor, Community Medicine, MAMC, Delhi, 4Assistant Professor, Statistician, Community Medicine, MAMC, Delhi, 5Professor & Head, Community Medicine, SMS&R, Greater Noida, UP

Abstract

Objective
The present study was carried out with the objective to determine and compare the prevalence of CHD in urban and rural communities of Delhi.

Methods
A community-based epidemiological study from Delhi. The study was carried out by conducting a house-to-house survey. WHO Rose Questionnaire was used to detect angina among the study subjects. ECG was recorded using single-channel BPL ECG machine and was reported by a cardiologist.

Results
The overall prevalence of Coronary heart disease was observed to be 12.0% (95%CI: 9.71-14.29). The prevalence of Coronary heart disease was observed to be significantly higher among urban study subjects (14.8%; 95%CI: 11.05-18.55) as compared to that among rural study subjects (9.7%; 95%CI: 6.88-12.52) (p<0.05). Out of the total 92 cases of Coronary heart disease in the present study, 51 (55.5%) cases were detected either on the basis of ECG tracing or Rose questionnaire, while rest of the 41 (44.5%) cases were known cases of Coronary heart disease.

Conclusion
The present study clearly indicates that the prevalence of CHD in India is a major problem of public health importance in India.

Keywords
CHD, prevalence, urban, rural, WHO Rose questionnaire, ECG, public health.

Introduction
Coronary heart disease (CHD) is now the leading cause of death worldwide1. More than 60% of the global burden of CHD occurs in developing countries. Retrospective analysis of previous epidemiological studies on CHD has shown an increase in its prevalence in India. The prevalence of CHD has increased in urban areas of India from 1% in the year 1960 to 11% in the year 2001. Similarly, in the rural areas too, its prevalence has increased from 2% in the year 1974 to 5% in the year 20022,3. To monitor the trends in the prevalence of coronary heart disease, periodically well organized systematic epidemiological studies are required in various geographical areas of India. The present study was carried out with the objective to determine and compare the prevalence of CHD in urban and rural communities of Delhi.
Table 1: Socio-demographic profile of study subjects

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Urban (n=500)</th>
<th>Rural (n=500)</th>
<th>Total (N=1000)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20-29</td>
<td>93 (18.6)</td>
<td>95 (19.0)</td>
<td>188 (18.8)</td>
</tr>
<tr>
<td>30-39</td>
<td>112 (22.4)</td>
<td>128 (25.6)</td>
<td>240 (24.0)</td>
</tr>
<tr>
<td>40-49</td>
<td>87 (17.4)</td>
<td>118 (23.6)</td>
<td>205 (20.5)</td>
</tr>
<tr>
<td>50-59</td>
<td>100 (20.0)</td>
<td>65 (13.0)</td>
<td>165 (16.5)</td>
</tr>
<tr>
<td>60 and above</td>
<td>108 (21.6)</td>
<td>94 (18.8)</td>
<td>202 (20.2)</td>
</tr>
<tr>
<td>Mean ± SD</td>
<td>45.42±15.22</td>
<td>44.32±15.79</td>
<td>44.87±15.52</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>211 (42.2)</td>
<td>224 (44.8)</td>
<td>435 (43.5)**</td>
</tr>
<tr>
<td>Female</td>
<td>289 (57.8)</td>
<td>276 (55.2)</td>
<td>565 (56.5)</td>
</tr>
<tr>
<td>Educational status*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Illiterate</td>
<td>116 (23.2)</td>
<td>111 (22.2)</td>
<td>227 (22.7)**</td>
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<td>Primary/Just literate</td>
<td>61 (12.2)</td>
<td>90 (18.0)</td>
<td>151 (15.1)</td>
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<td>Middle school</td>
<td>73 (14.6)</td>
<td>69 (13.8)</td>
<td>142 (14.2)</td>
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<td>High school</td>
<td>67 (13.4)</td>
<td>114 (22.8)</td>
<td>181 (18.1)</td>
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<td>Intermediate/Post High</td>
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<td>73 (14.6)</td>
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<td>school diploma Graduate</td>
<td>83 (16.6)</td>
<td>28 (5.6)</td>
<td>111 (11.1)</td>
</tr>
<tr>
<td>PG/Professional</td>
<td>31 (6.2)</td>
<td>15 (3.0)</td>
<td>46 (4.6)</td>
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<tr>
<td>Occupation*</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Unemployed</td>
<td>272 (54.4)</td>
<td>288 (57.6)</td>
<td>560 (56.0)b</td>
</tr>
<tr>
<td>Unskilled</td>
<td>24 (4.8)</td>
<td>31 (6.2)</td>
<td>55 (5.5)</td>
</tr>
<tr>
<td>Semi-skilled</td>
<td>37 (7.4)</td>
<td>41 (8.2)</td>
<td>78 (7.8)</td>
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<tr>
<td>Skilled</td>
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<td>57 (11.4)</td>
<td>103 (10.3)</td>
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<td>100 (20.0)</td>
<td>75 (15.0)</td>
<td>175 (17.5)</td>
</tr>
<tr>
<td>Farm-owner</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Semi-professional</td>
<td>14 (2.8)</td>
<td>7 (1.4)</td>
<td>21 (2.1)</td>
</tr>
<tr>
<td>Professional</td>
<td>7 (1.4)</td>
<td>1 (0.2)</td>
<td>8 (0.8)</td>
</tr>
<tr>
<td>Income group**</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lower</td>
<td>36 (7.2)</td>
<td>47 (9.4)</td>
<td>83 (8.3)d</td>
</tr>
<tr>
<td>Upper lower</td>
<td>151 (30.2)</td>
<td>211 (42.2)</td>
<td>362 (36.2)</td>
</tr>
<tr>
<td>Lower middle</td>
<td>137 (27.4)</td>
<td>161 (32.2)</td>
<td>298 (29.8)</td>
</tr>
<tr>
<td>Upper middle</td>
<td>82 (16.4)</td>
<td>52 (10.4)</td>
<td>134 (13.4)</td>
</tr>
<tr>
<td>Upper</td>
<td>94 (18.8)</td>
<td>29 (5.8)</td>
<td>123 (12.3)</td>
</tr>
</tbody>
</table>

Figures in parenthesis are percentages
*According to Modified Kuppuswamy classification
** According to Mahajan-Gupta Socio-economic scale
a. p value not significant; b. p<0.05; c. p<0.01; d. p<0.001

1. Clinically documented coronary heart disease
2. WHO Rose-questionnaire positive angina
3. ECG changes (ST, T or Q waves)4,5

If any one or more than one of these criteria were present, the study subject was considered to be a case of CHD.

Data collected was tabulated and analyzed by using statistical package for social science (SPSS) software. The prevalence was recorded in percentages. Chi-square test or Fisher Exact test was applied for statistical analysis.

**Results**

Table 1 depicts the socio-demographic profile of the study subjects. The study subjects in the urban area ranged from 20-87 years of age, whereas, the subjects in the rural area ranged from 20-90 years of age. The mean age of the urban subjects was 45.42±15.22 yrs while the mean age of the rural subjects was 44.32±15.79 yrs. Amongst all study subjects, 43.5% of the subjects were males and 56.5% of the subjects were females. The overall literacy rate among study subjects was observed to be 77.3%. More than half of the study subjects in both urban and rural areas were housewives/unemployed. Among those who were employed, a large number of subjects were either semi-skilled/skilled workers (18.1%) followed by clerical/shop-owner/farm-owner (17.5%). A small number of study subjects were unskilled workers (5.5%) and very few subjects were either semi-professionals/professionals (2.9%). Majority of the subjects belonged either to lower income group (44.5%) or middle income group (43.2%). Only 12.3% of the subjects belonged to the upper income group.

Table 2 depicts prevalence of Coronary heart disease among the study subjects. Out of 1000 study subjects, 755 subjects underwent ECG (337 in the urban area and 418 in the rural area). Out of the remaining 245 subjects, 12 subjects were either known cases of coronary heart disease or had Rose-questionnaire positive Angina (8 in the urban area and 4 in the rural area). Thus, in total, 767 subjects were screened for coronary heart disease (345 in the urban area and 422 in the rural area). The overall prevalence of Coronary heart disease was observed to be 12.0% (95%CI: 9.71-14.29). The prevalence of Coronary heart disease was observed to be significantly higher among urban study subjects (14.8%; 95%CI: 11.05-18.55) as compared to that among rural study subjects (9.7%; 95%CI: 6.88-12.52) (p<0.05). Out of the total 92 cases of Coronary heart disease in the present study, 51 (55.5%) cases were detected either on the basis of ECG tracing...
or Rose questionnaire, while rest of the 41(44.5%) cases were known cases of Coronary heart disease.

Discussion

The prevalence of Coronary heart disease in the present study was observed to be 14.8% in the urban area and 9.7% in the rural area. The prevalence of Coronary heart disease in the urban area in the present study was observed to be higher than that in the previous studies conducted in the urban populations by Gupta et al.6 in Jaipur in 1995 (7.5%) and in 2002 (7.3%), by Singh et al.3 in Moradabad (8.5%) and by Chadha et al.6 in Delhi (9.6%). The prevalence in the rural area in the present study was also observed to be higher than the earlier studies conducted in the rural populations by Singh et al.3 in Uttar Pradesh (3.0%), Gupta et al.6 in Rajasthan (3.5%), Wander et al.7 in Punjab (3.0%) and Gupta et al.6 in Himachal (5.0%). This increasing prevalence of Coronary heart disease in both urban and rural populations could be attributed to increasing longevity, urbanization and lifestyle changes.

The limitation of the present study is that it cannot project a representative picture of the problem of the whole country, considering the wide variation in socio-economic groups, dietary habits and cultural patterns existing in the country. The prevalence rates estimated in this study may be applied to urban and rural population of northern India, where socio-economic and other environmental factors are almost similar.

Conclusions and recommendations

The present study clearly indicates that CHD, as assessed by both clinical history and ECG criteria is of a magnitude that makes it a major problem of public health importance in India. The prevalence of coronary heart disease reported in the rural area is much higher in the present study than that reported in the earlier studies which necessitates that the IEC activities regarding the prevention of CHD should focus not only in the urban areas but also in the rural areas.

To obtain a composite picture for the whole country, community based epidemiological studies will have to be undertaken in the urban and rural areas of different parts of the country. Moreover, the diverse diagnostic criteria of coronary heart disease and the varying survey methods used in various studies make it difficult to estimate the trends of the disease over time, or the variations between different geographical regions. Thus, more periodically well organized systematic epidemiological studies with uniform methodology and uniform diagnostic criteria of CHD are required in various geographical areas of India to monitor trends of the disease over time and to study the variations between different areas.

Acknowledgements

I am thankful to the Department of Community Medicine, Maulana Azad Medical College, Delhi for providing me the help and support during the study.

Conflict of interest: NIL

References

A comparative study of random urine protein: creatinine ratio with 24 hour urine protein excretion in diabetic nephropathy

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Abstract
Renal involvement in diabetes is one of the long term complications leading to morbidity and premature mortality. Proteinuria is the most widely accepted clinical sign of diabetic nephropathy. The present study was a correlation between 24 hour urine protein excretion and random urine protein: creatinine ratio in diabetic subjects for predicting renal involvement. A total of 69 diabetic subjects were studied. It was observed that protein excretion in 24 hr sample correlated significantly with random urine protein: creatinine ratio in physiological and also in nephrotic range of proteinuria. This may have important clinical applications as single urine specimens, which can be collected easily in outpatient clinics and field studies, could replace the more traditional timed urine collections that have been used to assess the risk of clinical diabetic renal disease.

Key Words
Diabetic nephropathy, protein: creatinine ratio.

Introduction
Diabetic nephropathy is a devastating complication of diabetes mellitus and is among the leading indications for dialysis and kidney transplantation. It is the commonest cause of end stage renal disease. Occurrence of nephropathy is around 40% in type 1 and 25% in type 2 diabetes mellitus1.

Degree of proteinuria is the useful marker for renal involvement and response to treatment. Protein excretion rate is ordinarily determined from a 24-hour urine collection. Random specimens vary considerably in protein concentration. 24-hour urine sample collection could be inconvenient and cause frequent collection errors. Random urine sampling for protein:creatinine ratio would be more acceptable and less time consuming. The protein: creatinine ratio takes into account the fact that creatinine excretion remains fairly constant in presence of a stable Glomerular Filtration Rate. The protein excretion would also be fairly stable. Therefore the ratio of the two in a single voided sample would reflect the cumulative protein excretion over the day, as the two stable rates would cancel out the time factor2.

The protein:creatinine ratio in a single voided urine samples is an accurate, convenient, inexpensive and reliable estimate of total proteinuria in the vast majority of patients. Hence protein: creatinine ratio in single voided urine samples may be even more reproducible than 24 hours urinary protein excretion3. The protein: creatinine ratio of a randomly obtained urine specimen correlates with 24-hour urine protein in patients with type 1 diabetes and may be a useful tool in screening a patient for proteinuria or estimating range of proteinuria in patients4.

Methods
Total of 69 diabetics were selected. 12 were with 24 hours urinary protein levels > 3g/day, 39 with protein levels 0.2 – 3.0 g/day and 18 were with < 0.2g/day. Detailed medical history and relevant clinical examinations were carried out in these patients. Patients with chronic renal failure, glomerular nephritis due to other systemic conditions and hypertensives were excluded from the study. 24 hour urine was collected in the container having 4ml of 10% thymol in isopropanol as a preservative for 24 hours. This was thoroughly mixed and a sample of 2ml was taken for evaluation of proteins. Total volume was noted and calculation was done for 24 hours. A random urine sample of 5ml was collected on next day any time just before the analysis. The biochemical parameters were analyzed.

Results
Of the 69 diabetic subjects studied , 16 subjects with 24 hour urine protein < 0.2 g/day had P:C ratio of < 0.2, 37 subjects with 24 hour urine protein 0.2 – 3.0 g/day and 9 subjects with 24 hour urine protein > 3.0 g/day had P:C ratio of > 3.0 g/day. It is evident that correspondence of 24 hr urine protein and protein:creatinine ratio in diabetic subjects is highly significant (p<0.001).

Discussion
The reference method for estimating urine protein concentration is accurately timed 24-hour urine specimen which is considered to be difficult in some circumstances because of difficulties associated with obtaining a complete collection. An alternative approach
of estimating urine protein:creatinine ratio in random specimen can be considered. Protein:creatinine ratio in single voided urine samples correlates well with measurements of 24-hour urinary protein, this may be even more reproducible than 24-hour urinary protein excretion. More over a random urine sample in an outpatient tested usually represents renal excretion during two to four hours of varied activity and most of the daily urinary excretion of protein of an outpatient takes place during activity not recumbency. Correlation between random urine protein:creatinine ratio and 24 hour urine protein excretion was significant and strong protein:creatinine ratio is highly useful test in outpatient clinical setting but its precision and accuracy may be affected by patient's physical activity.

Urinary protein excretion is routinely expressed as the amount of protein excreted per unit of time per unit body surface area. Since creatinine production and excretion are also related to body size, it is possible that good correlation exist between protein:creatinine ratio and quantitative protein excretion in individual patients. It is noted in earlier studies and also in present study that patients who excreted > 3 gm protein in 24 hour had protein:creatinine ratio that exceed 3.0 which predicted nephrotic proteinuria. Similarly those patients who excreted < 200 mg protein in 24 hour had protein:creatinine ratio of < 0.2, which reflect insignificant or physiological protein excretion. Hence a highly significant correlation exist between 24-hour urinary protein concentration and protein:creatinine ratio levels in random urine sample.

By careful choice of cut-off, protein:creatinine ratio can be used in patients with kidney disease to rule out abnormal 24 hr loss of protein. Random samples can be used as surrogate for 24 hour sample.

**Conclusion**

Proteinuria is an important clinical sign of diabetic nephropathy. The definitive method for quantifying urinary protein excretion is 24-hour urine sample, but obtaining these samples is cumbersome, time consuming and inaccurate.

The present study suggests estimating protein:creatinine ratio in random urine sample for renal involvement in diabetic subjects provide a convenient method for early diagnosis and intervention of diabetic nephropathy. Hence the single voided test is simple, reliable and useful in screening, assessment and follow-up of diabetic nephropathy.

**References**


**Table 1: Correlation Between Protein: Creatinine Ratio In Random Urine Sample And 24 Hour Urinary Protein Concentration**

<table>
<thead>
<tr>
<th>Protein : creatinine ratio in random urine sample</th>
<th>24 hour urinary protein concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 0.2 (g/day)</td>
<td>&lt; 0.2 - 3.0 (g/day)</td>
</tr>
<tr>
<td>&lt; 0.2</td>
<td>16</td>
</tr>
<tr>
<td>0.2 - 3.0</td>
<td>2</td>
</tr>
<tr>
<td>&gt; 3.0</td>
<td>-</td>
</tr>
<tr>
<td>Total</td>
<td>18</td>
</tr>
</tbody>
</table>

X² = 92.0

p < 0.001 HS (highly significant)
Diagnostic imaging in implantology: From conventional to newer paradigms

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Abstract

Imaging is an useful adjunct for many dental procedures in general but is essential in the clinical practice of implantology. Apart from ruling out local bone and tooth pathology, imaging helps clinicians to determine bone quality and quantity and give a fair idea of implant orientation. Unlike endodontics, which can rely on conventional intra oral periapical assays (IOPA), imaging in implantology require definite cross sectional views to appreciate buccal and lingual bone plates and overall bone quality and quantity bucco-lingually. From conventional IOPAs and orthopantomograms (OPG) to cone beamed computed tomography (CBCT) and three dimensional implant planning software with computer aided designing and machining, the diagnostics in implantology has gone through a sea change. Without these advancements, certain concepts like flapless surgery and ideal positioning of the implant would have not seen the light of the day. Overall, the precision in patient and implant selection, implant placement, evaluation of the need for hard and soft tissue augmentation has been greatly enhanced with the help of these advancements.

This review series covers the conventional techniques and also deals with the advancements like multiplanar reformatted computed tomography and three dimensional imaging techniques as used in implantology today with clinical guidelines to adhere in the end.

Keywords

Imaging, IOPA, OPG, CBCT, three dimensional implant planning software.

Introduction

The influence of implants in clinical dentistry has been phenomenal. These are in a way unique as they provide a range of additional artificial abutments – a possibility which never existed before. The success of implants in providing optimum esthetics and function is unparallel by any other treatment modality in prosthodontics1,2,3.

In the past few years, the advances in science related to implant diagnosis and treatment planning has also been incredible. From conventional periapical films to three dimensional implant planning software with computer aided designing and machining, the diagnostics in implantology has gone through a sea change. Without these advancements, certain concepts like flapless surgery and ideal positioning of the implant would have not seen the light of the day. Overall, the precision in patient and implant selection, implant placement, evaluation of the need for hard and soft tissue augmentation has been greatly enhanced with the help of these advancements.

This review series covers the conventional techniques and also deals with the advancements like multiplanar reformatted computed tomography and three dimensional imaging techniques as used in implantology today with clinical guidelines to adhere in the end.

Fig. 1: The stepwise procedure from examination to placement.
implant cases as given by Rosenfeld and MecallS helping clinicians to place implants with clinical precision. With the help of diagnostic wax up, correct alignment of the replacing tooth can be finalised. Barium coated templates will help clinicians to determine the exact implant alignment and multiplanar reformatted computed tomography will help clinicians to determine the need for any hard or soft tissue augmentation. Although recommended, this protocol may not be possible in every implant case because of economic, time, space and availability concerns. Standard intraoral periapical assays (IOPA) and orthopantomograms (OPG) although do not provide cross sectional imaging but still are of significance as these can be used as screening and also for non complicated cases having abundant bone.

Various imaging modalities in implantology

a) Parallel IOPA:

Conventional IOPA are used mainly to rule out any local bone or dental disease but has limited value in determining bone quality and quantity. Normally, a no.2 size dental film captures 25 x 40 mm of view, hence certain landmarks like inferior alveolar canal and maxillary sinuses are beyond its view. In addition, there is difficulty in obtaining an undistorted film in areas of shallow palate and reduced sulcus depth. Long cone paralleling technique limits distortion and magnification to 10%. Low cost and low radiation dosage is an advantage. Recommended in:

1) Single tooth implants with abundant bone width.
2) Estimating crestal bone loss over a period of time. However, it’s to be noted that vertical bitewing intraoral x ray is better than IOPA in estimating crestal bone loss
3) To check fit of the abutment and sulcus formers.
4) Diagnosis of a failing implant.

b) Digital IOPA:

The charged couple device systems use rigid sensors placed intraorally to capture the image. When radiograph is made, the remnant x ray beam exiting the patient is captured to scintillator coating a silicon chip. The resulting image flashes at the monitor in seconds of exposure and hence allows operator to make necessary adjustments if required during surgery. Advantages include better sharpness and contrast and less exposure than conventional IOPAs.

In a study comparing diagnostic potential of direct digital and conventional intraoral radiography in the evaluation of perimplant conditions, it was observed that there is no statistically significant difference between the two. However, there was a tendency of more consistent evaluation in digital radiographs. Also, the speed of delivery of view and low radiation exposure make the digital IOPA the suitable option in today’s scenario.

In a study on imaging of peri-implant bone levels of implant with buccal bone defects, it was concluded that periapical assay does not provide valid data on the circumferential bone level in implants with buccal bone defects but in general reflects the bone level on the lingual side, hence overestimating the bone anchorage of these implants.

c) Lateral cephalogram:

Demonstrates geometric skeletal anatomy, interarch relationship, anterior crown to implant ratio and clear view of the sinuses. However as only the median area is shown, it has got limited use in implant placement.

d) Oblique lateral cephalogram:

In extreme mandibular atrophy it can be difficult to obtain intraoral radiographs of adequate diagnostic quality. Extraoral oblique lateral cephalometric radiographs (OLCR) can then be an alternative and also reproducible images of large parts of mandible can be obtained. In vitro, the results of densitometry using periapical films and OLCR were shown to be similar.

Tomography:

Tomography otherwise known as body section radiography, planigraphy, laminography or stratigraphy

Fig. 2: IOPA showing crestal bone loss

Fig. 3: Oblique lateral cephalometric radiograph
is a process of using motion of a x ray source and film in generating images in one plane at a time.

It’s basically of two types i.e linear and complex. Complex again can be divided into many types like circular, spiral, elliptical, figure-8, trispiral and hypocycloidal.

Each of these motions has advantages regarding the way in which out of plane structures are blurred. For example, a linear structure which is aligned with the linear motion of a linear tomograph, will not appear blurred, except at the ends, whereas such a structure will be blurred by the circular motion of a circular tomograph.

e) Pantomography:

It is a special tomography technique where panoramic roentgenograms of curved surfaces are obtained by rotating the X ray tube and film-screen holder around the patient.

It’s the most widely used imaging modality but not the most diagnostic. It’s considered as a screening radiograph which provides a good overall picture, but are subject to distortion and magnification. Vertical magnification is 10% while horizontal is 20% and also variable.

Uses:
1) Opposing landmarks are easily identified
2) Vertical bone height can be assessed
3) Procedure performed with great ease and speed
4) Gross anatomy and related pathology can be assessed

A,B- from alveolar crest to nasal fossa in the maxillary anteriors.

C,D- from alveolar crest to maxillary sinus floor in maxillary posteriors.

E- from alveolar crest to inferior alveolar canal in mandibular posteriors.

F- from alveolar crest to lower border of the mandible in mandibular anteriors.

G- from alveolar crest to mental foramen in mandibular premolar region.

Disadvantages:
1) Does not demonstrate bone quality
2) Misleading due to magnification
3) Little use in depicting the spatial relation
4) Least useful in deciding angulation.
5) Overlapping in maxillary anterior region.

To reduce errors and to aid in orientation, stents are often used. These may be copies of a denture incorporating steel balls of known dimension, twists of wire or vaccum moulded splints painted with radiopaque material.

The formula for calculating the bone height is as follows:

\[
\text{Radiographic bone height} = \frac{\text{Radiographic dimension of the metal ball}}{\text{Actual dimension of the metal ball}}
\]

In a study relating to the effectiveness of an OPG in clinical placement of mandibular implants, it was found that there was no permanent sensory disturbances of the inferior alveolar nerve in any of the cases and there were only two cases of postoperative paresthesia, representing 2/2584 (0.08%) of implants inserted in the posterior
segment of the mandible or 2/1527 (0.13%) of patients. These sensory disturbances were minor, lasted for 3 and 6 weeks and resolved spontaneously11.

In a study to know the reliability of crown root ratio, linear and angular measurements on panoramic radiographs, it was observed that these parameters can be taken of the same patient at different times with consistent accuracy12.

f) Transverse saggital section:
The cross sectional imaging in implantology is required essentially so as to visualize buccal and lingual bone morphology which is otherwise not possible in traditional imaging modalities previously discussed.

However, the structures are not very clear and also limited by distortion, the exact measurements can only be made after magnification correction. But with limited cost, this transverse section has gone along way in redefining clinical implantology as it gives a view of buccal and lingual walls which were never seen before.

g) Computed tomography:
Invented by sir Hounsefield in 1972, it has revolutionized imaging in implant dentistry. It’s a digital and mathematical imaging technique which reduces blurring to a great extent. In this modality axial images are produced and can view both hard and soft tissues. This treatment modality fulfills all the five objectives of implant imaging. The density of bone and soft tissues is categorized on the basis of Hounsfield units.

<table>
<thead>
<tr>
<th>Density</th>
<th>Hounsfield units</th>
</tr>
</thead>
<tbody>
<tr>
<td>D1</td>
<td>≥ 1250</td>
</tr>
<tr>
<td>D2</td>
<td>850-1250</td>
</tr>
<tr>
<td>D3</td>
<td>350-850</td>
</tr>
<tr>
<td>D4</td>
<td>150-350</td>
</tr>
<tr>
<td>D5</td>
<td>&lt;150</td>
</tr>
</tbody>
</table>

Major disadvantage of CT is of metal artifacts from tooth filling was overcome by use of axial plane and hence left the image of the jaw bone undistorted. Device is capable of acquiring thin slices of 1.5mm or less. Conventional spiral tomography plays an important role in pre-surgical treatment planning, increasing clinician’s certainty of the need of additional surgical procedures (bone grafting, sinus lifting, and others) in pre-surgical treatment stage13.

h) Cone Beam CT:
The cone-beam technique involves a single 360° scan in which the x-ray source and a reciprocating area detector synchronously move around the patient’s head, which is stabilized with a head holder14.

Advantages of CBCT over CT:
- Rapid scan time.
- Display modes unique to maxillofacial imaging,
- Reduced image artifact
- X-ray beam limitation.
- Cost of equipment is approximately 3-5 times less than traditional conventional CT.
- The equipment is substantially lighter and smaller,
- No special electrical requirements needed,
- No floor strengthening required,
- The room does not need to be cooled,
- Little technician training is required,
- Feasible choice for Claustrophobic individuals.
- The upright position is also thought by many to provide a more realistic picture of condylar positions.
during a TMJ examination,
• Both jaws can be imaged at the same time (depending on the specific cone beam machine),
• Radiation dose is considerably less than with a conventional CT.

i) Three dimensional Implant planning software:
First preoperative implant planning software was developed in 1993 in Columbia under the name Simplant. These kind of softwares have revolutionized imaging and preoperative planning in implantology as choosing the exact length and diameter of the implant, exact site of placement and knowing the exact density of the bone have become increasingly easy. As the software provides three dimensional imaging it is even possible to do a virtual mock surgery. Dentascan and Implan are the other similar softwares available. These softwares also help us in fabrication of CT based surgical guides which are made on the precise measurements obtained by a three dimensional scan and hence precise placement of the implant. Nobleguide and M guide are two such examples.

Dosages in various imaging modalities

<table>
<thead>
<tr>
<th>Imaging Modality</th>
<th>Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Full mouth IOPA</td>
<td>13–100 µSv</td>
</tr>
<tr>
<td>OPG</td>
<td>10 µSv</td>
</tr>
<tr>
<td>CT</td>
<td>1,320–3,324 µSv</td>
</tr>
<tr>
<td>CBCT</td>
<td>36.9–50.3 µSv</td>
</tr>
</tbody>
</table>

Guidelines and conclusion
The conventional two dimensional imaging modalities discussed in this article may not provide an exact overall picture as computed tomography and other three dimensional implant planning softwares but are very useful, handy, economic and efficient provided magnification is corrected. These can be used in relatively simple cases and in patients having abundant bone. The advancements like multiplanar reformatted computed tomography and three dimensional imaging planning softwares are essential in cases where hard and soft tissue augmentation is needed and multiple implants are planned. Cone beam computed tomography has become a standard of care owing to its advantages of being economical, in office design and image accuracy.

References
Abstract

Background
Periodontal dressing are advocated post operatively to afford wound healing and prevent patient discomfort. And these materials are not readily modified by addition of antibiotics or antiseptics to provide antibacterial properties.

Material and Methods
Subjects for this study consisted of 21 patients of both sexes in the age group of 20 to 40 years, requiring periodontal therapy.

Results
The present study showed that there was marked reduction in plaque index, gingival index, gingival exudates measurement and the degree of inflammatory cells on Chlorhexidine treated side compared to the control side.

Summary and Conclusion
Further longitudinal experimental studies are necessary to delve into the usefulness of Chlorhexidine gluconate as a post-operative surgical dressing.

Key Words
Periodontal Dressing, Chlorhexidine, Flap surgery, Gingival index, Gingival exudates.

Introduction
Periodontal dressing are used following surgery with the aim of providing comfort to the patient during wound healing. However, relatively few dressing have any well documented evidence to substantiate their efficacy. With the advent of Chlorhexidine gluconate and its proven effectiveness as an anti-bacterial agent much interest has been shown in its use for post-operative care. Currently, Chlorhexidine gluconate is being used either in the form of topical application or mouth wash in gingival inflammatory condition. Jorgensen et al. and Addy et al. have reported better healing after periodontal surgery with the use of periodontal dressing incorporating Chlorhexidine and have attributed this to anti-bacterial property of Chlorhexidine.

However, no study so far has been undertaken to determine the efficacy of Chlorhexidine as a plaque inhibiting agent to promote better gingival healing after periodontal surgery.

Therefore, a clinical and histological study has been planned with the following objectives.

1. To comparatively evaluate the effect of topical Chlorhexidine and conventional eugenol containing periodontal dressing on gingival healing after flap surgery.
2. To throw more light on the possible role of Chlorhexidine as plaque inhibiting agent in post surgical wound care.

Method and Materials
Subjects for this study consisted of 21 patients of both sexes in the age group of 20 to 40 years, who visited the department of periodontics, requiring periodontal therapy. Subjects with periodontal pockets of 4 to 6mm were included in the present study.

The clinically parameters included were, quantitative measurement of gingival exudates, histological examination of gingiva, plaque index system of Silness and Loe for oral hygiene status, gingival index of Loe and Silness and the pocket depth were measured by using William's periodontal probe.

Full thickness flap surgery was chosen as the treatment of choice to eradicate periodontal pocket in the selected subjects. After the surgical procedure the wound was covered with a thin film of 2% Chlorhexidine gluconate as a post operative dressing with the help of a swab on one half of the operated site and conventional eugenol containing pack was placed on the other half as a control. Sutures were removed on the 8th day and information regarding any notable difference in pain, discomfort, sensitivity to hot and cold, between experimental and control side was recorded.
Quantitative measurement of gingival exudates

Gingival exudates measurement was performed on the mid-labial aspects of gingiva in all the three of both experimental and controlled teeth separately. The exudates measurements were made using modified intracrevicular method of Loe and Holm Pederson. After the gingiva and surrounding tissue were carefully dried with air, strips of filter paper Whatman No.3 of 1.5mm in width and several millimeters in length were gently placed parallel to the long axis of the tooth at the entrance of the gingival crevice avoiding physical irritation to the crevicular epithelium. The strips remained in place for 3 minutes (Fig 1). They were then removed, dried and immediately stained with an alcoholic solution of 0.2 % ninhydrin Brill7 (Fig 2). Only the area which absorbed crevicular fluid was then measured with travelling microscope and mean scores were calculated.

Histopathological examination of gingiva to assess the degree of inflammation by taking biopsies

Histopathological examination of gingiva was done by taking biopsy from identical sites that is between 21 and 21 in both experimental and control sides. Biopsies obtained were immediately transferred into bottled containing 10% formalin and allowed to fix for 24 hours. Later, they were embedded in paraffin blocks and sections of 5 microns thickness were cut with the help of a rotating microtome. The sections were stained with hematoxyline and Eosin and mounted with a cover slip by using Canada balsm. The tissue sections were examined under a microscope. The density of inflammatory cells infiltration was assessed at 100 x magnification according to the following criteria.

O – No inflammatory cells in the connective tissue and epithelium.
1 – Sparse distribution of inflammatory cells.
2 – Moderately dense accumulation of inflammatory cells in isolated areas, sparse distribution in other areas.
3 – Dense aggregation of inflammatory cells throughout the subepithelial connective tissue.

Oral hygiene, gingival status and quantity of gingival exudates were periodically assessed post-operatively on 15th,22nd and 29th day. However; gingival biopsy was repeated, only on 29th day, from the area between 32 and 23.

Statistical Analysis

The mean, standard deviation of oral hygiene index, gingival index and gingival exudates measurement of different post-operative intervals were then calculated and correlated with clinical and histopathological evidence of degree of inflammation for both experimental and controlled sides. The different parameters of experimental sites were then compared with that of the controlled sites. Student’s test was applied to find out the statistical experimental and control areas both clinically and histopathologically.

Results and Discussion

Periodontal dressings are advocated post-operatively to afford wound protection and prevent discomfort. But there is still lack of agreement as to whether these dressings have any specific effect on plaque control and enhance better gingival healing. However, no study so far has been undertaken to determine the efficacy topical Chlorhexidine as a plaque inhibiting agent to promote better gingival healing after periodontal surgery. There is also a lacuna in the information on the histopathological evidence to confirm the efficacy of Chlorhexidine in promoting better gingival healing. Therefore a comparative clinic-histopathological study was undertaken to find out the role of topical Chlorhexidine gluconate on wound healing after flap surgery in comparison with conventional eugenol containing periodontal dressing.

The results of the present study are tabulated in the tables I to V.

In the present study most of the patients had less or no discomfort and compared to that of control side. This is in agreement with the finding of Jorgensen et al, Newman et al and Addy and Dolby. However, during the rest of the post-operative period there was no noticeable difference between both the sides.
Plaque Index

The mean and standard deviation of plaque index was less on the side where Chlorhexidine was topically applied compared to the control side with the conventional periodontal dressing on the 8th post-operative day. This again confirms the earlier reports by Loe and Schiot, Lindhe et al, Flotra et al, Hoyos et al, and Sally et al, who have observed a reduction in mean plaque index with the use of Chlorhexidine as a plaque inhibiting agent either in the form of mouth wash or as a topical application or being incorporated into the gel. The mean plaque index during the rest of the post-operative period was same on both sides.

Gingival Exudate:

The gingival exudates measurement was less on the experimental side on 8th and 15th post-operative days compared to the control side. This confirms with the above finding of Lindhe et al and Jorgensen et al. However, there was no statistically significant difference on both sides during the rest of the post-operative period.

Gingival Index

A statistically significant reduction in gingival index was noted on chlorhexidine treated side. The above finding are in agreement with the earlier findings of Hoyos et al, Loe and Schiot, Jorgensen et al and Newman et al. Hoyos et al has reported a marked reduction in plaque index and gingival index scores with the use of coronosyl dental gel in 56 children after one month with regular oral hygiene methods.

Jorgensen et al also observed less gingival exudates and decreased tendency to bleeding with the use of Chlorhexidine containing surgical dressing during healing of reverse beveled flap operations.

Histopathological measurements

The distribution of inflammatory cells were minimal.

Table I: Subjective Symptoms.

<table>
<thead>
<tr>
<th>No of patients</th>
<th>Period of study</th>
<th>Groups</th>
<th>Number of patients with different subjective symptoms</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Pain</td>
</tr>
<tr>
<td>21</td>
<td>8th day</td>
<td>Control</td>
<td>13</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Expt</td>
<td>5</td>
</tr>
<tr>
<td>21</td>
<td>15,22 &amp; 29 days</td>
<td>Control</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Expt</td>
<td>-</td>
</tr>
</tbody>
</table>

Table II: Plaque index.

<table>
<thead>
<tr>
<th>No of Patients</th>
<th>Period of Study</th>
<th>Groups</th>
<th>Mean and Standard Deviation</th>
<th>t-test</th>
<th>Statistical significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>21</td>
<td>Pre-surgical</td>
<td>Control</td>
<td>2.00±0.11</td>
<td>0.06</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>1.96±0.17</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>8th day</td>
<td>Control</td>
<td>0.66±0.07</td>
<td>7.59</td>
<td>P &gt; 0.001</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.30±0.03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>15th day</td>
<td>Control</td>
<td>0.46±0.05</td>
<td>1.91</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.32±0.05</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>22nd day</td>
<td>Control</td>
<td>0.33±0.05</td>
<td>1.58</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.30±0.05</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>29th day</td>
<td>Control</td>
<td>0.28±0.05</td>
<td>0.41</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.25±0.05</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table III: Gingival exudates measurement.

<table>
<thead>
<tr>
<th>No of Patients</th>
<th>Period of Study</th>
<th>Groups</th>
<th>Mean and Standard Deviation</th>
<th>t-test</th>
<th>Statistical significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>21</td>
<td>8th day</td>
<td>Control</td>
<td>2.34±0.09</td>
<td>1.92</td>
<td>P &gt; 0.01</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>1.97±0.09</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>15th day</td>
<td>Control</td>
<td>1.4±0.07</td>
<td>2.12</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>1.19±0.07</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>22nd day</td>
<td>Control</td>
<td>0.73±0.05</td>
<td>1.41</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.70±0.05</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>29th day</td>
<td>Control</td>
<td>0.39±0.05</td>
<td>0.16</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.40±0.04</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
or sparse in most of the patients on the side with Chlorhexidine dressing on 8th post-operative day. However, on the 29th post-operative day the degree of inflammation was very minimal and almost the same on both the sides (Fig 3,4,5,6). The above finding are in agreement with the reports of Robert et al15 who demonstrated microscopically that 2% Chlorhexidine gluconate applied topically each day to the teeth and gingival margin of dogs, resulted in significant reduction in inflammation in healing gingival wound.

The difference in the different parameters noted on both experimental and the control sides indicate that the use of 2% Chlorhexidine gluconate topical application as a post-operative dressing on the experimental side showed better healing of gingival wounds after periodontal surgery. Davies et al16, Lindhe et al11 and Schiot17 have shown that topical application of 2% chlorhexidine gluconate on teeth can completely inhibit the plaque formation. Hence from the present study it can be concluded that the beneficial effect of Chlorhexidine on wound healing is due to its potent antibacterial activity as well as its non-irritating character.

From the above discussion it may be inferred that the decrease in the gingival index and the gingival exudates measurement on the experimental side in the present study may be due to antibacterial and plaque inhibiting effect of chlorhexidine. The clinical-histopathological finding during the initial healing phase following the flap surgery shows that a better results was noticed on the experimental side where topical 2% Chlorhexidine gluconate was used compared to the control side where a conventional eugenol containing dressing was given.

**Summary and Conclusion**

The present study showed that there was marked reduction in plaque index, gingival index, gingival exudates and the degree of inflammatory exudates on the experimental side, whereas the control side showed no significant change. This indicates that chlorhexidine gluconate has a significant role in the healing of gingival wounds following periodontal surgery.

**Table IV:** Gingival Index.

<table>
<thead>
<tr>
<th>No of Patients</th>
<th>Period of Study</th>
<th>Groups</th>
<th>Mean and Standard Deviation</th>
<th>t-test</th>
<th>Statistical significance</th>
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</thead>
<tbody>
<tr>
<td>21</td>
<td>Pre-surgical</td>
<td>Control</td>
<td>2.06±0.10</td>
<td>0.07</td>
<td>P &gt;0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>2.05±0.10</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>8th day</td>
<td>Control</td>
<td>0.82±0.09</td>
<td>4.95</td>
<td>P &gt; 0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.35±0.03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>15th day</td>
<td>Control</td>
<td>0.16±0.02</td>
<td>1.52</td>
<td>P&gt;0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>0.12±0.02</td>
<td></td>
<td></td>
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<tr>
<td>21</td>
<td>22nd day</td>
<td>Control</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>21</td>
<td>29th day</td>
<td>Control</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>-</td>
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</tbody>
</table>

**Table V:** Histopathological Assessment of degree of Inflammation.

<table>
<thead>
<tr>
<th>No of Patients</th>
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<th>Groups</th>
<th>Number of patients with different grading</th>
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<td></td>
<td></td>
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<td>Grade 0</td>
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<tr>
<td>21</td>
<td>8th day</td>
<td>Control</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>8</td>
</tr>
<tr>
<td>21</td>
<td>29th day</td>
<td>Control</td>
<td>8</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Experimental</td>
<td>10</td>
</tr>
</tbody>
</table>

Table V shows histopathological assessment of degree of inflammation in 21 patients on 8th and 29th post-operative periods. The distributions of inflammatory cells were minimal or sparse in most of the patients on the side with Chlorhexidine dressing on 8th post-operative day. However, on the 29th post-operative day the degree of inflammation was very minimal and almost the same on both sides.

**Figure 3:** Photomicrograph of gingiva showing minimal inflammation in the underlying epithelium and connective tissue on 8th post-operative day (expt side). H & E stain in low power.

**Figure 4:** Photomicrograph of gingiva showing moderate inflammation in the underlying epithelium and connective tissue on 8th post-operative day (control side). H & E stain in low power.
cells on Chlorhexidine treated side compared to the control side where conventional eugenol containing periodontal dressing was given.

Results obtained in the present study shows at the end of first post-operative week there was statically significant difference in different parameters on Chlorhexidine treated side. During the observation period, the gingiva which had Chlorhexidine dressing showed less gingival exudates decreased bleeding tendency and less inflammatory cells indicating of that topical application of 2% Chlorhexidine gluconate as a post operative dressing increased the healing rate.

However, further longitudinal experimental studies are necessary to delve into the usefulness of Chlorhexidine gluconate as a post-operative surgical dressing.

References

Effect of two tongue cleaning methods on oral mutans Streptococci level

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Abstract

Background
Tongue scraping and brushing have been appreciated for hundreds of years but are still appreciated or used by the public. Scientific evidence has validated the need to practice habitual and tongue cleaning as part of daily home oral hygiene procedures.

Objective
To assess and compare the effect of tongue scraping and tongue brushing on oral Mutans streptococci level.

Methods
20 healthy subjects aged 14 to 15 years were randomly selected. Flat plastic tongue scraper and Nylon multitufted small headed tooth brush are the two tongue cleaning devise used. Unstimulated salivary samples were obtained at intervals from each individual. Salivary samples were inoculated on Mitis Salivary Agar Plate and Sorbital Broth was used for identification of Mutans streptococci group.

Results
Paired and unpaired ‘t’ test were employed. Reduction in the Mutans streptococci level from 48.4X10⁴ CFU and 38.3X10⁴ CFU at baseline in tongue scraping and tongue brushing group respectively to 0.34X10⁴ and 0.39X10⁴ CFU after 7th day. Conclusion: Both tongue coating removal methods evaluated were efficient in reducing Mutans streptococci level. This implies that physical removal of the coating on the dorsum of the tongue is important and not the method used for the same.

Key Words
Tongue scraping, Tongue brushing, Tongue cleaning, Mutans streptococci.

Introduction
The most common concept concerning individual health is the harmony of one’s physical, mental, and social well being. The simple absence of disease is not accepted as an indication of health. Tongue cleaning being an ancient habit, is practiced for centuries in many Eastern and Oriental cultures, though not very popular in the Western Civilizations¹.

Recent literature has shown that tongue cleaning leads to healthy oral Environment². Tongue is a small but powerful organ of the body since it performs the function of -Taste, speech, mastication and deglutition. Hence the need for tongue cleaning has become a part of daily oral hygiene. Gilmore and Bhaskar presented convincing evidence that plaque forming streptococci counts increased ten folds after a week of not brushing the tongue².

Tongue Care in Antiquity
Although largely an unknown practice in the west, tongue brushing and scraping have been used since antiquity and are still used by natives of Africa, Arabia and India. In the early civilizations, oral cleansing often had religious ritual significance. The Hindus regarded ‘mouth as the gateway of the body, therefore it was necessary to keep it scrupulously clean’. The ancient Hindus used tongue scrapers with sharp curved edges made of Gold, Silver, Ivory or Tin¹.

The Mohammedans used Siwak wood brush once a day in a manner specified in the Koran. Prophet Mohammed said ‘you shall clean your tongue for that is the way to praise God’ and in Mohammedans the final stage of oral cleansing involved vigorous tongue brushing¹.

From 15th-19th century, tongue cleaning was known to be practiced primarily by the affluent leisure class. More recently, during the 20th century, tongue cleaning was not a popular concept, and only a few references are mentioned in the literature¹.

Why Clean the Tongue
Tongue, because of its surface texture contributes significantly in plaque formation and accumulation, has remained a neglected part in the oral cavity⁴.

The dorsal posterior part of most tongues has a coating of millions of microorganisms. Studies have shown that Dorsum of the tongue is an important reservoir for Mutans streptococci².
Further more, studies have also found a significant correlation between the prevalence of Mutans streptococci in saliva and its prevalence on the dorsum of the tongue

Mutans streptococci (MS) are one of the most virulent cariogenic pathogens in the oral cavity. Tooth brushing alone is effective in reducing bacterial counts in the mouth, but not dramatically. Tongue cleaning seems to have a more dramatic effect on the salivary levels of caries-causing bacteria, Such as Mutans streptococci5. With tongue scraping becoming established as an excellent tool for reducing the levels of Mutans streptococci in the oral cavity, it would be of great interest to compare the efficacy to tongue brushing method for decreasing the bacterial count in the oral cavity.

In this regard the effect of mechanical oral hygiene techniques on the levels of microorganisms, especially Mutans streptococci, is of great interest to dentists focused preventive care.

**Objectives of the Study**

1. To assess the effect of tongue scraping and tongue brushing on oral Mutans streptococci level.
2. To compare the two methods of tongue cleaning on the reduction of oral Mutans streptococci level.

**Material and Methods**

Study design: Double blinded Randomized Controlled Trial

Study population: 20 healthy subjects with similar food habits aged 14-15 years were selected and randomly distributed into two groups:

Group A-10 subjects, Group B-10 subjects.

Following tongue cleaning devices had been selected.

- Flat plastic tongue scraper - group A subjects.
- Nylon multitufted small headed toothbrush - group B subjects.

Inclusion criteria:

- Subjects with permanent dentition were included.
- All levels of Oral hygiene and dental caries were accepted those who had either rampant tooth decay or very poor oral hygiene were also included in this study this was important to see if the protocol was effective for all ranges of oral hygiene or not.

Exclusion criteria:

- Subjects suffering from Tonsillitis
- Subjects with any contributing medical history
- Subjects who have performed any type of tongue cleaning habits.

**Method of collection of salivary samples**

Unstimulated salivary samples were collected in the test tube by spitting method from all subjects prior to start of the experiment to establish base line Mutans streptococci level, after routine tooth brushing.

Demonstration to perform tongue scraping and tongue brushing was given to both the groups by a single examiner. Subjects were told to clean the tongue every morning after routine tooth brushing for 7 days.

Group A, involving 10 subjects, were given a tongue scraper and asked to scrape the dorsum of the tongue along the linea mediana and at each lateral part of the tongue once every morning after routine tooth brushing, every day for 7 days. Group B, consisted of 10 subjects were asked to brush the tongue with forward and backward strokes along the linea mediana and at each lateral borders of the tongue once every morning after routine tooth brushing, every day for 7 days.

Upon scraping or brushing the tongue, the patients were asked to spit out the excess saliva that had accumulated on the tongue. Then unstimulated salivary samples were obtained at 1 hour, 3rd day, and 7th day after the start of the experiment. A total of 4 samples were collected from each individual. During the entire study participants continued their habitual oral hygiene and were instructed not to take any antibiotics without prior information.

One ml of salivary samples were collected in the test tube and transported through Thyroglycolate broth media. 10 micro liters of saliva from each sample was inoculated on Mitis Salivary Agar Plate and Incubation was done at 37°C in 5-10% carbon dioxide for 48hrs. Sorbital broth was used for identification of Mutans streptococci group.

**Ethical Clearance and informed consent:**

Before staring the study, ethical clearance was obtained from ethical review committee board of K.L.E’s Institute of dental science, Belgaum, Karnataka state India. Oral consent was obtained from all the children and written informed consent was obtained from the parents.

**Statistical Analysis**

Statistical test employed for the obtained data in our study were Paired and unpaired ‘t’ tests. There were 20 participants in this research study. Paired ‘t’ test was used to compare within the group at different intervals and unpaired ‘t’ test was used to compare between the two groups.

**Results**

Reduction in the Mutans streptococci level from 48.4X104 colony forming units (C.F.U) and 38.3X104 C.F.U at baseline in tongue scraping and tongue brushing group respectively to 0.34X104 and 0.39X104 C.F.U after 7th day (p<0.01).

Graph I show the comparisons of mean number of Mutans streptococci in tongue scraping and tongue brushing groups at baseline. Mean number of Mutans streptococci
in tongue scraping group was 48.8 X 10^4 and in tongue brushing group it was 38.3X10^4 colonies forming units. The results demonstrate that there were no significant differences between group means at base line. This simply states that the groups were statistically equivalent before the start of the treatment. Base line measurements were compared to ascertain if there were any differences among the groups before the start of the treatment.

Graph II shows the mean decrease in Mutans streptococci level in tongue scraping group at different time intervals. When baseline value was compared with 1 hour, third day and seventh day value (p<0.01) and 1 hour value with 7th day value statistically highly significant difference was found. But when 1 hour value was compared with 3rd day value and 3rd day value with 7th day value no significant difference was found.

Graph III shows the mean decrease in Mutans streptococci level in tongue brushing group at different time intervals. When baseline value was compared with 1 hour, third day and seventh day value (p<0.01) and 1 hour value with 7th day value statistically highly significant difference was found. Only when 3rd day value was compared with 7th day value no significant difference was found.

Graph IV shows the gradual reduction of microorganisms from baseline to one hour, 3rd day and 7th day (p<0.01) but when it was compared between the groups at the final stage (after 7th day) no significant difference was found.

Discussion

The data obtained from this research project showed very clear trends and highly significant results. In the present study, there was a gradual reduction in the Mutans streptococci count from base line to 7th day after tongue scraping which is in close agreement with the study conducted by Almas.et.al2, White GE.et.al5 and Bordas A.et.al7. Since this method is performed every day there will be a gradual reduction of microorganisms from the tongue coating. Hence by 7th day the count reduced significantly. Bordas A. et.al7 reported that while mechanical tongue cleaning with or without chemical intervention can reduce bacterial load on the tongue, this effect is transient, and regular tongue cleaning is required to provide a long lasting (overnight) reduction in bacterial numbers. Nevertheless, tongue cleaning is an oral hygiene procedure that is little practiced due to discomfort and/or lack of awareness on the part of dental professionals and their patients.

In a study conducted by Quirynen. M.et.al8 no significant reduction in bacterial load was found when using toothbrush or scraper to clean the tongue. This may be because of different methods employed in collection of microbial sample from the tongue and analysis of non specific bacteria, where as in the present study, only Mutans streptococci level has been evaluated and saliva samples were collected by spitting method.

Hence this study is unique, being the first to compare two different methods of tongue cleaning and evaluating its effects on the reduction of specific microorganisms. Thus the clinical significance of this study should not be overlooked as research has proved the need to include the tongue in all oral hygiene measures. Thorough preventive measures need to include an effective means of reducing the pool of Mutans streptococci inhabiting the dorsum
of the tongue if one is to truly expect a striking reduction in caries.

**Conclusion**

In summary, both tongue coating removal methods evaluated were efficient in reducing Mutans streptococci level. This implies that physical removal of the coating on the dorsum of the tongue is important and not the method used for the same. On the basis of literature there appears to be enough data to justify the necessity to clean the tongue on a regular basis and as part of daily home oral hygiene practice.

Tongue cleaning is simple, fast and the benefits for most people far out weigh the small investment and time required to accomplishing this procedure. Therefore oral hygiene measures should include the dorsum of the tongue, especially in high-risk patients, who have endogenously high levels of Mutans streptococci residing in the oral cavity.

**References**

A study of oxidative stress and altered endothelial cell function in preeclampsia

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Abstract
Preeclampsia is a complex multisystem disorder characterized by hypertension and proteinuria. It is one of the most common and potentially fatal complications of pregnancy. A case control study was carried out to assess the levels of homocysteine, lipid peroxidation and antioxidant status in patients with preeclampsia. Fasting venous samples were collected during antepartum period and serum levels of homocysteine (Hcy), malondialdehyde (MDA), ascorbic acid (vitamin C) and uric acid were measured. In the preeclamptic group, Hcy and MDA levels were significantly raised while antioxidant ascorbic acid level was significantly reduced (p < 0.01) and uric acid concentration was increased significantly (p <0.01). These findings suggest that Hcy and lipid peroxidation are associated with preeclampsia. In preeclampsia, antioxidants are extensively utilized to counteract the cellular changes and endothelial dysfunction mediated by oxidative stress. Placental oxidative stress which results from the ischemic reperfusion injury is reported to be involved in the etiopathogenesis of preeclampsia.

Keywords
Preeclampsia, Homocysteine, lipid peroxidation, antioxidants.

Introduction
Preeclampsia is a pregnancy specific disorder which complicates 7-10% of all gestation1. This common disorder, which is more prevalent in first pregnancies, is associated with approximately 10-15% of maternal and fetal morbidity and mortality, with serious outcomes occurring in developing countries2.

Preeclampsia is a triad of oedema, hypertension and proteinuria occurring primarily after the 20th gestational week and most frequently near term3.

It is associated with defective placentation, in which the dislodging of extravillous trophoblast plugs in the maternal spiral arteries. This leads to the onset of blood flow into the intervillous space, causing an oxidative burst that generates reactive oxygen species (ROS)4. Also, the reduced uteroplacental perfusion due to the aberrant placentation leads to ischemic reperfusion injury to the placenta5.

Preeclampsia is characterized as a state of oxidative stress resulting from increased generation of free radicals and decreased levels of antioxidants6.

The free radicals thus, generated are capable of exerting mediating processes such as tissue remodeling, hormone signaling oocyte maturation, folliculogenesis, tubal function, ovarian steroidogenesis and germ cell function. Its increased levels can inflict significant damage to cell structure7.

Oxidative stress has been proposed as a promoter of endothelial cell dysfunction and lipid peroxidation8 which results in production of lipid peroxidation products i.e. malondialdehyde (MDA) and lipid hydroperoxides9. Homocysteine is a sulfur containing amino acid derived from the metabolic demethylation of methionine10.

Homocysteine by autooxidation generates superoxide and hydrogen peroxide, both of which damage the arterial endothelial lining, causing endothelial dysfunction11.

Recently, homocysteine has been observed as an independent risk factor for vascular endothelial cell injury in cardiovascular factor disease and common obstetric problems i.e. preeclampsia12.

It is observed that oxidative stress leads to focal collagen damage in the fetal membranes and results in preterm labour13.

Significantly decrease levels of antioxidants i.e. vitamin C, vitamin E and uric acid have been reported in patients with preeclampsia14.

In this context, the present study has been undertaken to determine the changes in serum levels of total homocysteine, lipid peroxidation product, MDA and antioxidant levels with vitamin C and uric acid.

Material and Methods
A case control study was conducted in Department of Biochemistry, Santosh Medical College and Hospital, Ghaziabad. Fifty severe eclamptic patients (criteria of severe preeclampsia: systolic blood pressure of 160 mmHg or more and diastolic blood pressure of 110 mmHg or more, persistent proteinuria of at least 2+ by dipstick or 24 hour urinary excretion of 2 g or more, persistent headache, oliguria, nausea, vomiting, epigastric pain, pulmonary oedema, thrombocytopenia) and fifty normotensive healthy pregnant controls.
Subjects free from pre-existing hypertension, cardiovascular disease, cerebrovascular disease, diabetes mellitus, renal disease, liver disease and hypothyroidism were included in the study.

Study subjects of both groups were matched with respect to maternal age, gestational age and BMI. Sociodemographic features of subjects are given in Table I.

Ethical clearance for the study was taken from the concerned authorities. Informed written consent was taken from the subjects.

With all aseptic precautions morning blood samples from all the study subjects were collected before any medication was given. Samples were centrifuged at 3000 rpm; serum was separated and stored at -70°C.

Serum homocysteine concentration was analysed by fluorescence polarization immunoassay (FPIA) method by Abbott's Ax SYM system.

Serum MDA levels were measured by Thiobarbituric acid reactive substances assay (TBARS) at 532 nm wavelength by spectrophotometer.

Ascorbic acid concentration was measured by Dinitrophenyl Hydrazine (DNPH) method at 520 nm wavelength by spectrophotometer and uric acid concentration was measured by Uricase method using Erba Mannheim Diagnostics kit at 550 nm.

Statistical analysis was performed using Mann Whitney U test and Spearman Correlation analysis.

**Results**

Regardless of the preeclampsia none of the subjects had any chronic disease or pregnancy complications. No significant difference was found between the clinical features of the preeclamptics and the normotensive pregnant controls other than hypertension and proteinuria Table I.

Table I: Sociodemographic features of the subjects

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Parameters</th>
<th>Cases (n=50) (Mean ± SD)</th>
<th>Controls (n=50) (mean ± SD)</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Age</td>
<td>25.30±2.59</td>
<td>26.75±2.12</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>2</td>
<td>Gestational age (days)</td>
<td>235 ±13</td>
<td>226 ±14</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>3</td>
<td>Pregnancy (Number)</td>
<td>4.3 ±1.9</td>
<td>4.6 ±2.2</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>4</td>
<td>Blood pressure (mmHg) Systole Diastole</td>
<td>164 ±12 114 ±6</td>
<td>116 ±11 76 ±4</td>
<td>&lt;0.05*</td>
</tr>
<tr>
<td>5</td>
<td>Proteinuria (mg/24 hours)</td>
<td>5300 ±1250</td>
<td>54 ±24</td>
<td>&lt;0.05</td>
</tr>
</tbody>
</table>

Serum homocysteine MDA and antioxidant status concentrations of preeclamptics and normotensive pregnant controls were given in Table II.

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Parameters</th>
<th>Cases (n=50) (Mean ± SD)</th>
<th>Controls (n=50) (mean ± SD)</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Homocysteine (µmol/l)</td>
<td>18.78 ±6.71</td>
<td>12.59 ±5.06</td>
<td>&lt;0.05*</td>
</tr>
<tr>
<td>2</td>
<td>Malondialdehyde (nmol/l)</td>
<td>5.46 ±0.20</td>
<td>3.46 ±0.18</td>
<td>&lt;0.05*</td>
</tr>
<tr>
<td>3</td>
<td>Ascorbic acid (mg/dl)</td>
<td>0.72 ±0.21</td>
<td>0.98 ±0.17</td>
<td>&lt;0.01*</td>
</tr>
<tr>
<td>4</td>
<td>Uric acid (mg/dl)</td>
<td>8.16 ±1.68</td>
<td>4.72±0.91</td>
<td>&lt;0.01*</td>
</tr>
</tbody>
</table>

(*: significant)

**Discussion**

Preeclampsia is one of the leading causes of maternal and perinatal mortality in the developing countries. It affects as many as 8,370,000 causes worldwide per year. Though the etiopathogenesis of preeclampsia is largely not understood, oxidative stress and a generalized inflammatory state forms the principle factors contributing to preeclampsia15.

Our study suggests that Hcy and a marker of free radical, MDA were significantly increased in patients of preeclampsia.

Preeclampsia has been proposed as a two stage disorder. In the first stage, the placenta produces a cytotoxic factor and in the second stages the increased mitochondrial activity of the placenta results in increased generation of ROS16.

Abnormal placenta in preeclampsia results in the placental ischemia17. The ischemia reperfusion injury to the placenta leads to generation of placental oxidative stress and hence, increased synthesis of ROS. Increased superoxide ion generation in the placenta has been detected with the direct electron paramagnetic spin resonance technique18.

The deleterious effects of free radicals include the initiation of lipid peroxidation, cellular dysfunction and leukocyte activation leading to endothelial dysfunction19.

Numerous independent studies demonstrating a significant association between preeclampsia and the levels of various biomarkers of oxidative stress have strengthened our evidence for lipid peroxidation in preeclamptic patients20,21.

Serum concentrations of Hcy decrease in normal pregnancy either due to physiological response to pregnancy, increase in estrogen or hemodilution22.
Our study demonstrates an increase in the Hcy levels in the patients with preeclampsia along with a positive correlation in between MDA and Hcy in these patients. In a study conducted it was shown that hyperhomocysteinemia decreases total vessel surface and disrupts placental perfusion in preeclampsia. It is also seen that vascular damage in maternal uteroplacental circulation is present in endothelial dysfunction and smooth muscle cell proliferation, both of which are characteristic features of preeclampsia.

Along with these findings it is corroborated that there is decreased endothelial nitric oxide (NO) synthesis and activity due to effect of increased Hcy levels. In consistent with present findings, Vanderget et al. and Rajkovic et al. also found Hcy concentration to be raised in preeclamptic patients. Preeclampsia is associated with increased utilization of the antioxidants. Several studies have demonstrated presence of decrease serum levels of ascorbic acid as compared to normal pregnant women and same findings were found in the present study.

There is a significant increase in the serum uric acid levels in the present study. It implicates antioxidative response related to the pathogenesis of preeclampsia. Hence, in conclusion, preeclampsia predisposes the vascular endothelium to oxidative stress and elevated Hcy levels. There is increase in total oxidant and decrease in antioxidant activities of these patients leading to endothelial injury and vasospasm. There is need to further evaluate the Hcy lowering effect of vitamin B6, vitamin B12 and folate supplementation. Also, the results of ongoing trials using vitamin C and E supplementation may be able to delineate the role of oxidative stress in the pathophysiology of preeclampsia.

References


Knowledge, attitude, practice regarding breast feeding practice among mothers attending Alluri Sita Rama Raju Academy of Medical Sciences, Eluru, Andhra Pradesh

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Abstract

Background
Breast feeding practice rates vary in different places. Little is known about breast feeding practice and barriers women experience. In addition information is scant on how organisations can best promote breastfeeding. It is estimated that a breastfed baby is 14 times less likely to die from diarrhoea, 4 times less likely to die from respiratory diseases and 2.5 times less likely to die from other infections than a non breast fed infant.

Objectives
1) To determine knowledge, attitude, and practice of rural mothers regarding breastfeeding. 2) To find association of knowledge, attitude, and practice of rural mothers regarding breast feeding with demographic variables.

Methodology
The hospital based descriptive study was undertaken in departments of Paediatrics and Gynaecology, Obstetrics in both the outpatient and inpatient wards in Alluri Sita Rama Raju Academy of Medical Sciences, Eluru, Andhra Pradesh. In this study 214 mothers who had children less than 2 years of age were selected. A structured questionnaire was applied and children above two years of age group were excluded from the study.

Results
About 32.7% of the mothers had the knowledge regarding 6 months of breast feeding is enough. 86.9% of the mothers gave the opinion regarding attitude, breast feeding is a good way to decrease family expences. Out of 214 mothers who were interviewed 27 mothers are illiterates, 39 mothers have primary schooling, 105 mothers have high school education, 23 mothers had secondary education, and 20 mothers are graduates. 78.9% mothers were > 24 yrs of age and 20.1% mothers were < 24 years of age. Of the 214 mothers interviewed 189 mothers gave colostrums, whereas 25 mothers (11.81%) discarded colostrums. In mothers who delivered by vaginal delivery out of 140 mothers 118(84.2%) initiated breastfeeding within 12 hrs after delivery whereas out of 70 mothers who delivered by caesarean section 49(70%) initiated breast feeding 12 hrs after delivery.

Conclusions
In the present study conducted on 214 mothers showed that high proportion of population gave exclusive breast feeding. However lot of gap is seen between literate and illiterate mothers in breast feeding practices which need to be changed. The positive changing trends are seen in aspects of colostrum feeding. Hence there is a need for awareness programs regarding breast feeding in this area.

Key Words
Knowledge, Attitude, Practice Regarding Breast Feeding.

Introduction
Children are our future and most precious resources. After the birth the health of the baby depends upon the nurturing practice adopted by the family. Breast feeding is the first step in life which ensures that infants and young children get a healthy and nutritious state in life. It is one of the few consistent sources of energy dense food. The first milk is most suitable for new born, the sastras call it Peeyusha and Western sciences use the word Colostrums. It is the infant’s first immunization. For most of the children breast feeding makes the difference between life and death and it is the infants Passport to life. Exclusive breast feeding can save many lives by preventing malnutrition and reducing the risk of infections and hypothermia. Exclusive breastfeeding should be practiced for at least four months and preferably six months in poor countries since they have a high risk of infection through contaminated water and food. In the year 2001-2002 a survey conducted in a rural area of district Dehradun, Uttaranchal suggested that 56.3% mothers reported to have fed colostrums, 74.1% mothers put their children on full breast feeding while only 26.0% with partial feeding. Breast feeding practice rates vary in different places. Little is known about breast feeding and barriers of women experience. In addition information is scant on how Organizations can best promote breastfeeding. To improve rates of full
breastfeeding, specific information about the beliefs and practices that influence this outcome is needed. The duration of breastfeeding varies with rural or urban residence, literacy, socio-economic status and other factors. Keeping in mind the above viewpoints this study is conducted to assess various socio-economic and cultural factors associated with knowledge, attitude and practices regarding breastfeeding in a rural community of Eluru.

**Objectives**

1. To determine knowledge, attitude, and practice of rural mothers regarding breast feeding.
2. To find association of knowledge, attitude, and practice of rural mothers regarding breast feeding with demographic variables.

**Material & Methods**

A hospital based descriptive study was undertaken in departments of Paediatrics and Gynaecology, Obstetrics in both the outpatient and inpatient wards in Alluri Sita Rama Raju Academy of Medical Sciences, Eluru, Andhra Pradesh during the period of 6(six) months.

Inclusion criteria: 214 mothers who had children less than 2 years of age were selected.

Exclusion Criteria: Above two years of children excluded.

A structured questionnaire with 40 items was used. It included

1. Demographic variables (12 items)
2. Related to knowledge (13 items)
3. Related to attitude (8 items)
4. Related to practice (7 items)

Pretested questionnaire and validity of the tool was established in consultation with guide, experts from the departments of obstetrics, gynaecology, and paediatrics. The statistical package Microsoft office excel 2003 was used for data processing and statistical analysis. Variables were described using frequency distribution for categorical and mean and standard deviations for continuous variables.

**Results**

Out of 214 mothers who were interviewed, of which, 78.9% mothers were > 24 yrs of age 20.1% mother were < 24 years of age. Regarding educational status of mothers, 12.6% of mothers were illiterates, 18.2% mothers have completed primary school, 49.1% mothers have completed high school education, 10.7% of mothers had secondary education, and 19.4% mothers were graduates. 91.6 % mothers were housewife and remaining mothers were working group. About 75.7% of mothers having > 2 children and remaining 24.3% were having > 2 children. Regarding the status of birth, about 65.5% deliveries taken place vaginal route and remaining 34.5% deliveries were caesarian section. About 19.6% of child birth needed Hospital admission.

Out of 214 mothers were interviewed, of which, each mother was given multiple answers regarding knowledge and attitude variables. About 32.7% of the mothers had the knowledge regarding 6 months of breast feeding is long enough. 86.9% of the mothers gave the opinion regarding attitude, breast feeding is a good way to decrease family expenses.

Table 1: Socio-Demographic characteristics of Study population:

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt;24</td>
<td>171</td>
<td>79.9</td>
</tr>
<tr>
<td>=24</td>
<td>43</td>
<td>20.1</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Education</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Illiterate</td>
<td>27</td>
<td>12.6</td>
</tr>
<tr>
<td>Primary Schooling</td>
<td>39</td>
<td>18.2</td>
</tr>
<tr>
<td>High School</td>
<td>105</td>
<td>49.1</td>
</tr>
<tr>
<td>Secondary Education</td>
<td>23</td>
<td>10.7</td>
</tr>
<tr>
<td>Graduation/Post Graduation</td>
<td>20</td>
<td>9.4</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Employment</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>House Wife</td>
<td>196</td>
<td>91.6</td>
</tr>
<tr>
<td>Employed</td>
<td>18</td>
<td>8.41</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>No. of Children</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>=2</td>
<td>162</td>
<td>75.7</td>
</tr>
<tr>
<td>&gt;2</td>
<td>52</td>
<td>24.2</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Type of Family</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Joint</td>
<td>121</td>
<td>56.5</td>
</tr>
<tr>
<td>Nuclear</td>
<td>93</td>
<td>43.5</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Type of Birth</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaginal</td>
<td>149</td>
<td>65.5</td>
</tr>
<tr>
<td>Caesarean Section</td>
<td>74</td>
<td>34.5</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Neonatal Hospitalization</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>42</td>
<td>19.6</td>
</tr>
<tr>
<td>No</td>
<td>172</td>
<td>80.4</td>
</tr>
</tbody>
</table>

Table 2: Mothers’ knowledge and attitude about breast feeding:

<table>
<thead>
<tr>
<th>Item</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Knowledge</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6 Months of Breast Feeding is long enough</td>
<td>70</td>
<td>52.7</td>
</tr>
<tr>
<td>Breast Feeding will provide immunity to the child</td>
<td>108</td>
<td>50.4</td>
</tr>
<tr>
<td>Breast Feeding will decrease diarrhoea</td>
<td>66</td>
<td>30.8</td>
</tr>
<tr>
<td>Regarding expression and storage of breast milk</td>
<td>62</td>
<td>28.9</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attitude</th>
<th>n</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Breast Feeding is convenient than feeding infant formula</td>
<td>214</td>
<td>100</td>
</tr>
<tr>
<td>It is not difficult for Breast Feeding mother to care her family</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Breast Feeding has no negative effect on marital relationship</td>
<td>183</td>
<td>85.5</td>
</tr>
<tr>
<td>Breast Feeding is a good way to decrease family expenses</td>
<td>186</td>
<td>86.9</td>
</tr>
<tr>
<td>Community Encourages Breast Feeding over feeding infant formula</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Doctors and Nurses encourage Breast Feeding</td>
<td>148</td>
<td>69.1</td>
</tr>
</tbody>
</table>

Table 3 reveals that of the 214 mothers interviewed 189 mothers gave colostrums, whereas 25 mothers (11.81%) discarded colostrums.
The significant association between educational status and colostrum feeding was tested using chi-square test. Using chi-square test, the test result was significant at 1% level of significance. X2 value is 6.86 and the ‘p’ value is <0.01. This showed that there was an association between education qualifications of the mother with feeding of colostrum.

Table 4 depicts that about 32% of the mothers given prelacteal feeds to child and remaining 68% of the mothers not given prelacteal feeds.

Table 5 shows that in the present study, 74.5% started giving breast feed before 12 hours after delivery whereas 25.5% mothers gave breast feed 12 hours after delivery. In literate mothers, 152 (81.34%) started breast feeding before 12 hrs after delivery, whereas 16(60%) of illiterate mothers started breast feeding before 12 hrs after delivery.

Table 6 reveals that in initiation of breast feeding, mode of delivery also played an important role. In mothers who delivered by vaginal delivery out of 140 mothers 118(84.2%) initiated breastfeeding within 12 hrs after delivery whereas out of 70 mothers who delivered by caesarean section 49(70%) initiated breast feeding 12 hrs after delivery.

Table 7 depicts that of the 214 mothers interviewed 122 mothers (57.2%) gave exclusive breast feeding up to 6 months period and 92 mothers (42.8%) did not give exclusive breast feeding. Out of 92 mothers, 77 mothers were literates and 15 were illiterates.

Discussion

This cross sectional study was conducted to determine the knowledge, attitude, practice regarding breast feeding mothers from rural population and also to find Association of knowledge, attitude, and practice with demographic profiles of rural Population. In this study 100% of mothers responded positively that breast feeding is Convenient, also women had better attitude regarding breast feeding.

This study showed that 11.81% discarded colostrums. The present study is not Coinciding with study of S. PSrivatsava, Vijay Kumar, and Sharma et al 1 where 82.91% of mothers discarded colostrums. The probable discordance may be due to increased awareness and also due to inclusion of more educated women in the present study because educated women had better practice. In this study 75.5% mothers started breast feeding before...
12 hours after delivery which accounted for 81.34% of literates and 60% of illiterates. In Tamilnadu a study at Chengalput by BPNI 71% of mothers initiated breast feeding at right time, which is similar to the present study. The 10th year plan goal of early initiation of breast feeding is to increase it from 15.8% to 50% by 2007 which is surpassed in this study. The mothers who delivered by caesarean section delayed initiation of breast feeding, where mothers rarely care for their babies in the first 2 days post-operatively.

The present study showed that practice of exclusive breast feeding is higher in rural and illiterate mothers when compared to literate mothers. Pragti Chhabra Vijay L. Grover et al 4 demonstrated in their study the same proportion which is similar to the present study. In the present study literate mothers introduced complementary feeds earlier than illiterate mothers. In a study by BPNI conducted at Chengalput and Chidambaram districts of Tamilnadu, the starting of complementary feeding at 6 months of age was around 74% and 88.3% respectively, which is similar to the present study. The early start of complimentary feeding by literate mothers is due to impact from media as they get attracted to fancy advertisements about complimentary feeds and also due to poor knowledge regarding expression and storage of breast milk among working women.

Majority of the participants reported that reason for stopping of breast feeding is that milk is not sufficient. Education of women is required in this aspect and also regarding time needed for colostrums to change to transitional milk and education regarding ways of successful breast feeding can be valuable in decreasing concern about milk supply. Women reported that community and health workers encouraged breast feeding.

A limitation of our study was its short period of study, cross sectional design and also active non intervention. For inclusion in this study, the child’s age is limited to a maximum of 2 years. This would have diminished the risk of recall bias. Further cohort studies are needed to explore the effect of socioeconomic development on children health and nutrition.

**Conclusions**

Based on the results observed in the present study, 214 mothers showed that high proportion of population gave exclusive breast feeding. However lot of gap was seen between literate and illiterate mothers in breast feeding practices which need to be changed. The positive changing trends are seen in aspects of colostrum feeding, early initiation of feeding. The changing trends are negative in aspects of exclusive breast and duration of breast feeding. Literacy rate plays a major role in this aspect even many mothers feel that they will spoil the health of the baby if they feed with expressed breast milk which needs to be changed. Hence there is a need for initiation of wide IEC programs regarding breast feeding practices.

**Acknowledgements**

I wish to express my sincere thanks to all study subjects, research funding agency ICMR and completed this study under Student short term fellowship from ICMR and HOD of OBG and HOD of Paediatrics of Alluri Sita Rama Raju Academy of Medical Sciences, Eluru, A.P for helping me to complete the required sample.

**References**

2. BPNI Report 2003 Status of infant and young child feeding in 49 districts (98 blocks) of India.
Comparison of sub mucosal diathermy and partial resection of inferior turbinate in the treatment of symptomatic nasal valve blockage

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Abstract

Objectives
To evaluate the efficacy of sub mucosal diathermy (SMD) and partial resection of inferior turbinate (PRIT) in the treatment of symptomatic enlarged inferior turbinate.

Study Design: Prospective

Methods
Sixty patients of age group 18 -56 yrs. with symptomatic enlarged inferior turbinate had given choices for SMD and PRIT. All the patients had history of failed medical treatment.

Results
Each thirty patients underwent SMD (group I), PRIT (group II), eight patients of group I, have anterior nasal packing after surgery for bleeding. Four patients complained of excessive rhinorrhoea for first 2 weeks while 4 patients of Group 1 complained of nasal blockage for 1 week even after intervention. In group 2, 8 patients have re-anterior nasal packing after pack removal. Both groups followed up for 6 months. 13 patients were lost in follow up, so excluded from the study. Following 6 months of follow up, 8 patient of group I had recurrence with nasal blockage and in gr. II none had recurrence.

Conclusion
PRIT is better than SMD in long course; nevertheless it should be reserved for failed SMD, not as a primary option. Ink described the nasal valve in 1903. The nasal valve is formed medially by the septum and laterally by the caudal edge of the upper lateral cartilage and it accounts for approximately 50% of total upper airway resistance. The anterior tip of the inferior turbinate is found in the nasal valve region, and hypertrophy of this structure can cause exponential increase in airway resistance.

Introduction
Nasal valve is area of greatest constriction throughout the entire respiratory tract; limited medially by the septum, inferiorly by the floor of the nose and by the anterior portion of inferior turbinate. Nasal valve is a dynamic valve as swelling of the venous erectile tissue of the inferior turbinate and nasal septum can cause complete obstruction of the nasal passage. Enlargement of the inferior turbinate is mainly due to swelling of the sub mucosa and rarely due to enlargement of the bone itself. Hypertrophy of inferior turbinate caused by dilation of sub mucosal venous sinusoids is the cause in intrinsic rhinitis, and responds to decongestant. Sometimes the inferior turbinate enlargement due to sub mucosal fibrosis does not respond to decongestant. In few cases of inferior turbinate hypertrophy, the venous sinusoids become atonic and also do not respond to decongestant. When inferior turbinate hypertrophy is symptomatic, it needs treatment. There are different modalities of treatment but most popular and effective are SMD and PRIT. This paper aims to compare the efficacy of these methods in the treatment of symptomatic inferior turbinate hypertrophy.

Material and Methods
All together 60 patients (36 women and 24 men) with symptomatic inferior turbinate hypertrophy were included for the study. The patients were of age group 18-56 and they had history of failed medical treatment. After counseling all the patients and discussing all the pros and cons of both surgical interventions, patients were given choices to select their surgical procedure themselves. 30 patients (24 women and 6 men) underwent SMD (sub mucosal diathermy) under sedation and given a tag Group 1, and 30 patients (12 women and 18 men) underwent PRIT under general anesthesia and were given tag of Group 2. After surgical intervention in Group 1, nasal cavity were filled with antibiotic ointment and patients were discharged on the same day, whereas Group 2 patients were discharged after anterior nasal packing removal on the 3rd post operative day. In both group broad spectrum antibiotic and NSAID were given.

Results

Group 1
- Out of 30 patients of Group 1 (SMD) 8 patients needed anterior nasal packing after surgical intervention and discharged then after.
- 4 patients complained of excessive rhinorrhoea for the first 2 weeks.
- 4 patients complained of nasal blockage just after the surgical intervention.

**Group 2**
- Out of 30 patients 8 had to have reanterior nasal packing after pack removal after 48 hours of operation, for the bleeding and their discharge was delayed.
- 3 patients complained of nasal dryness and excessive crusting for 2 months.
- Both the groups of patients were followed up regularly- weekly for the first 2 weeks, 2 weekly for one month, and then every month till 6 months.
- At the end of 6 months we lost 13 patients (7 of Gr1 and 6 of Gr2) and one patients of Group 2 died in road accident. At the completion of 6 months 8 patients of Group 1 had recurrence of nasal blockage and in Group 2 none had recurrence.

**Discussion**
Symptoms of nasal obstruction may persist despite maximal medical management. In many patients who continue to complain of nasal obstruction, inferior turbinate hypertrophy can be confirmed by physical exam and rhinometry, though the latter is infrequently performed in clinical settings. It has been shown that inferior turbinate enlargement can prevent adequate medical management by preventing the transmission of topical steroids and topical antihistamines to the superior nasal cavity (4). So surgical procedures that reduce the size of the inferior turbinate can not only improve symptoms, but can also potentiate medical management of rhinitis. Numerous procedures exist for this purpose, and controversy abounds as to which is the best. There are very few randomized studies comparing different procedures to each other, and those that exist are generally not long-term studies. Procedures can be classified as those that address bony causes of nasal obstruction, and those that address mucous and sub mucous swelling. Patients with symptomatic nasal obstruction due to sorts of medical treatment, some sort of surgical intervention is recommended. The classically performed procedure for inferior turbinate hypertrophy was total turbinate resection. This procedure involves clamping the inferior turbinate at its base to achieve haemostasis, followed by the use of nasal scissors or endoscopic instruments to resect the entire turbinate along its base. This procedure definitively widens the nasal airway and has been shown to be one of the most effective procedures in achieving long-term nasal patency, with a retrospective study by Ophir et al showing that 80% of 150 patients had subjectively improved nasal breathing and 91% had widely patent nasal airways at an average follow-up time of 2.5 years (range 1 to 7). The most common complication of total inferior turbinectomy appears to be haemorrhage. The procedure often requires nasal packing after completion. Also, nasal crusting, synechiae, and discomfort are frequent occurrences for several months afterward because of exposed bone at the lateral nasal wall. A 1985 retrospective study by Moore et al condemned total inferior turbinectomy, reporting that 66 percent of their 18 patients had ozena, or advanced atrophic rhinitis characterized by chronic crusting and dysosmia even leading to anosmia due to destruction of olfactory cells. Others, such as Ophir, have refuted this notion and report that atrophic rhinitis is a rare and even insignificant complication of total turbinectomy. However, many otolaryngologists today have abandoned this procedure. Partial turbinectomy is a procedure developed to remove the anterior part of the inferior turbinate. It is directed at relieving obstruction at the nasal valve, while leaving a portion of the turbinate to continue its function of air conditioning. Nasal patency rates show great subjective improvement immediately after surgery, with one retrospective study suggesting that 70 of 76 patients reporting improvement at about 8 years (6). However, other studies have suggested decreased effectiveness with time (7), similar to nonresection procedures. Complications are similar to those for total turbinectomy, though the crusting is usually less severe, as is the risk of haemorrhage. Atrophic Rhinitis with this procedure is rare. Electrocautery has been used successfully in the ablation of inferior turbinates. Two forms of the procedure exist- submucosal diathermy, and mucosal cautery. Both procedures can be performed in the office under local anaesthesia. Mucosal cautery, as the name implies, utilizes the electrocautery device to burn from posterior to anterior along the inferior turbinate. This causes more pain and greater risk of haemorrhage. It also damages mucosa with subsequent increase in mucosal transport time. Submucosal diathermy avoids those risks. It involves inserting a bipolar cautery to cause a submucosal lesion along the inferior border of the inferior turbinate. The device frequently has two sharp points that are used to pierce the inferior portion of the inferior turbinate SMD- is an effective method of treatment for symptomatic inferior turbinate hypertrophy (Wenself, gleasa and siodlan, 1986). It reduces nasal blockage by 65 % (Jones et al 1989), but in our study it is 60%. Many rhinologist advocate SMD in cases where inferior turbinate shrinks with an alphareceptor agonist (Jones et al 1989).But nevertheless SMD in our study 4 complained of excessive rhinorrhoea for the first 2 weeks, then goes off automatically. Another 4 patients from the very beginning complained nasal blockage, initially it may due to post operative oedema, but as symptom lingers on for more then 2 months it indicate its negative aspects. In 6 months follow up we have 8 cases of failure, the cause may be fibrosis in the submucosal plain. Other popular method of surgical intervention is resection of inferior turbinate- partial resection or radical resection. PRIT is preferred one as radical trimming can cause unwanted results as atrophic rhinitis (Martinez et al 1983). In our study 3 patients complained of excessive
Custing for first 2 months, then we treated with nasal douching, which eventually disappeared. Up to 6 months we didn’t noticed any cases of atrophic rhinitis and we had no recurrence of nasal blockage in Group 2 (PRIT). It shows that anterior trimming is equally effective in reducing nasal blockage as radical operation (Weight, Jones, Clegg 1988) with less side effective. As PRIT has no recurrence of nasal blockage, and had nasal crusting in 3 patients (15%) in the initial period in our study; PRIT is a safe and effective procedure with minimal side effect (weight, Jones and Buckingham, 1990).

**Conclusion**

SMD should establish as a procedure in all patients with inferior turbinate hypertrophy unresponsive to medical treatment. If inferior turbinate hypertrophy recurs following SMD, partial resection of inferior turbinate (PRIT) should be carried out.

**References**

Patients’ perception towards professionalism in dentistry at K.L.E.S’s Institute of Dental Sciences, Belgaum, India

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Abstract

Background
Dentist - Patient relationship is the heart of successful dentistry. Physical appearance, behavior and communication skills of professionals might influence doctor-patient relationship.

Aim
The purpose of the present study was to determine how patients perceived about professionalism in dentistry.

Study Design
Cross-sectional survey.

Methods
The study was conducted on 305 subjects who visited the Department of Preventive and Community Dentistry, K.L.E.S’s Institute of Dental Sciences, Belgaum, Karnataka, India. It was conducted using a self-designed standard close ended questionnaire which included provision for recording demographic characteristics and questions pertaining to the appearance and professionalism of the dentists. The data was collected, tabulated and subjected to statistical analysis using chi – square test. Results: Attire and physical appearance of the dentists had an influence on the patients’ comfort and anxiety levels, and majority of patients agreed that the first impression of dentists affected their confidence level. Hairstyle, skin complexion and facial hair (for male dentists) appeared to have little effect on patients’ opinion of the various dental care providers.

Conclusion
Good human relationship between dentists and patients is essential for imparting positive attitude towards dentist and dentistry.

Key Words
Professionalism, dentistry, attire, behavior

Introduction
The term “relationship” means nothing more than indicating the relative position of two objects in time and space or in a classification system. What is life? When J. Krishna Murthy, a philosopher was asked by a person, he replied “Life is full of relationships”. From birth till death we all live a life of relationships. We establish, sustain, break, renew, rejuvenate and retreat our relationships with the world. Establishing and sustaining good and healthy relationship helps for better interaction. Human relationship implies that the participants have some understanding of each other’s behavior and act in ways that take this into account. Participants have expectations about each other, which influence the particular form of actions that will be taken.

Relationship between a doctor and his or her patient is one that is oriented toward the doctor helping the patient to deal effectively with a health problem. Patients cooperate with doctors and doctors attempt to return patients to as normal level of functioning as possible. Doctors usually take some type of action to satisfy the patients’ expectations. Doctors should be attentive not only to patients’ complaints but also to other factors that brought them to the doctor.

Profession refers to having special knowledge or skill in a particular field. Bernard Shaw rightly said, “Skill is not enough, trust is essential”. The layman might imagine that the primary concern of professionals would be professional competence. However, physical appearance, behavior and communication skills of professionals might influence doctor-patient relationship, especially for those patients who have never met the dentist previously.

Patients’ attitude towards the dental care becomes important because patients’ satisfaction with their dentists is a primary determinant to seek dental care, people who are dissatisfied tend to avoid care and jeopardize their dental health. Dentist - Patient relationship is the heart of successful dentistry. So, what does the public think of dentistry and what do dental patients think of dentists is vital for complete understanding of the image of dentistry. Sparse data are available to answer these questions, so the current study is carried out to assess the patients’ attitudes towards dentists and the dental profession.
Material and Methods

The study was conducted using a self-designed standard close ended questionnaire format to collect required and relevant information. To aid in the development of our questionnaire, previous surveys were reviewed and also observations done by us on various dentists were considered. Thus, the questions that addressed our concerns were selected. A copy of the questionnaire has been enclosed. Ethical clearance was obtained from the Local Ethical Committee and written informed consent was obtained from the participating subjects. The study was conducted at the Department of Preventive and Community Dentistry, K.L.E.S’s Institute of Dental Sciences, Belgaum, Karnataka, India.

Questionnaires were offered to all patients who visited the department to seek treatment for the first time in our department. Reason for including only the first time visitors was to avoid having too many patients with a long time relationship with their dentist, which could lead to better or worse patient satisfaction thus avoiding any possible bias in our results. Informed consent was obtained from all patients prior to enrolment. Questionnaires were prepared in three different local languages i.e., Kannada, Marathi and English, to enable the patient understand the questions appropriately and easily. Before patients filled their questionnaire format, the purpose of the study was explained to all of them. Each patient was given sufficient time to complete the questionnaire. Those who were unable to understand the questions properly were explained by the interviewer himself.

Questions were adapted to elicit demographic characteristics, education level, occupation, socioeconomic status (revised Kuppuswamy scale for socioeconomic status was slightly modified as per our convenience in our study) of the patients and also pertaining to the appearance, time management and professionalism of the dentist. The questionnaire contained eighteen statements and used 3 point likert scale to assess dentists professionalism (1-Agree, 2-Disagree, 3-Neutral). Patients were asked to classify the dentist based on their physical characteristics and behavior. Patients were also requested to provide their perception of the overall treatment procedure including comfort, anxiety, and confidence in the dental care provider and regarding their first impression on dentist.

The data was collected, tabulated and subjected to statistical analysis. As the data obtained in the present survey was- non parametric (nominal) data which is represented in frequencies, chi – square test was applied. The level of significance (α) was fixed at 5% (p = 0.05). SPSS version 12 was used for the present analysis.

Results

The study was conducted to know the patients’ perceptions of professionalism in dentistry in 305 patients (males 151 and females 154). The data obtained from the study was subjected to statistical analysis and the following results were obtained. Table 1 shows that majority of study subjects belonged to 20-29 years age group (33.8%). Males (49.5%) and females (50.5%) were almost equally distributed. Table 2 shows that majority of study subjects (34.8%) belonged to university level of education. Table 3 denotes that lower socioeconomic status comprised greater part of the study group (55.1%).

Data related to physical appearance of dentists shows that majority of study subjects agreed that skin complexion, hairstyle, attire and overall appearance of dentist affected their confidence level on them (Table 4), yet not statistically significant. Dentist’s built, facial hair of the male dentist and facial expression were found to

Table 1: Distribution of Study Subjects Based on Age and Gender

<table>
<thead>
<tr>
<th>Age</th>
<th>Males</th>
<th>Females</th>
<th>Total N (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 – 19</td>
<td>11</td>
<td>29</td>
<td>40 (13.1)</td>
</tr>
<tr>
<td>20 – 29</td>
<td>47</td>
<td>56</td>
<td>103 (33.8)</td>
</tr>
<tr>
<td>30 – 39</td>
<td>30</td>
<td>24</td>
<td>54 (17.7)</td>
</tr>
<tr>
<td>40 – 49</td>
<td>27</td>
<td>21</td>
<td>48 (15.7)</td>
</tr>
<tr>
<td>&gt;50</td>
<td>36</td>
<td>24</td>
<td>60 (19.7)</td>
</tr>
<tr>
<td>Total</td>
<td>151</td>
<td>154</td>
<td>305 (100)</td>
</tr>
</tbody>
</table>

Table 2: Distribution of Study Subjects Based on Different Level of Education

<table>
<thead>
<tr>
<th>Level of Education</th>
<th>Number of Subjects n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Illiterate</td>
<td>70 (22.9)</td>
</tr>
<tr>
<td>Elementary</td>
<td>59 (19.4)</td>
</tr>
<tr>
<td>High school</td>
<td>70 (22.9)</td>
</tr>
<tr>
<td>University</td>
<td>106 (34.8)</td>
</tr>
</tbody>
</table>

Table 3: Distribution of Study Subjects Based on Socioeconomic Status

<table>
<thead>
<tr>
<th>Socioeconomic Status</th>
<th>Number of Subjects n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Upper</td>
<td>8 (2.6)</td>
</tr>
<tr>
<td>Middle</td>
<td>129 (42.3)</td>
</tr>
<tr>
<td>Lower</td>
<td>168 (55.1)</td>
</tr>
</tbody>
</table>

Table 4: Response Given by Participants to Various Questions Related to the Physical Appearance of the Dentists

<table>
<thead>
<tr>
<th>Questions</th>
<th>Agree n (%)</th>
<th>Disagree n (%)</th>
<th>Neutral n (%)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Skin Complexion</td>
<td>130 (42.6)</td>
<td>128 (42)</td>
<td>47 (15.4)</td>
<td>0.12</td>
</tr>
<tr>
<td>Hairstyle</td>
<td>155 (50.8)</td>
<td>102 (33.5)</td>
<td>48 (15.7)</td>
<td>0.18</td>
</tr>
<tr>
<td>Dentist’s Dress</td>
<td>229 (75.1)</td>
<td>63 (20.6)</td>
<td>13 (4.3)</td>
<td>0.13</td>
</tr>
<tr>
<td>Dentist’s overall appearance</td>
<td>254 (83.2)</td>
<td>21 (6.9)</td>
<td>30 (9.9)</td>
<td>0.1</td>
</tr>
</tbody>
</table>
influence their opinion about dentists. Distribution was done based on socioeconomic status and the results were statistically significant. (Table 5)

Opinion about time management, dentist’s working unit, treatment procedure, appraisal of expenditure of treatment, dentist’s voice, first impression of the dentist (Table 6) revealed that majority of the study subjects agreed that these variables were of major concern for them but the results were not statistically significant. Dentists’ politeness, attentiveness, behavior and communicative skill (Table 7) were found to profoundly influence the patients and when these variables were stratified according to socio economic status they were statistically significant.

**Discussion**

Professionalism is the basis of medicine’s contract with society. It entails placing the interest of patients above those of the health care workers, while setting and maintaining clear standard of behavior that emphasize compassion, competence and integrity. For maintenance of professionalism in our health care environment, we must look out for some very specific outcomes. These outcomes will ensure professional behavior as a norm in our health care environment. Effective practice of dentistry requires not only technical competence but also behavioral knowledge and skills to achieve one’s full professional potential.

This study was conducted to explore patient’s perceptions of professional attire and behavior of dental care providers as indicators of the professional integrity and competence. Professional integrity is the ability of the health care provider to be worthy of trust, have concern for others, and have a substantial commitment to the patient. Professional competence is the health care provider’s ability to provide adequate dental care. A total of 305 patients (151 males and 154 females) were surveyed and the results are tabulated and compiled. Total valid comparisons could not be done between the present study and other studies reported in the literature due to wide variations observed with respect to selected age groups and methodologies employed. However, an attempt is done to compare and discuss to the extent possible and permissible.

Majority of the patients in our study agreed that dentist’s built (p<0.001), skin complexion (p=0.12), facial hair of male dentist (p<0.001), hair style (p=0.18) affected their confidence in them. (Table 4 and 5) The general patient preference toward traditional/ formal physician dress observed by Kanzler and Gorsulowsky6, and Gjerdingen et al6 was confirmed by our data. More than 75% of the patients agreed that formal attire was appropriate for dental care providers and should be required to wear clean trousers and shirt with polished shoes and white apron. Despite a recent movement towards more relaxed, casual attire in the general public and within the office
setting, it appears that patients continue to expect a formally dressed dental health care provider.

The patients’ responses were largely in the segment of “AGREE” by more than 80%, when questions related to facial expression (p=0.04), time management (p=0.07), treatment procedure (p=0.3), politeness (p=0.005) and attentiveness (p=0.03) were asked (Table 5, 6 and 7 respectively). This reflects positive attitude and professional competence exhibited by the dental care providers in the department clinic while rendering health services.

In the present study majority agreed that, dentist’s voice, behavior, working unit was satisfactory while providing the dental treatment. The first impression created in the mind of the patient about the physician can have strong influence in the future relationship and can positively or negatively reflect the rapport. Majority of patients agreed that their first impression of dentist affected their confidence level.

Successful doctor is one who knows how well to communicate with his/her patient. The doctor who is able to communicate with his/her patient is bound to give maximum psychological satisfaction to their patients. In the current study it was found that majority of patients were neutral with respect to the question asked related to the dentist’s communicative skill (p=0.001) (Table 7). It is a pointer towards improvement in communication skills. Positive patient attitudes can lead to an increase in exchange of information between the patient and the dental health care provider, in turn leading to more efficient and improved delivery of dental care.

Our findings indicate that patients are satisfied with the dental care they received, value their dentist’s competence and caring attributes. In brief, patients are pleased with their dentist. Dr. Michael Perich said, "Dentistry is a one-to-one situation. Each individual dentist, properly treating his or her patient, enhances dentistry’s image". Dentists can improve the image of dentistry by providing skillful dentistry, communicating articulately and clearly with patients, and showing patients that they are cared for. It is also important that the profession frequently monitors patients’ attitudes towards dentistry and dentists.

Practicing dentists who are aware of potential consumers concern can use this knowledge to improve their skills in treating patients and acquire an increased sense of control in their attempts to improve their practices.

**Conclusions**

1. Majority of the study subjects opined that dental care providers’ professional appearance and behavior influenced their confidence.
2. Patients preferred more formal dress attire from the dental care providers.
3. Attire greatly affected both the comfort and anxiety levels of the patients.
4. Hairstyle, skin complexion and facial hair (male dentist) appeared to have little effect on patient’s opinion of the various dental care providers.
5. Majority of the study subjects opined that dental care providers’ voice affected their anxiety level.
6. 50% of the study subjects were neutral in their opinion regarding dentist’s communicative skills.

Probing questions which can provide further information related to the influence of independent variables selected in the study on patients’ satisfaction are required to have better understanding of patients’ perception towards dentist and dentistry.

**Acknowledgements**

We would like to thank all the subjects who participated in the study.
References

Telavancin: A promising weapon against antimicrobial resistance

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Abstract
Antimicrobial resistance threatens the clinical effectiveness of many current antimicrobials, while jeopardizing important advances being made against major infections. This has led the World Health Organisation (WHO) to adopt Antimicrobial resistance as its theme for World Health Day 2011. While rational prescribing practices will help decrease the problem of resistance, we must have reserve drugs in our hands to fight this ever-increasing menace.

Over the past few years, there has been an intensified focus on the emergence of methicillin-resistant Staphylococcus aureus (MRSA), including both community-acquired and hospital-acquired strains, against which vancomycin became the drug of choice. Increased frequency of staphylococcal and enterococcal resistance to vancomycin has prompted a search for newer anti-MRSA agents.

Telavancin, a semi-synthetic derivative of vancomycin, was approved in September 2009 by the FDA for the treatment of complicated skin and skin structure infections, (cSSSI), caused by susceptible gram-positive bacteria, especially methicillin-resistant S. aureus (MRSA). It is also being proposed for treatment of nosocomial pneumonia, with focus on MRSA infections.

Major advantage of telavancin over vancomycin is its therapeutic potential against vancomycin-resistant organisms, including vancomycin-resistant enterococci and vancomycin-resistant S. aureus (VRSA). Telavancin inhibits bacterial cell wall synthesis and is ten times more active than vancomycin in this respect. However, unlike vancomycin, which has a slow, primarily bacteriostatic action telavancin demonstrates rapidly bactericidal properties. It also disrupts bacterial cell membrane potential and increases permeability and outflow of essential ions. It shows considerable penetration into skin blister fluid, thus accounting for its use in cSSIs. With predictable and linear pharmacokinetic profile and low level of resistance against it, telavancin is a welcome addition to the clinicians’ armoury in the fight against antimicrobial resistance.

Key Words
Antimicrobial resistance, MRSA, Telavancin, Complicated skin and skin structure infections.

Introduction
The widespread danger of antimicrobial resistance threatens the continued effectiveness of many current medicines to treat the sick, while jeopardizing important advances being made against major infectious killers. This has led the World Health Organisation (WHO) to adopt Antimicrobial resistance as its theme for World Health Day 2011. While rational prescribing practices will help decrease the problem of resistance, we must have reserve drugs in our hands to fight this ever-increasing menace.

Over the past few years, there has been an intensified focus on the emergence of methicillin-resistant Staphylococcus aureus (MRSA), including both community-acquired and hospital-acquired strains. Staph aureus is associated with non-invasive skin and soft tissue infections to invasive infections of the bone, cardiovascular system, blood and lungs. Shortly after the introduction of penicillin, resistant strains of S. aureus were isolated. This led to the development of methicillin to treat the resistant strains; however, MRSA was quickly noted and glycopeptide antibiotic vancomycin emerged as the drug of choice for MRSA infections1. Increased frequency of staphylococcal and enterococcal resistance to vancomycin has prompted a search for newer anti-MRSA agents like linezolid, daptomycin and quinupristin/dalfopristin to treat some types of MRSA infections2,3. Telavancin, a newer agent, demonstrates discernible advantages over some of the existing anti-MRSA drugs.

Telavancin, a semi-synthetic derivative of vancomycin, was approved in September 2009 by the FDA for the treatment of complicated skin and skin structure infections, (cSSSI), caused by susceptible gram-positive bacteria, especially methicillin-resistant S. aureus (MRSA)4.

Antimicrobial Spectrum
The spectrum of activity of telavancin is similar to that of vancomycin, with activity against staphylococci, streptococci, and enterococci. However, its major advantage over vancomycin is its therapeutic potential against vancomycin-resistant organisms, including vancomycin-resistant enterococci and vancomycin-resistant S. aureus (VRSA). Combination therapy, particularly with gentamicin, may improve bacterial killing against certain strains4.
Telavancin shows concentration-dependent activity in vitro against gram-positive organisms including multi-drug resistant isolates such as MRSA. It is also active against vancomycin-susceptible Enterococcus species in vitro. Strains of enterococci, primarily Enterococcus faecium, which contains the Van A gene, are vancomycin resistant and telavancin is similarly ineffective in vitro against such species. Telavancin may be active against vancomycin-intermediate Staphylococcus aureus (VISA). For vancomycin-resistant Staphylococcus aureus (VRSA), higher minimum inhibitory concentrations (MICs) of telavancin are needed.

As with vancomycin, telavancin is also effective against gram-positive anaerobic bacteria, including Clostridium, Corynebacterium, Lactobacillus, Propionibacterium and Peptostreptococcus species.

**Structure and Classification**

Telavancin is a lipoglycopeptide antibiotic, structurally derived from the glycopeptide vancomycin which has enhanced potency against gram positive organisms, especially those developing resistance to vancomycin.

Vancomycin and other glycopeptide antibiotics inhibit cell wall synthesis in susceptible bacteria by forming a carboxylate binding pocket that imparts strong affinity for peptidoglycan precursors containing the D-alanyl-D-alanine (D-Ala-D-Ala) motif. Telavancin is distinguished from these glycopeptide antibiotics by N-linked long chain acyl-D-glucosamine decorations.

With telavancin, increased potency results from ability to anchor onto the binding sites on the growing cell wall primarily due to the presence of lipophilic side chains. These hydrophobic side chains place this newer glycopeptide into a separate subset of antibiotics, namely, the lipoglycopeptides.

The lipoglycopeptide molecules dimerize and thus stabilize binding to the peptidoglycan cell wall. The telavancin dimers can then anchor themselves to the bacterial membrane by their hydrophobic substituents, increasing binding affinity to the D-alanine-D-alanine site and increasing potency.

On binding to the target, a favorable interaction occurs between neighbouring drug molecules which also stabilizes binding to the peptidoglycan structure.

**Mechanism of Action**

Telavancin binds to C terminal D-Ala-D-Ala residues and inhibits cell wall synthesis. However, unlike vancomycin, which has a slow, primarily bacteriostatic action telavancin demonstrates rapidly bactericidal properties. Due to its strong binding to bacterial cell membranes, telavancin is ten times more active than vancomycin at inhibiting both the transglycosylation and the synthesis of peptidoglycan by interfering with polymerization and cross-linking of the peptidoglycan framework.

Telavancin also disrupts bacterial cell membrane potential and permeability, leading to leakage of cellular ATP and potassium ions.

This dual mechanism of action may confer advantages over traditional glycopeptides due to both increased potency and rapid bactericidal activity.

**Pharmacokinetics**

Telavancin displays linear and predictable pharmacokinetic properties. It has a pharmacokinetic profile similar to that of vancomycin, but with high protein binding of about 93%. Telavancin has a half-life of 7.5 hours, which allows for convenient once-daily dosing. Telavancin also has a small volume of distribution (0.12 L/kg) and a high level of renal excretion (72%). Dose reductions for patients with moderate to severe renal dysfunction are required.

Penetration into skin blister fluid is important for its use in complicated skin and skin structure infections. Telavancin demonstrated considerable penetration into skin blister fluid, with penetration ratios of 0.79 and 0.82, respectively after 3 and 4 days of treatment. Telavancin also showed good penetration into the epithelial lining fluid of a healthy subject with a fluid: plasma concentration ratio of 0.75.

**Pharmacodynamic Profile**

When compared with other antimicrobials, for the treatment of methicillin-susceptible S. aureus (MSSA), telavancin was sixteen times more potent than vancomycin and forty times more potent than oxacillin. Against MRSA, telavancin was four and thirty times more potent than vancomycin and linezolid, respectively.

Postantibiotic effect (PAE) is the delayed growth of bacteria after removal of an antibiotic so that its concentration is below the MIC. In one study, there was a significant PAE for telavancin against S. aureus (including MSSA, MRSA and VISA) in vitro with a duration of 4–6 hours. Vancomycin has a PAE of only one hour.

Coagulase-negative staphylococci are often the cause of infections related to indwelling medical devices. These infections which may affect the skin and skin structures are commonly associated with a biofilm adherent to the device. They are difficult to treat because of either a low growth rate or a lack of antibiotic penetration into the biofilm. In a study, telavancin was compared with vancomycin, teicoplanin, linezolid, and moxifloxacin for the treatment of MSSA, MRSA, and glycopeptide intermediate S. aureus biofilms. Telavancin outperformed the glycopeptides and linezolid against all strains, but moxifloxacin proved to be superior to telavancin.
IN VIVO Animal Studies
Animal models suggest that telavancin may be effective in the treatment of soft-tissue infections, bacteremia, endocarditis, meningitis, and pneumonia caused by gram-positive pathogens.

Clinical Trials
Phase II trials
The FAST and FAST 2 studies were randomized, double-blind, active-controlled multicentre phase II trials conducted to evaluate the treatment of cSSSIs.

The study aim of FAST was to determine the efficacy of IV telavancin at a dose of 7.5 mg/kg/day as treatment for cSSSIs caused by gram-positive organisms compared with vancomycin 1 gm every 12 hours or anti-staphylococcal penicillins (nafcillin 2 gm every six hours, oxacillin 2 gm every six hours, or cloxacillin 0.5-1 gm every six hours).

FAST 2 was an identical study with a dose of telavancin 10mg/kg/day.

The results in both studies were similar and demonstrated comparable cure rates to the other agents, and no statistical difference was found.

A big limitation of the FAST studies was that in treating S. aureus (MSSA and MRSA) infections, the administered standard initial dose of vancomycin 1 gm 12 hourly IV does not achieve goal trough levels for most patients with normal renal function.

Phase III trials
1. Two pivotal, Phase III clinical trials (ATLAS I and II) were conducted to evaluate the efficacy and safety of telavancin in the treatment of cSSSI. The study aims were:

- To evaluate if telavancin is non-inferior to vancomycin for the treatment of cSSSI, including those infected with MRSA.
- If non-inferiority is established in both phase III clinical trials, a secondary goal was to test for superiority of telavancin to vancomycin in patients infected with MRSA for the treatment of cSSSI in pooled analyses.

These two phase III trials were identical parallel, active-control, randomized, double-blind multicentre trials. Patients were randomized to treatment with telavancin (10mg/kg IV every 24 hours) or vancomycin (1 gm IV every 12 hours) for 7-14 days.

In ATLAS I (n=1867), overall cure rates were 88% for the telavancin group and 87% for the vancomycin group. Again, cure rates for complicated SSSIs caused by MRSA were higher with telavancin than with vancomycin (90% and 85%, respectively).

ALTAS II was a retrospective sub-study of telavancin versus vancomycin for treatment of cSSSI associated with surgical procedures. The subset (n=194) of the original patients of ATLAS I was evaluated for clinical efficacy and microbiological efficacy on follow up (7-14 days at completion of therapy).

The results of both ATLAS I and ALTAS II did not demonstrate statistical difference between the groups.

The authors concluded that telavancin was non-inferior to vancomycin for the treatment of cSSSI, associated with a surgical site. The results of these trials proved that telavancin was at least equivalent to vancomycin for the treatment of cSSSIs, including those caused by MRSA; these results were the basis for telavancin receiving FDA approval.

2. Two phase III trials (ATTAIN1 and ATTAIN 2) evaluated the use of telavancin in patients with hospital-acquired pneumonia, with a focus on MRSA infections. The two trials, with a total of 1503 patients, aimed to prove non-inferiority in the clinical cure rate of nosocomial pneumonia with intravenous telavancin 10mg/kg once daily against intravenous vancomycin 1g 12 hrly. In both studies, telavancin achieved the objective of non-inferiority in the all-treated and clinically evaluable patient populations.

In one study, the antimicrobial activity of telavancin against 2279 clinical gram-positive cocci obtained from patients with nosocomial pneumonia including those with ventilator-acquired pneumonia (VAP) located in numerous medical centres worldwide was evaluated. Telavancin was highly active against Staphylococcus aureus, coagulase-negative staphylococci, Streptococcus pneumoniae, viridans group streptococci, β-haemolytic streptococci and vancomycin-susceptible enterococci. Telavancin inhibited all staphylococci at ≤0.5 mg/L. Among enterococci non-susceptible to vancomycin (all Enterococcus faecium), telavancin was active against isolates exhibiting a Van B phenotype, but less potent against Van A strains. Telavancin demonstrated equal or greater potency than the comparators (vancomycin, teicoplanin, dapto, linezolid and quinupristin/dalfopristin) against Gram-positive pathogens implicated in nosocomial pneumonia.

Phase IV trials (Post-marketing study)
The pharmaceutical sponsors of telavancin will conduct a prospective study over a five-year period after introduction of telavancin to the market to determine if decreased susceptibility to telavancin is occurring in the target population of bacteria.

Indications for USE
- Telavancin is approved by the FDA for the treatment of adult patients with complicated skin and skin structure infections (cSSSI) caused by susceptible Gram-positive bacteria i.e., Staphylococcus aureus (methicillin susceptible and methicillin-resistant isolates), Streptococcus pyogenes, Streptococcus...
agalactiae, Streptococcus anginosus group (S. anginosus, S. intermedius, S. constellatus) or Enterococcus faecalis (vancomycin-susceptible isolates only).

- A recent new drug application for telavancin was submitted to FDA for nosocomial pneumonia. However, the FDA issued a complete response letter indicating that additional data and analyses were required to support efficacy and safety of telavancin for treatment of hospital-acquired pneumonia\textsuperscript{15}.

### Dosage and Administration

Telavancin is supplied as 250- and 750-mg single-use vials and should be stored in a refrigerator at 2–8 degrees C. Before administration, the contents of the vial should be reconstituted to a concentration of 15 mg/mL and the appropriate amount removed and diluted to 100–250 mL. The recommended dose of telavancin is 10 mg/kg once daily administered over one hour for 7 to 14 days\textsuperscript{16}.

### Adverse Drug Reactions (ADRs)

The most common adverse events observed were taste disturbance (metallic or soapy taste), nausea, vomiting, headache and foamy urine. Decreased appetite, diarrhea, stomach pain, chills, pruritus, dizziness and pain or redness at the site of administration are other adverse effects\textsuperscript{16}. Serious adverse events include raised serum creatinine and nephrotoxicity, hypokalemia, QT prolongation, and infusion-related reactions (ie “Red-man syndrome”).

### Precautions and Contraindications

- **Renal Impairment:** Monitoring of renal function is required; if renal function decreases, we need to consider risk/benefit of continuing telavancin or changing to alternative agent. There is decreased efficacy of the drug in patients with moderate to severe renal impairment (CrCl \leq 50\text{mL/min}) compared to patients with CrCL \geq 50\text{mL/min}. Available recommended dosage adjustments in renal impairment are as follows:

<table>
<thead>
<tr>
<th>Creatinine Clearance (mL/min)</th>
<th>Dosage Regimen</th>
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<tbody>
<tr>
<td>30-50</td>
<td>7.5 mg/kg once daily (24 hours)</td>
</tr>
<tr>
<td>10- &lt; 30</td>
<td>10 mg/kg once in two days (every 48 hours)</td>
</tr>
</tbody>
</table>

- **Hepatic Impairment:** No dosage adjustments are recommended in patients with mild or moderate hepatic impairment.

- **Pregnancy:** Telavancin is classified as Pregnancy Category C, with potential risk of fetal development toxicity if pregnant female is exposed to it. A serum pregnancy test is recommended prior to the administration of telavancin along with effective contraception during therapy (if pregnant test result was negative). It has caused adverse fetal effects, including limb abnormalities, in animal studies. Women should avoid the use of telavancin during pregnancy unless potential benefit to the patient outweighs potential risk for the fetus.

### Lactation

Unknown whether telavancin is excreted in human milk.

### Cross Reactivity

Telavancin should be avoided in patients with known hypersensitivity to vancomycin.

### Infusion reactions

As with vancomycin, “Red-man syndrome” (with flushing, urticaria, pruritis, or rash) may occur with rapid infusion of telavancin\textsuperscript{1}.

### CDAD

Clostridium difficile-associated diarrhea (CDAD) may be considered in all patients with diarrhea following use of telavancin. Warning signs are watery or bloody stools, stomach cramps, or fever up to two or more months after stopping treatment\textsuperscript{16}.

### QTc prolongation

We should avoid telavancin in patients with a history of congenital QT syndrome, known prolongation of the QTc interval, uncompensated heart failure, or severe left ventricular hypertrophy, and caution is warranted in patients concomitantly receiving other drugs known to prolong the QTc interval.

### Drug Interactions

### Drug-drug interactions

- Renal dysfunction can occur on concurrent administration of telavancin with angiotensin-converting enzyme (ACE) inhibitors; angiotensin receptor blockers (ARBs) or with aspirin and other non-steroidal anti-inflammatory drugs (NSAIDS).

- Concurrent administration of telavancin with antiarrhythmic drugs like amiodarone, disopyramide, dofetilide, procainamide, quinidine and sotalol, antibiotics like erythromycin, sparfloxacin, moxifloxacin and gatifloxacin, antipsychotics such as pimozide and thioridazine and prokinetic drugs like cisapride carry the risk of QTc prolongation.

- In vitro studies have shown that telavancin inhibits CYP 3A4/5 at potentially clinically relevant concentrations.

### Drug-lab interactions

- Effects of Telavancin on Coagulation Test Parameters: Telavancin binds to the artificial phospholipid surfaces added to anticoagulation tests and interferes with the ability of the coagulation complexes to assemble on the surface of phospholipids and promote clotting in vitro. Thus, a false increase in prothrombin time (PT), international normalized ratio (INR), activated partial thromboplastin time (aPTT), activated clotting time (ACT) and coagulation based factor Xa tests may occur. It is recommended to obtain blood samples within 6 hours prior to the patient’s next dose of telavancin.
Effects of Telavancin on Urine Protein Tests:
Telavancin interferes with the urine qualitative dipstick protein assays and quantitative dye methods; however, it does not interfere with concomitant use of microalbumin assays.

Conclusion
Telavancin is a new lipoglycopeptide, for which phase III trials have been completed and is approved by the FDA. It is rapidly bactericidal and possesses activity against a broad spectrum of gram-positive organisms, including methicillin-resistant and vancomycin-resistant strains and anaerobic species. In clinical studies, efficacy was shown in complicated SSSIs and is also considered potential treatment of hospital-acquired pneumonia.

Key advantages of telavancin include activity against vancomycin-resistant strains, rapid bactericidal activity and a low level of resistance. As antimicrobial resistance continues to increase worldwide, telavancin will add to the clinicians’ armoury in the fight against resistance caused by gram positive bacteria.

References
Developing training modules for nurses in safe motherhood
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Abstract

Background
Maternal mortality remains a challenge in India. The current maternal mortality rate in India is 254 (India statistics, WHO, 2005). In Andhra Pradesh, the maternal mortality rate is 154 (2009) and the 2015 target is to reach below 100. The Each One save One (EOSO) program believes that no mother and child should lose their life in the natural process of childbirth. EOSO aims to reduce the incidence of maternal mortality in Andhra Pradesh by training student nurses on safe birthing practices.

Aims
The goals of the program are 1) to increase knowledge on safe birth practices among student nurses in Andhra Pradesh, India. 2) To improve the ability of nurses to identify complications during labor and to take quick action. And the objectives were that at the end of the program, 80% of the nurse participants at Modern Government Maternity Hospital will have an increase in knowledge about safe childbirth practices and complications.

Material and Methods
The program was run from 1st July 2010 to 30th August 2010. The program was targeted to the final year Bsc students of NIMS College and Laxmi College of nursing. The classes were conducted at the Modern Government Maternity Hospital, Hyderabad. There were a total of 61 nurses who attended the training modules. The program includes four educational training modules.

Result
The mean score on pretests was 7, and the mean score on post test was 10. A paired t-test conducted on participants who took both pre and post test, showed the result to be statistically significant (p < 0.05). By the end of the program, 91% of the nurses had increase in knowledge, when compared with the pretest.

Conclusion
The Each One Save One program can be used to educate more student nurses throughout Andhra Pradesh and the other states of India, where there is a lack of instructors. Our study shows that there was significant increase in knowledge among the nurses following the intervention program. This knowledge when converted to action will help in reducing maternal mortality rates.

Keywords
Maternal mortality, Nurses, Educational modules.

Introduction
1. Program Rationale
The Eight Millennium Development Goals (MDG) was created to get the world on the right path to better health. In September of 2000, leaders of 189 countries all over the world signed the Millennium Declaration which committed their countries to striving towards ending extreme poverty by 2015 through the 8 different Millennium Development Goals. The fifth goal is to improve maternal health. The two targets for the 5th Millennium Development Goal are:

a) Target 1: Reduce by three quarters, between 1990 and 2015, the maternal mortality ratio.

b) Target 2: To achieve, by 2015, universal access to reproductive health.

Maternal mortality remains a challenge for India. The current maternal mortality rate in India is 254 (India statistics, WHO, 2005). India's maternal mortalities account for about 25% of all maternal deaths in the world (Aparajita, Ramanakumar, 2005). In hopes to improve these indicators, India agreed to pursue the achievement of the MDGs by 2015. In India, 65% of births occur at home and one in seven was attended by medical personnel (Biswas, 2003).

A survey conducted in 2005-2006 showed that approximately 76% of women who gave at least a live birth received Antenatal care, and 74% did so from a skilled provider. It has been surveyed that three quarters of all maternal deaths occur during delivery and in the immediate post-partum period. Hence one of the most significant interventions for safe motherhood is to ensure skilled care provided by skilled professionals during pregnancy and childbirth. Although 75% of births were
reported to occur in rural areas, just about 37% of them were assisted by Skilled Birth Attendants (WHO 2008). In Andhra Pradesh, the current maternal mortality rate is 154 (2009) and the 2015 target is to reduce it below 100. The Each One save One (EOSO) program believes that no mother and child should lose their life in the natural process of childbirth. ESOO aims to reduce the incidence of maternal mortality in Andhra Pradesh by training student nurses on safe birthing practices. Nursing students were selected as they numbered more in the region and there is an acute shortfall of physicians (Prakasamma, M. 2009). Nursing professionals deal with maternity care and it is anticipated that by increasing their knowledge and skills of nurses, the availability of skilled birth attendants with safer birthing practices result in fewer maternal and neonatal deaths.

2. Literature Review

As more than 60% of births are domiciliary deliveries, India needs to come up with an option to provide skilled birth attendance at community level. Lack of qualified midwives is a major human resource constraint for providing locally accessible skilled delivery care for rural women. Any country with a political pledge to reducing maternal mortality has to focus on well-trained midwives in the hospital and the community. With the change in the role of ANMs and program priorities, comprehensive services have been ignored. The National Family Health Survey (2006) shows that only 52% of women receive three antenatal contacts and 42% receive any postnatal care.

In Andhra Pradesh (A.P), India, the incidence of maternal mortality is 154, compared to national level of 254. The major factors for the steadily high Maternal Mortality Rates are the low awareness among women and families about risks and low access to resources and services. Only 68.8% of women in Andhra Pradesh delivered in an institution (Prakasamma, M. 2009). The dysfunction of the public-health system at the grassroots level to put forward a skilled birth attendant for every pregnant woman is a vital factor in the existing high maternal mortality rate. There is a smaller amount of priority in training the traditional birth attendants, resulting in scarcity of skilled care in the community.

Under NRHM and RCH –II (2005-2010), the government of India is actively pursuing the goals of reduction of maternal mortality by focusing on the four major strategies of essential obstetric care and new born care to all, skilled attendance at birth, emergency obstetric care for those having complications and referral services (India country report, 2007).

The nurse’s community has to join hands with the government to come up with a long term plan of action to reduce maternal deaths through skilled midwifery services. Unless every PHC has a 24 hour availability of a qualified and skilled nurse midwife, childbirth assistance cannot become a reality in rural India (Regu, M., Tabish, S.A., 2002).

3. Description of the Intervention Venue

The Modern Government Maternity hospital is a 462 bedded Obstetrics and Gynecology Tertiary care Hospital. The hospital has an everyday patient load of 1000 patients. The hospital caters to pregnant women from all over the State of Andhra Pradesh, together with referrals of high risk cases from neighboring states of Karnataka and Maharashtra.

4. Description of Target Audience

The primary target group of the “Each One save One” (EOSO) program were student nurses studying in the Final year BSc. Program. ESOO aimed to reduce maternal mortality, by increasing the knowledge on safe birth practices among student nurses in Andhra Pradesh, and improve their ability to identify complications during labor and to take quick action. The nurses posted for their clinical rotations, at the Modern Government Maternity Hospital were requested to participate in weekly teaching programme for one month duration. The participants were from the Nizam Institute of Medical Sciences (NIMS) College and Laxmi College of Nursing in Hyderabad. A total of 61 nurses participated in the program, 54 were BSc. Nursing students, 5 were MSc. nursing students and two were nursing tutors. The student nurses consisted of both male and female nurses.

Material and Method

1. Program Description

The purpose of this program was to increase the nurse’s knowledge about safe childbirth practices. This intervention program involved teaching educational modules to student nurses from Bsc. Nursing final year. The program consisted of four educational training modules, which were taught once a week for length of approximately 75 minutes. The sessions were conducted in the auditorium at the Modern Maternity Government Hospital, Hyderabad. The medium in which the classes were conducted was English and each session was conducted by a health educator.

Four modules were developed keeping in line with the maternity cycle. In the first module, covered topics related to safe motherhood and antenatal care, both essential and trimester based care. In the second module the topics covered were, danger signs during pregnancy and labor and immediate management that a nurse can perform in case an emergency arises. In the third module, an elaborate description was given about stages of normal labor, mechanism of normal labor and identifying abnormal labor in terms of delay or arrest in progress of labor. A video on normal labor was shown to emphasize the stages of labor. In the last module, the main intent was to educate on the current maternal mortality status
in India and make them aware about the 5th millennium development goal of reducing maternal deaths. The main causes of maternal deaths in Andhra Pradesh (eclampsia and post partum hemorrhage) were discussed at length. A video on maternal death in India was shown to sensitize the nurses about the current maternal mortality scenario in rural areas.

During each session, brochures were distributed to the nurses. These brochures summarized essential points from each session. Outcome evaluation of the program was done with the help of, a pre test conducted prior to the start of the first training session and a post test at the end of the program. The pre test consisted of 15 objective types of questions which were made from topics covered in the four modules. The same questionnaire was used to conduct a post test at the end of the fourth training session, to assess if the program made any difference in knowledge among the student nurses. Another evaluation done was a post program survey. The questionnaire was developed to evaluate the entire program “Each One Save One Program”.

2. Evaluation Tools

To ensure a thorough evaluation of the program, the evaluation consisted of process, impact and outcome evaluations in the following stages:

Planning phase
- Process Evaluation: A needs assessment was done prior to implementing the program. It was done based on a thorough literature review. A lesson plan for each training module was developed, which helped the sessions proceed in an organized manner and made sure that the health educator covered all the necessary topics.

Implementation phase
- Process Evaluation: Sign in sheets were used to monitor attendance of the nurses. An evaluation form was distributed at the end of the last session to measure the quality of the training session. This method of verification can help improve the program.
- Impact Evaluation: Pre tests and were conducted before the first session and post tests were conducted at the end of the last session. This method of verification helped to look for any variation in knowledge among the nurses after the training sessions.

End of Program Phase
- Process Evaluation: The survey conducted at the end of the program was used to measure the quality of the entire program. This included teaching methods, handouts and there was a provision for nurses to comment/suggest how to improve the program.
- Impact Evaluation: A post test was given at the end of the last session, which contained same questions as in the pretest.

- Outcome/Program Evaluation: After conducting the pre and post tests, the results were evaluated based on the paired t-test, which showed there was a significant variation in knowledge after the sessions were completed. The mean, mode and median scores were calculated from the pretests and post tests.

Statistical Analysis

There were a total of 61 participants in the program. During the first session there were 25 participants, hence the initial round of pretest was conducted among these students. During the following week, another 24 new participants joined the program and they too took the pre test. There were a total of 49 pretest questionnaires completed. The participants, who did the pretest on a later date, were also taught the module that they missed.

The mean score on the pretest was 7 out of a total of 15. Total score on pretests for 49 students was 344. The median score was 7.25 and mode was 9.

There were a total of 39 students who attended the last session. Hence only these 39 students completed the post test questionnaire and post survey of the entire program. The mean score on the post test was 10.5 out of 15. Total score on post test for 39 students was 411. The median score was 10 and the mode obtained was 13 in the post test.

Table 1: Comparison of results between pretest and posttest

<table>
<thead>
<tr>
<th></th>
<th>Mean</th>
<th>Median</th>
<th>Mode</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-Test</td>
<td>7</td>
<td>7.25</td>
<td>9</td>
</tr>
<tr>
<td>Post-Test</td>
<td>10.5</td>
<td>10</td>
<td>13</td>
</tr>
</tbody>
</table>

The paired t-test was conducted on the students who did both the pre and post test, there were 35 such participants. The p value obtained after conducting the paired t-test was 1.46 X 10-10. The test was statistically significant (p < 0.05). The post test surveys were completed by 39 participants.

Result

The evaluation from pre and post test found an increase in knowledge among the nurses about safe birth practices. This was achieved as the mean score on pretest was 7, while mean score seen on post test was 10.5. This increase in score was significant (p< 0.05) as evaluated by a paired t-test, showing a significant increase in knowledge among the nurses. Another significant finding noted at the end of the program was that 91% of the nurses fared better as compared by the pre test and post test.

Discussion

The program EOSO was carried on from 1st July 2010 to 31st August 2010 on 61 student nurses of the final year Bsc
Nursing from NIMS College and Laxmi College of nursing. These nurses were trained on safe motherhood and safe birth practices which was designed into four educational training modules. The program was in collaboration with staff at the Indian Institute of Public Health, Hyderabad and Modern Government Maternity Hospital, Hyderabad.

The aim of the program was to facilitate reduction of maternal mortality in Andhra Pradesh which was at 154 per lakh live births at the time of intervention. In attempts to reduce maternal mortality, everyone in the health sector needs to act in unity to help reach the MDG of reducing maternal mortality rate to below 100 by 2015. Hence nursing students were chosen as the target population for this intervention, as they are the future nurse midwives and will work within the community.

This program was an effort made in reducing the maternal deaths by educating the nurses on practical aspects of pregnancy.

The program “Each One save One” can be seen as a pilot study, which was proven to be useful among the student nurses as the outcome evaluation showed a statistically significant result. This training model along with changes suggested by the participants can be used for training nurses in other hospital settings initially in Hyderabad and then in the rest of Andhra Pradesh.

**Conclusion**

Training of the nurses on safe birth practices and safe motherhood has led to an increase in knowledge among the nurses. Sixty one student nurses in their final year were trained on four interactive educational modules at the Modern Maternity Government Hospital, Hyderabad, India. Pre and post tests were conducted. The data obtained was compiled and statistically analyzed. The program found a significant increase in knowledge among the nurses after training on four modules. Hence it can be concluded that training nurses on safe motherhood and safe birth practices significantly increased their knowledge level.

**References**

Co-relation between chronic periodontitis and anemia – A pilot study

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Abstract

Aim
Anemia of chronic disease, a cytokine-mediated anemia, is a frequent complication of many chronic inflammatory conditions. The present case control study was aimed to evaluate levels of systemic hematological markers indicative of anemia in patients with generalized, severe, chronic periodontitis.

Methods
A convenience quota sample of 110 systemically-healthy, urban, male patients of a town Modinagar district Ghaziabad (42 Kms from New Delhi), comprised two groups, based on full mouth periodontal examination: group A patients (n = 50) were diagnosed with generalized, severe, chronic periodontitis, and group B patients comprised the control group (n = 50), which included patients with a clinically-healthy periodontium. Out of these 10 patients were eliminated due to diagnosis of systemic diseases. Laboratory blood investigations included hemoglobin (g%), total number of erythrocytes (red blood cells), hematocrit/packed cell volume, erythrocyte sedimentation rate, mean corpuscular volume of erythrocytes, and mean corpuscular hemoglobin concentration. An analysis was done to compare the mean values of hematological parameters within groups.

Results
The mean values of hemoglobin, red blood cells, packed cell volume, and mean corpuscular hemoglobin concentration were significantly lower, while the mean corpuscular volume of erythrocytes and erythrocyte sedimentation rate were significantly higher in group A patients compared to those in group B, indicating mild anemia.

Conclusions
Severe periodontal disease can be linked with anemic status.

Key Words
Anemia of Chronic disease, Periodontitis, Anemia.

Introduction
The initiation and progression of gingivitis and periodontitis may be affected by certain systemic conditions. The converse side of the relationship between systemic health and oral health has also been demonstrated. This means that there may be potential effects of periodontal disease on a wide range of organ systems. Anemia of chronic disease is defined as anemia occurring in chronic infections, inflammatory conditions or neoplastic disorders that are not due to marrow deficiencies or other diseases, and occurring despite presence of adequate iron stores and vitamins. Recently, Hutter has concluded that periodontitis, like other chronic conditions, lead to anemia.

Periodontitis, a chronic, infectious disease associated with gram-negative microorganisms that exist in a subgingival biofilm, is one of the most widespread diseases worldwide. The prevalence of periodontitis is reported as affecting 30–50% of the population, with approximately 10% having a severe form, and it is more prevalent in developing countries. Chronic periodontitis is the most common form of periodontal disease, which progresses relatively slowly and is more common in adults.

Anemia is defined as a state of reduced Hb concentration, reduced number of circulating erythrocytes in the blood, or both. Anemia is a common and serious health disorder among both sexes and all age groups, although it has a higher prevalence among women than men in India. Anemia of chronic disease (ACD) is a form of anemia that occurs in chronic infections and inflammatory conditions; it occurs in the presence of adequate iron stores and vitamins, and is not due to marrow deficiency.

Anemia of chronic disease (ACD) has been described in the literature, and seems to be one of the most common forms of anemia observed in clinical medicine. ACD is defined as the anemia occurring in chronic infections, chronic inflammatory processes or tumor formation that is not due to dysfunction of bone marrow cells or other diseases, and occurring despite the presence of adequate iron stores and vitamins. A characteristic finding of the disorders associated with ACD was the increased...
production of the cytokines that mediate the immune or inflammatory response; such as tumor necrosis factor, interleukin-1, and the interferon. All the processes involved in the development of ACD can be attributed to these cytokines, including shortened red cell survival, blunted erythropoietin response to anemia, impaired erythroid colony formation in response to erythropoietin, and abnormal mobilization of reticuloendothelial iron stores. These cytokines are also released by periodontal tissues in response to bacterial infection, which suggests that periodontitis like other chronic disease may cause ACD.

However, conflicting results have been reported regarding the association of periodontal disease and anemia. The purpose of this pilot study was to compare the hematological parameters related to anemia in male patients with severe periodontal disease with that of periodontally healthy male patients, and thereby evaluate a possible association between severe periodontal disease and anemia.

Material and Methods

A convenience quota sample of 110 adult, male patients in the age range of 30–60 years, who reported to the Department of Periodontics and Oral Implantology in D.J. College of Dental Science & Research, India, were enrolled in the study. Out of these 10 patients were excluded due to detection of systemic diseases. The study protocol was approved by the institutional ethical committee. Informed consent was obtained from patients. Demographic and clinical data were collected by means of a questionnaire, survey of any previous medical records, and oral examination.

Inclusion criteria included presence of at least 20 teeth, body mass index of 18.5–25, and self-reported consumption frequency of at least four times per week of food from the following groups: milk or curd; pulses; fruits; dark green, leafy vegetables; eggs; chicken or meat; and fish. Exclusion criteria were a self-reported intake of vitamins and iron supplementation or any anti-inflammatory or antimicrobial drug within the previous 3 months; a self-reported history of acute or chronic medical conditions, including diabetes or viral, fungal, or bacterial infections; smokers or smokeless tobacco or betel (areca) nut and betel quid (paan) consumers; alcohol consumers; past periodontal treatment; or a recent history of trauma or tooth extractions. A full mouth periodontal examination was conducted (Figure 1).

Clinical periodontal parameters were recorded with a Williams graduated probe at six sites on each tooth for both case and control groups. These were probing depth (PD), clinical attachment loss (CAL), and bleeding on probing (BOP), as recorded on visual examination after 30–60 sec of probing. Based on the clinical examination, selected patients were grouped into case and control groups.

Fifty patients with at least 30% of sites with a CAL of 5 mm or more, along with 10% or greater number of sites with a PD of 6 mm or more were diagnosed with generalized, severe, chronic periodontitis. These patients made up group A. Fifty patients with 0% of sites with CAL were considered periodontally healthy and made up the control group, group B.

Blood collection and analysis

10 mL venous blood samples were collected by venepuncture under aseptic conditions in the antecubital fossa without excessive venous stasis between 9.00 and 12.00 hours for both case and control group patients. The blood was transferred into ethylenediaminetetraacetic acid containing bulbs. Centrifugation of blood using REMI R-8C laboratory centrifuge for measuring serum ferritin was assessed. (Figure 2)

The hematological parameters assessed in the present study were hemoglobin (Hb) (in g% by Sahli’s method), (Figure 2) total number of erythrocytes (red blood

Figure 1: CPITN probe used for measuring the pocket depth and clinical attachment level at baseline

Figure 2: Centrifugation of blood using REMI R-8C laboratory centrifuge for measuring serum ferritin
cells (RBC) (in million/mm³ by a Neubauer counting chamber), hematocrit/packed cell volume (PCV) (in mm by Wintrobe’s tube centrifuge method) (Figure 3), erythrocyte sedimentation rate (ESR) (in mm by Wintrobe’s tube centrifuge method), mean corpuscular volume of erythrocytes (MCV) (in cu µm), and mean corpuscular hemoglobin concentration (MCHC) (in %) (Figure 4). Student’s t-test and an analysis of covariance for age adjustment were performed with appropriate Microsoft Excel software for a comparative analysis of outcome parameters between groups A and B.

Results
The mean age of group A, which was 53.46 (±7.4) years, was significantly higher than that of control group B, which was 42.13 (±5.9) years. Thus, the analysis of covariance for age adjustment was applied to compare outcome parameters. The mean values of Hb and RBC were significantly lower in group A (11.19±1.97 g/dL and 4.81 ± .392 million/mm³) compared to group B (16±1.8 g/dL and 6.09±.30 million/ mm³) (P < 0.001). Similarly, the mean values for the PCV and MCHC were significantly lower in group A being 38.71±4.22 % and 32.88±2.1 %, respectively) compared to those recorded for group B 44.9±3.7 % and 37.11± 1.2 %, respectively (P < 0.001). The mean MCV in group A 97.51±7.29 cu µm) was significantly higher than that of group B 88.5±3.03 cu µm). The mean value of ESR was also noted to be higher in group A (19.71±9.82 mm) compared to group B (3.72±9.21 mm), which was statistically significant (P < 0.001) (Table 2). Even after adjusting for the age difference between both groups by the analysis of covariance, the differences between both groups remained statistically significant.

Discussion
Anemia of chronic disease is the second most prevalent form of anemia after nutritional, iron-deficiency anemia, and can coexist together, causing additional anemic burden. ACD is a cytokine-mediated anemia characterized by hypoferremia, with adequate reticuloendotelial iron stores and normal-to-elevated ferritin concentrations. It is a known, frequent complication of chronic inflammatory conditions such as rheumatoid arthritis. The pathogenesis is reported to be dysregulation of iron homeostasis, depressed erythropoiesis, and a blunted erythropoietin response caused by elevated levels of systemically-circulating pro-inflammatory cytokines resulting due to a local chronic inflammatory process.

Various studies have tried to evaluate the relationship between periodontitis and hemoglobin. Hutter et al. and Thomas et al. found that periodontitis patients have lower hematocrit, lower numbers of erythrocytes, lower hemoglobin levels and higher erythrocyte sedimentation rates when compared to healthy controls. Rai and Kharb found an increased in hemoglobin and RBC levels in patients with severe periodontitis after scaling and root planning. Considering the relatively high prevalence of anemia, as well as periodontal disease, in Indians, determining the etiological contribution of periodontitis to the presence of an anemic status assumes clinical significance.

Table I: Mean & SD and student ‘t’ test significance values for systemic Parameters within groups. (n=50)

<table>
<thead>
<tr>
<th>PARAMETERS</th>
<th>GRP. (A) mean (±SD) (n= 50)</th>
<th>GRP. (B) mean (±SD) (n=50)</th>
<th>P-VALUE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemoglobin</td>
<td>11.19±1.97</td>
<td>16±1.8</td>
<td>.0007*</td>
</tr>
<tr>
<td>RBC</td>
<td>4.81 ± .392</td>
<td>6.09±.30</td>
<td>.00002*</td>
</tr>
<tr>
<td>Packed cell volume</td>
<td>38.71±4.22</td>
<td>44.9±3.7</td>
<td>.00038*</td>
</tr>
<tr>
<td>Erythrocyte sedimentation rate</td>
<td>19.71±9.82</td>
<td>3.72±.921</td>
<td>.00000*</td>
</tr>
<tr>
<td>Mean corpus volume</td>
<td>97.51±7.29</td>
<td>88.5±3.03</td>
<td>.00001*</td>
</tr>
<tr>
<td>Mean corpus hemoglobin concentration or (P&lt;0.05)</td>
<td>32.88±2.1</td>
<td>37.11±1.2</td>
<td>.0005*</td>
</tr>
</tbody>
</table>

*P<.001 snows a high significant different between group A & B
significance. Smoking tobacco alters systemic cytokine levels and contributes to anemia, as well as increased periodontal destruction.

In our study, MCV levels in both groups were within the reference values, indicating normocytic anemia in group A, as commonly seen in ACD. As MCV levels greater than 95 have been reported as less likely to indicate iron deficiency anemia, and are suggestive of ACD with high predictability, we can attribute the anemia in group A as ACD. The normal range of MCHC is 28–33 g%, and the means for both groups fell within this range, although group A showed a significantly lower mean MCHC, indicating mildly hypochromic anemia. While hypochromia is commonly suggestive of iron deficiency anemia, a mildly hypochromic ACD has been reported in 30–40% of cases.

Although the RBC indices noted in our study are suggestive of ACD, an analysis of serum ferritin levels and the soluble serum transferrin receptor concentration, or a bone marrow examination, would be necessary to quantify iron stores and definitively distinguish between ACD, an analysis of serum ferritin levels and the soluble transferrin receptor concentration, or a bone marrow examination, would be necessary to quantify iron stores and definitively distinguish between ACD, anemia of chronic disease. Additionally, within the limits of the study, our results indicate a possible effect of severe periodontal inflammation on the severity of anemia, and we suggest that periodontal therapy might have a potential role in improving anemic status in periodontally-diseased individuals.

Long term, multicenter, prospective studies are essential to verify the magnitude and clinical relevance of this potential therapeutic effect.

Conclusion

Generalized, severe, chronic periodontal disease is positively associated with decreased levels of Hb, hematocrit, and RBC counts, along with increased ESR and MCV levels, suggesting that mild ACD is induced by the systemic effects of periodontal inflammation in patients with severe periodontal disease.

References

A Study on perceptions of key health care staff towards disease outbreak
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Abstract
To understand health care workers' perceptions towards a disease outbreak, a survey was conducted in 5 districts of Karnataka, India between August 2009 and December 2009, to understand the factors that may influence their ability and willingness to report to duty in such an event. The data suggest that nearly half of health care workers are not likely to report to duty during an outbreak. The stated likelihood of reporting to duty was significantly greater for clinical than technical and support staff, and perception of the importance of one's role in the district's overall response was the single most influential factor associated with willingness to report for work. The perceived risk among the health care workers was shown to be associated with several factors peripheral to the actual hazard of this event. Lacunae in knowledge identified serve as barriers to outbreak response and must be specifically addressed to enable effective local public health response to this significant threat.

Key Words
Perceptions, Outbreak, health care staff.

Introduction
The health and the medical education departments are considered the backbone of public health response plans for any and all disease outbreaks and emergencies. Emerging Infectious disease outbreaks are considered increasingly likely, and are now considered one of the most significant and urgent threats to India's public health preparedness infrastructure. The threat of emerging infectious diseases is a product of the globalization process. Changing lifestyles, patterns of behaviour and several such complex factors have led to the emergence and spread of disease in India. Outbreaks of diseases like SARS, Japanese encephalitis, dengue, chickungunya, malaria, bird flu, etc., in recent times, have critically influenced human lives in India. Added to this, is the recent pandemic of swine flu. A deliberate or a natural outbreak of disease would force the target state to mobilize the government machinery, human and financial resources while combating the disease1.

The recent outbreak of swine flu in India has brought to light deficiencies that exist regarding disease preparedness, prevention and lifestyle in India. The numbers of cases have been on a rise since May 2009 and many people have been affected2. Successful containment relies on effective system for outbreak detection, rapid data collection, analysis, assessment and timely reporting3. These activities would require an extensive prompt response by local health departments. Previous studies have shown that during extreme scenarios, a varying proportion of healthcare workers may be unable or unwilling to report to duty4-6. Data from even the developed countries like the United States indicate inconsistent and sometimes slow after-hours response by health departments to urgent events involving communicable disease7.

Risk perception theory provides a revealing framework for better understanding response limitations and needs of the public health workforce. The perceived risk, according to this theory, is a multifactorial phenomenon, involving the summation of actual risk and other peripheral influences independent of the actual risk, such as perceived authority, trust, and situational control; these peripheral influences have been termed “outrage” or “dread.”[8]. Based on these models, it was previously suggested that contributing factors peripheral to the actual risk will have a considerable practical impact on how the health care employees would respond in a crisis9.

Aside from physical and circumstantial barriers such as availability of transportation or dependency of family members, specific risk perception issues whose impact may be markedly high and of unique importance for the public health workforce's response to a crisis were identified. These factors arise from a number of features previously suggested to have been associated with elevated risk perception, including manageability of the threat; risk to future generations; direct personal impact; and sense of control over events.

Based on these factors, several major barriers to effective health workforce emergency response were suggested; these include uncertainty regarding working environment safety, unclear expectations of role-specific emergency response requirements, safety and well being of family members, inadequate emphasis on the critical value of each employee to the district’s response efforts, and insufficient emphasis on stress management techniques – all of which may heighten employees' sense of dread due to a lack of personal control9.
So we decided to assess health care personnel’s risk perception and likelihood of reporting to duty during a local disease outbreak, and to uncover the variables that affect these outcomes, thus providing a needed evidence base for health departments’ planning and training efforts.

**Methodology**

We conducted the study in 5 districts of Karnataka, India namely Shimoga, Chitradurga, Tumkur, Hassan and Kolar between August 2009 and December 2009. All the key health care personnel were identified both from the Health and Medical Education in these districts. Self-administered anonymous survey questionnaires were sent to all the identified personnel by us through proper channels. Completed questionnaires were directly mailed to the chief investigator at the government medical college, Shimoga.

The questionnaire included questions on personal characteristics such as age, sex and job classification. Questions were typed in the local language (Kannada) as well for getting accurate responses from the support staff. The health care personnel used a 5-point Likert scale for questions pertaining to a possible disease outbreak in the locality.

The job classification variable was categorized into technical/support staff and professional staff. The former included lab technicians, pharmacists, computer operators, data managers and drivers. The professional staff included district health officials, program managers, general duty medical officers, clinical staff in the medical college (e.g., physicians, nurses), public health communicable disease staff, public information staff, and other public health professional staff (e.g., health educators, LHVs, etc)

The responses to the job classification question were categorized as professional and technical/support groups. Questions about likelihood of reporting to work and disease outbreak related attitudes and beliefs were dichotomized into responses with a score two or less, and all other responses. Logistic regression was used to compute Odds Ratios to evaluate the association of demographic variables and attitudes and beliefs with self-described likelihood of reporting to work. Multivariate logistic regression was used to explore associations between attitudes and beliefs related to outbreak preparedness and self-described likelihood of reporting to work. Adjustment for age, sex, and job classification was done. Likewise, bivariate and multivariate (adjusted for age, sex, and job classification) logistic regression models were used to evaluate the association between the various attitudes and beliefs. In order to assess non-response bias, age, sex, and job classification distributions for the respondents and for all health care personnel were compared. SPSS version 14 was used to analyze the data.

**Results**

The overall response rate was 77% (n=616). The breakup of the responses from the 5 districts is shown in Table-1. Statistically significant difference in age and gender distribution between the respondents and all health department personnel was not found. A small yet statistically significant difference in the proportion of technical/support staff (vs. professional staff) was detected (22.4% vs. 32% in the study group and all personnel respectively, p = 0.003), yet no significant difference in the proportions of professional staff subgroups was detected.

Of the 606 who responded to the question about their likelihood of reporting during an outbreak related emergency, 326 (53.8%) indicated they would likely report to work during such an emergency. Age and sex did not have an association with likelihood of reporting. Professional staff indicated a higher likelihood of reporting (Multivariate OR: 2.5; CI 1.3–4.7) than technical/support staff (Table -2).

Only 40% of all respondents- 45.1% professional staff and 26.1% technical/support staff – felt it was likely they would be asked by their departments to respond to an outbreak related crisis. Perception of likely to be asked by the department to respond was associated with self-described likelihood of reporting (Multivariate OR: 8.5; 95%; CI 4.6–15.6). Only 33.4% (202) individuals thought of themselves to be knowledgeable about the public health impact of an epidemic (Table -3).

Perception of one’s existing knowledge about disease outbreak, and perception of having an important role in the district’s overall response were significantly higher among professional staff compared to technical/support staff (Figure -1).

In multivariate analysis, increased self-described likelihood of reporting to work during an influenza pandemic emergency was significantly associated with agreement with several constructs, most notably perception of the capacity to communicate risk effectively, perception of the importance of one’s role in the agency’s overall response, and familiarity with one’s role-specific response requirements in a pandemic influenza related emergency (Table -3).

The vast majority (83%) of the respondents felt they would benefit from additional training activities. A lower
perceived level of familiarity with one's role was not significantly associated with a higher perceived need for additional training (Multivariate OR: 1.4; CI 0.6–3.4). Most of the respondents also perceive psychological support during the event (57.1%) and post-event psychological support (61.3%) as important. Psychological support during and after the event was deemed more important by staff who considered themselves likely to be asked to report to duty during such an extreme public health crisis. In fact, most of the workers (and nearly three out of four technical/support workers) do not believe they will even be asked to report to work.

66% of the respondents perceived themselves to be at personal risk when performing their duties during such an event. Confidence in personal safety was associated with several constructs independently of one's job classification, including perception of existing knowledge about public health impact of disease outbreak (Multivariate OR: 4.1; CI 2.3–7.6); family preparation (Multivariate OR: 2.5; CI 1.4–4.3); department's perceived ability to provide timely information (Multivariate OR: 4.8; CI 2.6–9.0); perception of the capacity to effectively communicate risk (Multivariate OR: 4.1; CI 2.9–7.7); and familiarity with one's role-specific response requirements (Multivariate OR: 3.5; CI 1.8–6.2). The associations between self-identified likelihood of reporting to work and perception of one's capacity to effectively communicate risk were substantially stronger for technical/support staff compared to professional staff (Bivariate OR: 19.4; CI 2.4–160.4 vs. OR: 5.9 CI 2.9–12.2) respectively.

Discussion

Experience has shown, localized “hot spots” of increasing transmission can continue to occur even when the pandemic has peaked at the national level [10]. Existing plans account for personnel shortages within the healthcare settings. Our data suggest that, nearly half of the local health department workers are likely not to report to duty during such an extreme public health crisis. In fact, most of the workers (and nearly three out of four technical/support workers) do not believe they will even be asked to report to work.

It was found that the willingness to report to duty during an outbreak varies considerably according to the individual's job classification. Clinical staff state they are significantly more likely to report to duty, compared with all other workers. This difference correlates well with the single most influential construct associated with willingness to report to duty – the perception of the importance of one's role in the district's overall response. Less than a third of the respondents believed they will have an important role in the district's response to local outbreaks of an infectious disease, but within this subgroup, willingness to report to duty was as high as

Table 2: Demographic profile of the health care personnel

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>n (%)</th>
<th>Bivariate OR (95% CI)</th>
<th>Multivariate OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yrs)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20 - 25</td>
<td>40 (6.6)</td>
<td>Reference</td>
<td>Reference</td>
</tr>
<tr>
<td>25 - 30</td>
<td>96 (15.8)</td>
<td>1.2 (0.4–3.4)</td>
<td>0.9 (0.3–2.8)</td>
</tr>
<tr>
<td>30 - 35</td>
<td>204 (33.6)</td>
<td>1.3 (0.5–3.3)</td>
<td>0.8 (0.3–2.5)</td>
</tr>
<tr>
<td>35 - 40</td>
<td>214 (35.2)</td>
<td>1.3 (0.5–3.3)</td>
<td>0.9 (0.3–2.5)</td>
</tr>
<tr>
<td>&gt;40</td>
<td>54 (8.9)</td>
<td>0.9 (0.3–3)</td>
<td>0.5 (0.1–1.9)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>118 (19.1)</td>
<td>Reference</td>
<td>Reference</td>
</tr>
<tr>
<td>Male</td>
<td>498 (80.9)</td>
<td>0.7 (0.4–1.4)</td>
<td>0.6 (0.3–1.2)</td>
</tr>
<tr>
<td>Job</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Technical/Support Staff</td>
<td>138 (22.4)</td>
<td>Reference</td>
<td>Reference</td>
</tr>
<tr>
<td>Public Health Officials</td>
<td>14 (2.3)</td>
<td>2.6 (0.5–14.2)</td>
<td>1.9 (0.3–11)</td>
</tr>
<tr>
<td>General Duty Medical Officers</td>
<td>204 (33.1)</td>
<td>2.3 (1.2–4.4)</td>
<td>2.5 (1.3–4.7)</td>
</tr>
<tr>
<td>Clinical Staff in medical colleges</td>
<td>78 (12.7)</td>
<td>0.9 (0.4–1.9)</td>
<td>0.6 (0.2–1.4)</td>
</tr>
<tr>
<td>Program Managers</td>
<td>24 (3.9)</td>
<td>3.1 (0.8–12.4)</td>
<td>3 (0.7–12.1)</td>
</tr>
<tr>
<td>Public Information Staff</td>
<td>16 (2.6)</td>
<td>0.6 (0.2–1.4)</td>
<td>0.4 (0.1–1.9)</td>
</tr>
<tr>
<td>Other Public Health Staff</td>
<td>142 (23.1)</td>
<td>0.7 (0.3–1.3)</td>
<td>0.7 (0.3–1.3)</td>
</tr>
</tbody>
</table>

Table 3: Association of perceptions regarding disease outbreak preparedness with projected likelihood of reporting to duty

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Construct agreement =n</th>
<th>Bivariate OR (95% CI)</th>
<th>Multivariate Model OR (95%CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Perceived importance of training and education</td>
<td>508</td>
<td>3.8 (1.9–7.5)</td>
<td>3.4 (1.6–7.1)</td>
</tr>
<tr>
<td>Perception of the capacity to effectively communicate risk</td>
<td>160</td>
<td>7.1 (3.6–13.9)</td>
<td>6.6 (3.2–13.5)</td>
</tr>
<tr>
<td>Department's perceived ability to provide timely information</td>
<td>390</td>
<td>2.4 (1.5–3.8)</td>
<td>2.3 (1.3–3.8)</td>
</tr>
<tr>
<td>Perception of existing knowledge about public health impact of an outbreak</td>
<td>202</td>
<td>3.5 (2.1–5.9)</td>
<td>3.1 (1.8–5.5)</td>
</tr>
<tr>
<td>Familiarity with one's role-specific response requirements</td>
<td>142</td>
<td>7.2 (3.5–14.7)</td>
<td>7.6 (3.4–16.9)</td>
</tr>
<tr>
<td>Perception of the importance of one's role in the district's overall response</td>
<td>186</td>
<td>10.4 (5.3–20.3)</td>
<td>9.5 (4.6–19.9)</td>
</tr>
<tr>
<td>Family preparation</td>
<td>310</td>
<td>2.4 (1.5–3.8)</td>
<td>2.1 (1.2–3.4)</td>
</tr>
<tr>
<td>Confidence in personal safety</td>
<td>200</td>
<td>4.4 (2.6–7.6)</td>
<td>4 (2.2–7.2)</td>
</tr>
</tbody>
</table>
Belief in the importance of one's role was lowest among technical/support staff, program, and other non-clinical professional staff (15.1%, 18.4% and 18.8% respectively), groups in which willingness to report was shown to be lowest. Therefore we feel it is important that further efforts must be directed at ensuring that all local public health workers, but most notably non-clinical professional staff, understand in advance the importance of their role during an outbreak – otherwise they will fail to show up when they are most needed.

Lack of knowledge, ambiguity regarding one's exact tasks, and questionable ability in performing one's role as risk communicator were all significantly associated with a higher perceived personal risk and a two- to ten-fold decrease in willingness to report to duty; these factors proved to be more influential even than the perceived level of family preparedness to function in one's absence. It is therefore important to recognize that public health employees, who are intended to serve as purveyors of risk communication for their communities, themselves represent a community with specific perceptions that must be addressed in the context of emergency readiness training. It was only in the last one year; H1N1 strain became increasingly pandemic in Southeast Asia and as lethal infections with the virus occurred in an alarmingly increasing rate among humans, that the urgency of the situation was openly declared by national and international health authorities. The rapidity of this evolving situation may serve to explain why only one third of the respondents felt they were adequately knowledgeable on outbreaks, and why only one in five respondents felt capable in effectively communicating pandemic risks. Only 2 of the 78 technical/support staff workers who felt incapable of effective risk communication was willing to report to duty, even though most of them believed the health department will have the ability to provide timely information.

The study has some important limitations that must be factored into the overall analysis. First, the sample was limited to five non-randomly selected districts, and all of which have staff sizes under 300. The job classifications do not necessarily map neatly onto functional responsibilities in disaster response. For example, health educators may play as frontline a role as clinical staff, in terms of their degree of interface with the public in a disaster. The lack of significant difference in age and gender distribution, as well as the lack of significant difference in job classification other than technical/support staff indicates that the extent of such a bias in the study is probably limited.

This study show similar patterns to data on the willingness of urban healthcare workers from non-public health settings to respond to emergencies: a survey of 6248 employees from 47 healthcare facilities in the New York City area revealed that these workers were least willing (48%) to report to duty during an untreatable naturally-occurring infectious disease outbreak affecting their facility (SARS), compared to other disaster scenarios [6]. In the face of a pandemic H1N1 influenza threat, local health department employees' unwillingness to report to duty may pose a threat to the nation's emergency response infrastructure. Addressing the specific factors associated with this unwillingness is necessary to help ensure that existing local health department preparedness competencies

**Conclusion**

Most of the health care personnel feel they work under significant personal risk, in a scenario they are not adequately knowledgeable about, performing a role they are not sufficiently trained for, and believing this role does not have a significant impact on the district's overall response. These specific perceptions and needs must be attended, and specific intervention programs must be initiated. In order to reduce the perceived risk associated with the worker's role in a case of a disease outbreak, each worker must have better understanding of the scenario and importance of his or her personal role within these settings, confidence that the district will provide adequate protective equipment for its employees, psychological support and timely information, and a belief of being well-trained to cope with emergency responsibilities including the ability to communicate risk to others.

**Acknowledgement**

The authors are grateful to all the respondents who took part in the study

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Prevalence of ABO and Rhesus blood groups among blood donors

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Abstract

Background
ABO and Rhesus (Rh) blood group antigens are hereditary characters and are useful in population genetic studies, in resolving medico-legal issues and more importantly for the immunologic safety of blood during transfusion.

Objectives
To document the distribution pattern of the ABO and Rh blood groups among blood donors in and around Davangere (Karnataka).

Methods
The distribution of ABO and Rh blood group was analyzed among 19,413 blood donors, over a period of 5 years (2005 to 2009). The age group and sex of donors, frequency of ABO blood groups and Rh status were calculated.

Results
The predominant donors belonged to the age group between 18-35 years (86.18%). Male to female ratio among donors was 86:1. The most prevalent blood group was O (36.76%), followed by group B (29.85%) and group A (26.15%). The least common blood group was AB (7.24%). The prevalence of Rh positive and negative distribution in the studied population was 94.48 and 5.52% respectively. The highest frequency of coexisting ABO-Rh phenotypes was that of O positive (34.67%) followed by B positive (29.85%) and A positive (26.15%).

Conclusion: Knowledge of frequencies of the different blood groups is very important for blood banks and transfusion service policies that could contribute significantly to the National Health System.

Key Words
Blood groups, ABO, Rhesus, phenotypes, blood donors.

Introduction
Over 700 erythrocyte antigens have been reported in the literature and have been organized into 30 blood group systems by the International Society of Blood Transfusion. The two most significant blood group systems were discovered by Karl Landsteiner during early experiments with blood transfusion: the ABO group in 1901 and in cooperation with Alexander Wiener the Rhesus group in 1937. The discovery of the ABO system marked the beginning of modern blood banking and transfusion medicine. The ABO system is the single most important blood groups which hold a respectable position in view of the safety of blood/blood component transfusion to date.

Blood groups are genetically determined and exhibit polymorphism in different populations. The knowledge of the distribution of ABO and rhesus (Rh) blood group is essential for effective management of blood banks inventory, be it a facility of a smaller local transfusion service or a regional or national transfusion service. Apart from their importance in blood transfusion practice, ABO and Rh blood groups are useful in population genetic studies, researching population migration patterns, as well as resolving certain medico-legal issues, particularly of disputed parentage. It is, therefore, imperative to have information on the distribution of these blood groups in any population.

The frequencies of ABO and Rh blood groups vary from one population to another. There are very few documented works devoted to the study blood groups in India and no data is available for Davangere, Karnataka. Our aim was to determine the ABO and Rh blood group distribution pattern among blood donors in a tertiary care hospital and to compare our results with other studies in India and elsewhere.

Material and Methods
ABO and Rh blood group of 19,413 blood donors (includes both voluntary and replacement donors) who donated blood for various reasons at the S S Blood Bank, Davangere, were analyzed from the inception of blood bank in 2005 to 2009 (five years). Donors were also recruited from various blood donations camps held in association with Indian Red Cross society and district hospital Davangere. The blood group phenotypes were detected by the classic slide and tube method by the antigen-antibody agglutination test with appropriate positive and negative control by mixing whole blood
with appropriate anti-sera. In case of doubt, the test was examined under a microscope or the results were confirmed by reverse grouping using known group A and B red cells. Results of donors’ age, male to female ratio and frequency of ABO and Rh blood groups were analyzed.

Results

A total of 19,413 blood donor’s blood groupings were done. The donors were aged 18-60 years with the maximum donors between 18-35 years (86.18%) [Table 1]. Male donors (19,189, 98.85%) were more than females (224, 1.15%) with male to female ratio of 86:1. The number of voluntary and replacement donors were 1197 (6.17%) and 18,216 (93.83%) respectively.

Table 1: Age distribution of blood donors

<table>
<thead>
<tr>
<th>Age Group (Years)</th>
<th>No. of Donors</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>18-25</td>
<td>8950</td>
<td>46.11</td>
</tr>
<tr>
<td>26-30</td>
<td>5119</td>
<td>26.37</td>
</tr>
<tr>
<td>31-35</td>
<td>2659</td>
<td>13.70</td>
</tr>
<tr>
<td>36-40</td>
<td>1562</td>
<td>8.05</td>
</tr>
<tr>
<td>41-45</td>
<td>691</td>
<td>3.55</td>
</tr>
<tr>
<td>46-50</td>
<td>336</td>
<td>1.73</td>
</tr>
<tr>
<td>51-55</td>
<td>78</td>
<td>0.40</td>
</tr>
<tr>
<td>56-60</td>
<td>18</td>
<td>0.09</td>
</tr>
<tr>
<td>Total</td>
<td>19,413</td>
<td>100</td>
</tr>
</tbody>
</table>

The most common ABO blood group was found to be group O (7133, 36.76%), followed by group B (5795, 29.85%) and group A (5078, 26.15%). The least common blood group was AB group (1407, 7.24%) i.e., O > B > A > AB. Rh antigen was detected in 18,339 (94.48%) donors while Rh negative phenotype was found in 1074 (5.52%) donors [Table 2 and Figure 1].

Rh-D negative was common to the blood group O (2.09% of all blood groups) as compared to blood group of AB (0.30%). The Rh group negativity for blood groups A and B were close (1.54% and 1.59% respectively). There was no association between ABO blood group and Rh status.

Discussion

Both ABO and Rh blood grouping is very important for safe blood transfusion. Dangerous hemolytic transfusion reactions may occur when blood is transfused into an individual with an incompatible blood type. The severity of the resulting transfusion reaction may vary from an asymptomatic minor rise in the plasma bilirubin level to severe jaundice and renal tubular damage, with anuria and death. Another complication due to “Rh incompatibility” arises when an Rh negative mother carries an Rh positive fetus (erythroblastosis fetalis).

ABO and Rh genes and phenotypes vary widely across races and geographical boundaries despite the fact that the antigens involved are stable throughout life. The resultant polymorphism remains important in population genetic studies, estimating the availability of compatible blood, evaluating the probability of hemolytic disease in the newborn, resolving disputes in paternity/maternity and for forensic purposes.

In our study, maximum donors were between 18-35 years because young adults usually volunteer to donate blood than older age group. It was also evident that numbers of female donors were less because of their low body weight, low hemoglobin concentration, ignorance, lack of motivation and awareness and fear among the females regarding blood donation. The similar findings were noted by Ahad et al. Hence, general population needs more motivation and many voluntary blood donation camps need to be conducted to meet the increasing demand for blood.

We have compared the frequency of distribution of blood groups ABO and Rh to previous studies. The present study is useful in providing information about the status of ABO and Rh blood group among blood donors which might represent ABO and Rh frequency in general population.

As far as distribution of ABO blood group is concerned, the group O is the most frequently encountered phenotype in the population under study. This observation is in accordance with previous reports from other parts of the world.
South India 14,15,18. Another south Indian study conducted on the population of Chittoor district of Andhra Pradesh also showed similar pattern of distribution of blood groups 17.

With regard to the other ABO blood group phenotypes, the frequency of group 'A' in the present study is in proximity to the reported frequency among the Bangaloreans 14. Slightly higher frequency for group A was reported in Gangadikara Vokkaligas of Mysore 15. A proportion of group 'B' as in our study is similar to report by other studies from South India 14,18 whereas in north India blood group 'B' was found to be commonest ABO subtype 19,20. The discrepancy between our findings and that reported by Wadhwa et al 19 and Sidhu 20 may be attributed to the ethnic difference among the population of India, or it could be due to the smaller sample size as against a relatively larger sample size in the present study. A Study done by Nanu and Thapliyal 29 also reports that group B is the most predominant ABO group in the north Indian population as also in neighboring Pakistan 21. Bombay group was not found in our study.

The frequency of Rh negative phenotype is close to those reported from other parts of South India 14,18 and north-India 19,20. This indicates that frequency of Rh-D negative phenotype in India is around 5% in sharp contrast to the frequency of about 15% phenotype reported in other nations 6,11.

**Conclusion**

This study establishes that there is a significant difference in ABO blood groups between south and north India, and India as well as different parts of the world. Among blood donors, the various ABO and Rh blood groups in south India, group O is the commonest, followed by blood groups B and A, where as in north India group B is the commonest phenotype. The frequency of Rh-negative in India is similar to other series. The results of this study could contribute significantly to the National Health System in aiding the prediction of percussions of certain diseases related to blood groups, as well as the requirement for certain blood groups within the blood donation programme. Knowledge of the frequencies of the different blood groups in this part of India is very important for blood banks and transfusion service policies. Knowledge of blood group phenotype distribution is also important for clinical studies (for example disease association), as well as for population studies. Thinking about the life threatening complications and dangerous sequel of transfusion reaction we should be very careful regarding ABO and Rh blood grouping starting from blood collection, blood grouping and cross matching and up to transfusion to the recipient. Only sincerity and awareness can bring these dangerous transfusion reactions to zero.
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Comparative study of lipid profile in chronic smokers with and without acute myocardial infarction

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Abstract

Background
Smoking is one of the major risk factors in the genesis of coronary atherosclerosis and development of coronary heart disease. Smoking may alter normal plasma lipoprotein levels.

Objective
The present study was undertaken to compare the lipid profile between chronic smokers (Group A) and chronic smokers with myocardial infarction (Group B).

Material and Methods
Fifty apparently healthy chronic smokers and 50 smokers with myocardial infarction were selected for the study. Fasting venous blood sample was collected in controls, in AMI patients sample was collected before any intervention, and total cholesterol (TC), triglycerides (TG), and high density lipoprotein cholesterol (HDL-C) were measured. Low density lipoprotein cholesterol (LDL-C) and Very low density lipoprotein cholesterol (VLDL-C) were calculated by Friedwald’s formula. Statistical analysis was done by Z test.

Results
The lipid profile was compared between Group A and Group B.
There was a significant rise in TC, TG, LDL-C, VLDL-C and significant decrease in HDL-C in Group B as compared to Group A.

Conclusion
Cigarette smoking modulates the IHD risk through gene-environment interaction and smoking increases the risk of atherosis. Further studies are required to ascertain the gene environment interaction.

Key Words
Chronic smokers, Myocardial infarction, Lipid profile.

Introduction
Smoking is among the 10 greatest risk to health¹. Cigarette smoking done more than 10 per day for 6-8 years is strongly implicated risk factor and also leading cause of death for chronic obstructive pulmonary disease(COPD), lung cancer and atherosclerosis².

There is a mechanistic link between increased levels of oxidative stress associated with smoking and the onset and progression of cardiovascular diseases. Two major phases were identified in cigarette smoker: A tar phase and a gas phase. The tar phase of cigarette contains semiquinone radical which can react with oxygen to produce superoxide anion, hydroxyl radical and hydrogen peroxide. The gas phase contains alkyl and peroxy type. In one puff of cigarette, the smoker is exposed to more than 1015 free radicals in gas phase alone, with addition exposure in tar phase equal to more than 1017 free radicals per gram³.

Smoking is associated with increased of lipid, protein and DNA oxidation. Increased oxidative stress and conversion of LDL to oxidized LDL could lead to atherosclerotic lesions. Circulating products of lipid peroxidation and autoantibody titers to oxidized LDL are significantly increased in smokers. It has been described that nicotine increases the circulatory pool of atherogenic LDL via accelerated transfer of lipids from HDL and impaired clearance of LDL from plasma compartment. Therefore, it increases the deposition of LDL-C in the arterial wall [4]. The triglyceride/ high-density lipoprotein abnormalities have recently been suggested to be related to insulin resistance.

In smokers there will be significant increase in the levels of serum cholesterol, triglycerides and HDL-C, but HDL-C is lowered in smokers than non-smokers [4]. Many studies have been done to compare the level of lipid profile among smokers and non smokers, and the non smokers and smokers with myocardial infarction. None of the studies which we have reviewed, have compared between chronic smokers with out myocardial infarction and chronic smokers with myocardial infarction. Therefore, the focus of this study was to find any other component that could have a role to play in lipid profile among chronic smokers and subsequent complications. In this context, the present study was undertaken to compare the lipid profile in apparently healthy chronic
smokers (Group A) and chronic smokers with myocardial infarction (Group B).

Material and Methods

In this case control study, 50 apparently healthy chronic smokers were chosen as controls and 50 chronic smokers with diagnosed myocardial infarction were included as cases after appropriate matching. All of them were male subjects. Both chronic smokers and chronic smokers with myocardial infarction, who smoked 14 or more cigarettes per day for more than 12 or more years (Mean values are mentioned in the table 1) were selected from Hanagal Shri Kumareshwara Hospital, Bagalkot, Karnataka, India. Those excluded from the study were persons abusing alcohol, ex-smokers, patients with diabetes mellitus, hypertension, renal diseases, hepatic impairment, endocrine disorders, obese individuals and patients on drugs like β-blockers, lipid lowering drugs and thiazide diuretics.

Ethical Clearance was obtained from S.Nijalingappa Medical College Ethical Committee. Informed consent was obtained from all the subjects.

Each subject was interviewed and information about demographic details and smoking was obtained.

The demographic details included age, sex, body weight, body mass index (BMI). Smoking habit included smoking period and number of cigarettes smoked daily.

Fasting venous blood sample was collected, serum was separated and analyzed for the following parameters - Triglycerides (TG) were measured by GOP-PAP method\textsuperscript{5,6}, optical density was measured at 546 nm. Total cholesterol (TC) and High Density Lipoprotein cholesterol (HDL-C) were measured by CHOD-PAP method\textsuperscript{5,6,7,8,9}, optical density was measured at 500 nm. Low Density Lipoprotein cholesterol (LDL-C) and Very Low Density Lipoprotein cholesterol (VLDL-C) were calculated by Friedwald’s formula\textsuperscript{10}.

Statistical analysis was done by ‘Z’ test using SPSS version 15.0. P< 0.05 was considered as statistically significant.

Results

The demographic characteristics of all the subjects are shown in Table -1. All of them were male subjects in the study. There was no significant difference in mean age, body weight and BMI among Group A and Group B. Group A and Group B were hypertensive, both groups smoked 14 or more cigarettes per day for 12 or more years (Mean values are mentioned in the table 1).

Table 2 shows lipid profile in chronic smokers (Group A) and chronic smokers with MI (Group B). The mean TC in Group A and Group B was 196.38± 22.40 mg/dl and 224.52± 13.10 mg/dl respectively. Group A had a mean TG of 166.40± 12.10 mg/dl and Group B, 192.18± 13.40 mg/dl.

The mean HDL-C in Group A was 29.70± 2.90 mg/dl and in Group B, 28.48 ± 2.60 mg/dl. The mean VLDL-C in Group A and Group B was 31.40±3.20 mg/dl and 40.18±3.16 mg/dl respectively. The mean LDL-C in Group A was 122.18± 13.60 mg/dl and in Group B, 150.84± 8.83 mg/dl.

TC, TG, VLDL-C and LDL-C in Group B were significantly higher as compared to Group A (P<0.001 for all parameters).

But there was significant decrease in HDL-C in Group A and Group B (P<0.05).

Discussion

The present study showed that the lipid parameters, TC (P<0.001), TG (P<0.001), VLDL-C (P<0.001), LDL-C (P<0.001) were significantly higher and HDL-C was significantly lower in chronic smokers with MI than chronic smokers, even when both the groups were matched for age, body weight, BMI and smoking habits.

Although cigarette smoking is a well established risk factor for vascular diseases, the genetic mechanisms that link cigarette smoking to an increased incidence of stroke are not well understood.

The inter subject variability in the atherosclerotic process in smokers may be partially mediated by genetic variants. Either CYP1A1 MSP polymorphism or certain endothelial NO synthase intorn 4 polymorphisms increased the susceptibility to cigarette smoke exposure – related atherosclerotic diseases including multi-vessel CHD and MI\textsuperscript{11}.

C alleles were associated with higher risk of carotid plaque formation and were found to act synergistically with other inflammatory single nucleotide polymorphisms (SNPs) to increase carotid intima media thickness (IMT), specifically among the smokers\textsuperscript{12}.

Cigarette smoke has shown to increase the expression of proinflammatory cytokines including IL-6. Furthermore,

<table>
<thead>
<tr>
<th>Demographics</th>
<th>Chronic Smokers (Group A)</th>
<th>Chronic Smokers with MI (Group B)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No of subjects</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>Sex· Male</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td>MeanAge (Years)</td>
<td>46.3±8.1</td>
<td>48.4±9.3</td>
</tr>
<tr>
<td>Body Wt(Kg)</td>
<td>56±6.8</td>
<td>62±3.2</td>
</tr>
<tr>
<td>BMI (Kg/m²)</td>
<td>28.1±1.04</td>
<td>30.28±0.88</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>140.21±4.83</td>
<td>146.84±9.02</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>91.83±0.4420</td>
<td>98.93±4.91</td>
</tr>
<tr>
<td>Smoking status</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td>Smoking Period (Years)</td>
<td>12.3±1.6</td>
<td>13.4±2.1</td>
</tr>
<tr>
<td>No. of Cigarettes smoked daily</td>
<td>14±2</td>
<td>15±3</td>
</tr>
</tbody>
</table>
several studies have demonstrated that IL-6 polymorphism may mediate carotid artery intima-media wall thickness. Genetic variation in IL-6 may modify stroke risk, and that this increased risk may be due to synergistic effect between pro inflammatory genotypes and smoking. The inflammatory gene SNPs are associated with early onset ischemic stroke among African-American women and cigarette smoking may modulate stroke risk through a gene-environment interaction (IL6, CD14). Several environmental factors are documented to influence redox metabolism, relatively little known about genetic effects. TAS, an indicator of redox homeostasis, is under strong genetic control, especially in smokers, this could lead to the increase in lipid peroxidation products, ultimately results in increased in oxidized LDL-C levels in chronic smokers with MI than the chronic smokers.

In conclusion, the present study suggests that smoking may modulate the IHD risk through a gene-environment interaction and which causes significant rise in TC, TG, LDL-C and VLDL-C and significant decrease in HDL-C, thereby increasing the risk of atherosclerosis. The limitations of present study were the sample size is small. MDA is more accurate when it is applied for larger population during epidemiological surveys. diet history was not taken in depth, extent of smoking ie amount of smoke exposed to lungs is not document and which is very difficult to measure. all the subjects were males.

However, further studies and long period follow up is needed- a) to find whether Group A subjects end up with myocardial infarction in due course. b) to ascertain the gene environment interaction in Group A and Group B subjects. c) to answer why all chronic smokers (14 or more cigarettes per day for a period of 12 or more years) do not have an MI.

### References


### Table 2: Lipid profile of chronic smokers and chronic smokers with Myocardial Infarction (MI)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Chronic Smokers (Group A)</th>
<th>Chronic Smokers with MI (Group B)</th>
<th>Z Value</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>TC (mg/dl)</td>
<td>196.38±22.40</td>
<td>224.52±13.10</td>
<td>7.92</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>166.12±12.10</td>
<td>192.18±13.40</td>
<td>10.20</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>29.70 ±2.90</td>
<td>28.48±2.60</td>
<td>2.20</td>
<td>P&lt;0.05</td>
</tr>
<tr>
<td>VLDL-C (mg/dl)</td>
<td>31.40 ±3.20</td>
<td>40.18±3.16</td>
<td>13.90</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>122.18±13.6</td>
<td>150.84±8.83</td>
<td>9.90</td>
<td>P&lt;0.001</td>
</tr>
</tbody>
</table>
Bilateral Talon Cusp—An unusual presentation and its management

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Abstract
The talon cusp is a relatively rare developmental anomaly characterized by the presence of an accessory cusp-like structure projecting from the cingulum area or cemento-enamel junction of the maxillary or mandibular anterior teeth in both the primary and permanent dentition. The prevalence of talon cusp varies considerably among Populations. These are usually seen in maxillary incisors unilaterally. An uncommon presentation of bilateral maxillary palatal talon cusp and its management is reported. The talon cusp can occur as an isolated finding or in association with other dental anomalies and has been reported in patients with Rubinstein-Taybi syndrome, Mohr syndrome, Sturge-Weber syndrome, incontinentia pigmenti achromians and Ellis-van Creveld syndrome. The purpose of this article is to present a case of bilateral talon cusp on geminated maxillary incisors which caused clinical problems and to describe treatment modalities.

Key Words
Talons cusp; accessory cusp; dens evaginatus.

Introduction
Talon cusp is a well delineated cusp-like anomalous structure located on the surface of an anterior tooth; it is found in both primary and permanent dentitions and can be located on lingual or labial surface, it can be unilateral or bilateral1. This is a relatively rare anomaly, with a prevalence ranging from 0.04% to 10% and affects mainly the maxillary lateral incisor, unilaterally or bilaterally2. Talon cusp has been classified by Hattab et al.3 into three basic types according to its formation and extent:

- Type I (talon) is characterized by an additional cusp projecting from the palatal aspect of an anterior tooth and extending for half the distance between the cemento-enamel junction and the incisal edge
- Type II (semi-talon) is characterized by an additional cusp 1 mm in extent or more, extending from the cemento-enamel junction for less than half the distance to the incisal edge and
- Type III (trace talon) manifests as a prominent cingulum and its variations.

Presence of bilateral talons cusp is very less in comparison to unilateral condition. Thus an unusual case of bilateral maxillary palatal talon cusps in all incisors and its management is reported.

Case Report
A 14 year old girl reported to Dept. of Pedodontics and Preventive Dentistry, KM Shah Dental College & Hospital, Vadodara, with a complaint irregularly arranged front teeth. Intraoral examination revealed well defined accessory cusps (talon cusps) on the palatal aspect of both the permanent maxillary central and lateral incisors which were fully erupted (Figure 1). The accessory cusps were markedly developed and pyramidal. These accessory cusps in lateral incisors were slightly more prominent than in the central incisors. These extra prominent cusps extended almost from the cervical third of the crown to the middle third of the crown. The developmental grooves were pronounced between the talon cusps and the palatal surface of the crown. The talons cusps were not interfering with the alignment of the mandibular anteriors during occlusion.

Both intra-oral and extra-oral examination of the patient did not reveal any abnormalities and any kind of genetic syndromes. The trait was not present in any of the family members and relatives and so the case was considered as an isolated developmental abnormality. The medical and dental histories were noncontributory.

A periapical radiograph revealed multiple V shaped radiopaque cusp like structures without a pulpal extensions (Figure 2). On this basis a diagnosis of multiple Talons cusp was made. A conservative and preventive mode of...
treatment was planned. Study casts were prepared using the impressions and also for to check any interferences (Figure 3). A periodic reduction of the talon cusps was carried out at 4–6 week intervals, using a diamond bur in a high-speed water-cooled hand-piece. Following each grinding procedure, the exposed surface was treated with fluoride varnish as a desensitizing agent. In addition, the patient was scheduled for periodic dental examination and finally, the sealant was placed near the grooves to prevent any caries (Figure 4). The patient did not have any problem other than crowding of teeth and so was referred for orthodontic treatment.

Discussion

Mitchell was the first to describe a talon cusp in 1892, Mellor and Ripa named the accessory cusp as ‘talon cusp’ because of its resemblance in shape to an eagle’s talon. To differentiate a talon cusp from an enlarged cingulum it has been suggested that a talon cusp must extend at least half the distance from the Cementoenamel Junction (CEJ) to the Incisal edge.

Other names for this condition include dens evaginatus, interstitial cusp, tuberculated premolar, odontoma of the axial core type, evaginated odontoma, occlusal enamel pearl, occlusal anomalous tubercle and supernumerary cusp. The prevalence of talon cusp is low, with estimates ranging from less than 1% to approximately 8% of the population. The anomaly is commonly unilateral, but one-fifth of the cases are bilateral in occurrence. The association of talon cusp with syndromes includes Mohr syndrome, Sturge–Weber syndrome, Rubinstein–Taybi syndrome, Incontinentia Pigmenti Achromians and Ellis–van Creveld syndrome.

Talon cusp has been classified by Hattab et al. into three basic types according to the size of the accessory cusp and its extent. According to this classification, the talon cusps described in this paper were classified as follows:

- The anomalous cusp on the maxillary lateral incisors bilaterally were prominently projected from the lingual aspect and extended from the cementoenamel junction to the incisal edge and categorized as a type 1 talon cusp.
- The other anomalous accessory cusp on the maxillary central incisors extended half the distance from the cemento-enamel junction to the incisal edge and categorized as a type 2 semi-talon.

Small talon cusps (type 2 & 3) are usually asymptomatic and need no treatment. Large talon cusps (type 1) may cause clinical problems including occlusal interference, displacement of the affected tooth, irritation of the tongue during speech and mastication, carious lesion in the developmental grooves that delineate the cusp, pulpal necrosis, periapical pathosis, attrition of the opposing tooth and periodontal problems due to excessive occlusal forces. The treatment of talon cusp involves careful clinical judgment and review of whether the cusp contains or is devoid of a pulp horn. However, in general, the management depends on individual presentation and complications.

The prognosis of teeth with talon cusp depends on the timing of diagnosis. If it is diagnosed early, the accessory cusp may be progressively removed with polishing and the abraded area should be treated with fluoride varnish. At the last appointment, the reduced cusp should be covered with resin composite. This procedure can prevent premature contact and reduce the risk of caries. The same procedure was followed in the present case and there was
no caries and other endodontic or periodontal problems. To the best of our knowledge, English Literature reports or studies of bilateral talon cusps involving all incisors are very few. Little or no epidemiological information is available in this regard.

Summary and Conclusion

The talon cusp is a developmental anomaly characterized by the presence of an accessory cusp like structure projecting from the cingulum area of the maxillary or mandibular anterior teeth in both dentitions. Here we report a rare case of bilateral maxillary palatal talon cusp and its management is also discussed. The talon cusp can occur as an isolated finding or in association with many syndromes and other dental anomalies. Talon cusp is not an innocuous defect, as it may provide a substantial challenge during diagnosis and treatment planning to the clinician. Early diagnosis may minimize local problems such as caries, periodontal disease and malocclusion. The role of a pedodontist in managing such a case is utmost important because the minimal are the future complications, better is the prognosis. So it is essential to have precise diagnostic criteria and standardization of terminology for categorization of an accessory cusp as a talon cusp.

References

Synthetic drinks and ill health in children
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Abstract
The per capita consumption of carbonated soft drink has increased over the years especially among the pediatric population. This review attempts to give an insight of the impact of soft drinks on child’s health and suggests some alternative for a healthy life style. Various research studies have shown that children consuming soft drinks have developed various acute and chronic ill effects such as nutritional deficiency, obesity, dental diseases, bone pathologies & psychological illness. As rightly said “an ounce of prevention is better than a pound cure”, Government, parents, teachers, health professionals & mainly manufacturing companies should play a crucial role in solving the problems related to soft drink consumption.

Key Words
Synthetic drinks, carbonated drinks, obesity.

Introduction
Progressive globalization of the food supply chain and the increased consumption of soft drinks, snacks, fast foods form a significant part of daily life causing acute and chronic illness¹,².

Synthetic drinks are perceived as the “other” drinking problem in society in which the addicts are mostly kids. Intense marketing efforts by large soft drink manufacturers have ensured this addiction².

This review gives an insight of the impact of soft drinks on human health & suggests alternatives for a healthy life style.

Consumption Data
The per capita consumption of all carbonated soft drinks in united states increased from 24 gallons per year in 1970 3 to 52 gallons per year in 2005 an increase of 117%. Another report from an Australian soft drink industry says that the average per capita consumption of soft drink was 110 liters in 2003. This amount equates to approximately 300ml of soft drink consumed per person per day³.

Reasons for elevated soft drink consumption:
1) Cultural shifts play a major role in changing food habits.
2) There is strong association of high television viewing rate & unhealthy food habits leading to obesity in adolescents²,⁴
3) Watching movies has been another major influence.
4) Soft drinks being tasty & easily available, it is wrongly considered to be a stress reliever.

Synthetic drink & its ingredients
Synthetic drink or soft drink is a non alcoholic beverage containing water & a flavoring agent.
Carbonated soft drinks were discovered by Europeans in 18th Century & later bubbly drinks were patented in US in 1810.

Synthetic drink components:
1. Sweeteners: It can be in the form of sugar, saccharin & aspartame. Though aspartame in small doses is said to be harmless it is one of the most controversial additives. The uncontrolled consumption of soft drinks containing aspartame leads to acute & chronic methanol toxicity in humans.
2. Aromatic Substances: The pleasant taste in the soft drinks is due to aromatic substances. They also provide better stability to the drink.
3. Carbonated water: This is supposed to make the drink more refreshing. Carbonation is done by dissolving carbon dioxide in water under pressure in chilled temperature just before the bottle is capped.
4. Acids: Citric acid, Phosphoric acid & Malic acid are most commonly used. Acidity in the drink balances the sweetness.
5. Food Colorants: Color makes the drink appetizing & attractive. Since many colorants are toxic, the usage of colorants is strictly restricted, as it can trigger asthma, hives & attention deficit hyperactive disorders in children.
6. Preservatives: Preservatives like natriumbenzoate & potassium sorbate increases the shelf life of the product. Sodium benzoate is used as a broad spectrum antimicrobial, inhibiting bacteria, molds & yeasts. However preservative like sulfur dioxide can trigger asthma, rashes, hyperactivity, fainting, shock &coma.
7. Antioxidants: Ascorbic acid prevents reactions that destroy aromatic substances in soft drinks.
8. Additives: Emulsifying agents, stabilizing agents &
thickening agents such as pectin, alginates & carraghen are also added to keep the drinks even bodied.

9. **Caffeine**: Caffeine can be addictive. It can stimulate the central nervous system & increase the heart rate. It can increase the excretion of calcium.

10. **Undisclosed Ingredients**: These are ethyl alcohol, sodium alginate, bromine in vegetable oil & Caffeine. Sodium alginate is hazardous during pregnancy.

**Health hazards of soft drink consumption**

Various research studies have shown that children consuming soft drinks have developed acute and chronic ill effects.

1. **Displacement of healthier foods from diet**

A high level of soft drink consumption is associated with less intake of vitamins, minerals & dietary fiber. Another nutritional survey has shown that soft drink consumption by adolescents has increased leading to decline in milk consumption by 10%, as a consequence of which there will be decreased intake of calcium, magnesium, phosphorus, vitamin A and protein, resulting in short & long term bone diseases.

2. **Obesity:**

The major dietary factors associated with developing childhood obesity include, increased consumption of soft drinks, fat, oils & sodium. The most frequently encountered barriers in the management of obesity include consumption of fast food & soft drinks.

There is a significant association of soft drink consumption & obesity as shown by Ludwig et al & James et al in their studies. It was also shown that soft drink consumption has increased obesity by 57%.

In a study by National heart, lung & blood institute, where in over 2000 girls were followed from ages 9-10 years till 18-19 years of age, revealed that the average soda consumption increased almost 300% over 10 years of study. This study concluded that, soda was the only beverage that was associated with increased obesity (BMI)10.

Crawford PB et al study has concluded that, out of 28 dietary factors thought to be associated with obesity in children, sweetened beverages was the only dietary factor that was consistently linked to overweight in children.

Theoretically, daily consumption of one can of sweetened soft drink (500 KJ) over a period of 10 years could lead to a 50kg increase in weight; conversely by reducing daily intake of small quantity of energy or on increasing energy expenditure (energy gap), one may prevent unhealthy weight gain.

In children who increased their consumption of sugar sweetened drinks by one serving a day, their body mass index increased by 0.24kg/m2 & their odds of being obese significantly increased10.

3. **Dental Health**

Consumption of soft drinks in large quantity & increased frequency may cause damage to teeth in the form of enamel erosions & dental caries11.

Demineralization of dental enamel has increased in the recent past, due to larger usage of soft drinks, which eventually leads to calcium & phosphorus mobilization from the enamel, as a consequence of which collapse of the surface structures occurs. Another hypothesis is that, soft drinks may cause damage to teeth through acidogenicity & cariogenicity.

Dental erosion is the situation of a chronic loss of dental hard tissue that is chemically etched away from the tooth surface by acid & chelating agents without bacterial involvement. Compared with caries, dental erosion seems to have much stronger relationship with soft drinks16.

It is found that over time, exposing dental enamel to carbonated beverage weakens & permanently destroys enamel17.

According to Australian dental association, damage to tooth enamel by acid erosion was reported in 25-45% of those surveyed. 18 Sugared versions of soft drinks proved to be more erosive than their diet counterparts19.

The low PH & high acidity leads to erosion of enamel surface. The sugars in drinks are metabolized by plaque microorganisms to generate organic acids that adds to process of demineralization, leading to dental caries14.

A study at Iowa found that intake of regular soda pop was the strongest predictor of the severity of caries. There is also a strong association between the frequency of in-between-meal consumption soda pop & caries15.

4. **Bone health**

Commonly encountered problems following prolonged consumption of soft drinks are fractured bone, low bone density, osteoporosis & hypocalcaemia.

Several studies have shown that common bone problems occur due to displacement of milk from the diet & also due to direct effects of soft drinks. For example, caffeine causes loss of calcium in the urine20 leading to osteoporosis. In the market, available 375ml of soft drink contain 40-50 mg of caffeine.

Studies have also shown that, there is strong association between soft drink & kidney stones21.

5. **Psychological & behavioral illness**

Soft drinks containing caffeine can cause CNS disturbances like sleep disturbance, bed wetting, anxiety, headache, fatigue, decreased alertness, depressed mood & irritability22.

6. **Other long term implications on health**

Several studies have shown that consuming soft drinks for prolonged period can cause obesity, hypertension, impaired glucose tolerance & hypercholesterolemia.
Presence of Benzene 23 in soft drink can be carcinogenic. Framingham heart study has shown that, those who drank one or more sodas per day were 50% more likely to develop metabolic syndrome (a combination of risk factors, such as high waist circumference, high blood pressure, impaired fasting glucose or diabetes, that strongly predicts the likelihood of developing cardiovascular disease) than those who drank less than one soda per week.26

**Healthy alternatives**
1. Coconut water: Highly alkaline & easy to digest. It has all the properties of mother’s milk.
2. Sugarcane juice & melon drinks- These are rich in natural minerals.
3. Lime juice & Butter milk
4. Dates.

**Strategies to reduce soft drink consumption**

1. **Main goals should be to**
   a) Reduce intake of soft drink consumption by young children.
   b) Reduce frequency & quantity of soft drink consumption.
   c) Replace soft drinks with naturally available drinks.
   d) Replace sweetened soft drinks with milk based drinks.

2) **Recommendation for public & others**
   - Parents should be careful while choosing a beverage. It is important that “parents serve as role models”.
   - Parents & teachers should show more concern about the overall health of children & the foods consumed in the school.
   - Pediatricians, dentists, health care professionals & dieticians should promote & support a healthy school environment.
   - Schools should hold a thorough discussion before making any decision such as, for installing a vending machine in school to dispense a food or drink.
   - Strict policies regarding prohibition of the sale of soft drinks & unhealthy food stuffs inside the school campus.
   - Ban on advertisements of beverages in the school campus.
   - Promoting healthy food stuffs in the public places, where people gather in large numbers.
   - Manufacturing companies must acknowledge the problems of rising rates of obese children & work within their limits in influencing children from taking soft drinks.
   - Government should sponsor further research to explore the ill effects of soft drinks.
   - Additional taxes on each bottle of soft drinks to curtail consumption.

**Conclusion**
Several studies have proved beyond doubt that synthetic drink is no more a safe drink. "An ounce of prevention is better than a pound cure". Government, parents, teachers, health professionals & mainly manufacturing companies should play a crucial role in solving the problems related to soft drink consumption.

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Assessment of knowledge & practices among teachers regarding school health programme

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Abstract

Objective
To study the knowledge & practice among Teachers regarding school health programme.

Study Design
Cross sectional study.

Key Words
Minor ailments, Immunization, sex education, Nutrition.

Material Method
The study was carried out in 8 higher primary schools in Gulbarga city, by systematic random sampling technique the schools were selected & 100 Teachers were selected for study. The pre tested proformas were given to all 100 Teachers.

Results
100 Teachers selected for our study comprised 39 males & 61 females. Weighted average score system was used to assess the knowledge of teacher regarding school health. Mean Knowledge of male Teachers was 20.7±7.14 and female Teachers 23.77±5.18 which was statistically significant. 82 Teachers were against giving sex education in school, out of which 79% of male Teachers & 84% female Teachers which is statistically insignificant.

Conclusion
Attitude of teacher is one important determinant of state of many health topics.

Introduction
Every child should be taught early in his life that to preserve his own life & his own health and the lives and health of others is one of his most important and constantly abiding duties. The responsibility of this lies in the hands of Parents, Teachers, health administrators and the community1.

A systematically planned & implemented health education program, modification is essential to achieve the goals more objectively, this will help to reduce the increasing burden on existing curative services2.

Material Method
The present study was carried out in 37 government higher primary schools, i.e. 5th, 6th & 7th standard in Gulbarga city having 222 teachers3. All the schools are arranged in alphabetic order in those schools all teachers are arranged in alphabetic orders, to select a sample of 100 teachers from 222 teachers using systematic random sampling

i.e. sample interval = population size/sample size
= 222/100
= 2.22 ≈ 2

Every 2nd teacher was included in the study, one random number was selected between 2 numbers, i.e. 2, so the teachers 2nd, 4th, 6th & so on included in the study.

Results & Discussion
Health promotion & health education being the most important component of school health services for which teachers are the ideal role models to impart them. They can shape & instill good habits by various methods such as role play, during games & physical education as well as by way of story telling thus teachers play a very important role especially where there is shortage of health workers in maintaining the positive health of a child, who are future citizens of the country4.

Out of the 100 teachers under study 39 were males and 61 were females. Maximum 38 teachers were in the age group of 41-50 years and minimum i.e. 11 teachers were in the age group of 21-30 years.

Table 1 shows Knowledge assessment of the teachers regarding school health as a whole was done by weighted average score, the mean knowledge score of male teachers was 20.7±7.14 and female teacher was 23.77±5.18, the difference in mean score between male & female teacher was statistically significant, i.e. female teachers have good knowledge regarding health and disease as compared to male teachers.

Table 2 shows knowledge score regarding school health
services, which was maximum in the group of teachers with S.S.L.C qualification with training i.e. 23.90±4.88 and minimum in the group of post graduate with training i.e. 18.16 ± 4.26,which is statistically significant.

Table 3 shows there is no statistical significance in knowledge in relation to the length of service of the teachers.

Table 4 shows nearly 40% of the teachers don’t have the knowledge regarding one or other six killer diseases of the childhood which is statistically found highly significant.

Table 5 shows that out of 100 teachers 72 teachers generally knew about respiratory diseases, 66 about dental diseases, 61 about communicable diseases 48 about nutritional diseases 47 about water born diseases and 16 knew about diseases caused by the bacteria’s.

Table 6 shows nutritional importance of non-vegetarian diet was not known among 37 males and 61 female teachers, but most of the male & female teachers knew about the nutritional importance of the vegetarian diet.

Table 7 our study shows 82 teachers were against giving sex education,79% of male teachers and 84% of female teachers were against giving sex education, which is statistically insignificant.

**Conclusion**

The attitude of teachers is one of the most important determinants of state of many health topics it is only essential to find out what teachers know but also to explore their attitude and feeling towards any health education programme. Successful introduction and long term sustainability of school health program is heavily dependent on the support provided by the school teachers.

**Acknowledgement**

We express our gratitude to director of department of education Gulbarga for having given permission to conduct the study, and to the Principals and Teachers.

**Table 1:** Distribution of teacher’s knowledge regarding school health according to their age & sex

<table>
<thead>
<tr>
<th>Age &amp; Sex</th>
<th>No.</th>
<th>Knowledge score Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male/Female</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>21-30</td>
<td>01</td>
</tr>
<tr>
<td>31-40</td>
<td>12</td>
<td>13</td>
</tr>
<tr>
<td>41-50</td>
<td>16</td>
<td>22</td>
</tr>
<tr>
<td>51 &amp; above</td>
<td>10</td>
<td>16</td>
</tr>
<tr>
<td>Total</td>
<td>39</td>
<td>61</td>
</tr>
<tr>
<td>Grand Total</td>
<td>100</td>
<td></td>
</tr>
</tbody>
</table>

Male Vs Female  $t_{19}$ d.f =2.37  P<0.05 Significant

**Table 2:** Distribution of knowledge score according education qualification of Teachers

<table>
<thead>
<tr>
<th>Qualification With training</th>
<th>No.</th>
<th>Knowledge score Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1 S.S.L.C</td>
<td>66</td>
<td>23.90±4.88</td>
</tr>
<tr>
<td>Group 2 P.U.C</td>
<td>09</td>
<td>20.22±4.49</td>
</tr>
<tr>
<td>Group 3 Degree</td>
<td>19</td>
<td>20.73±4.55</td>
</tr>
<tr>
<td>Group 4 Post Graduate</td>
<td>06</td>
<td>18.16±4.26</td>
</tr>
<tr>
<td>Total</td>
<td>100</td>
<td>22.41±6.30</td>
</tr>
</tbody>
</table>

Group1 Vs Group 2 $t_{73}$ d.f=2.28  P<0.05 significant
Group1 Vs Group 3 $t_{83}$ d.f=2.64  P<0.01 significant
Group1 Vs Group 4 $t_{70}$ d.f=3.13  P<0.001 highly significant
Group2 Vs Group 3 $t_{26}$ d.f=0.28  P>0.05 Insignificant
Group2 Vs Group 4 $t_{13}$ d.f=0.89  P>0.05 Insignificant
Group3 Vs Group 4 $t_{23}$ d.f=1.27  P>0.05 Insignificant

**Table 3:** Distribution of knowledge regarding school health according to their service

<table>
<thead>
<tr>
<th>Service</th>
<th>No.</th>
<th>Knowledge score Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-10</td>
<td>19</td>
<td>23.21±5.38</td>
</tr>
<tr>
<td>11-20</td>
<td>42</td>
<td>22.14±6.59</td>
</tr>
<tr>
<td>21-30</td>
<td>30</td>
<td>22.36±6.26</td>
</tr>
<tr>
<td>31-40</td>
<td>09</td>
<td>22.89±7.2</td>
</tr>
<tr>
<td>Total</td>
<td>100</td>
<td>22.41±6.30</td>
</tr>
</tbody>
</table>

Group1 Vs Group 2 $t_{59}$ d.f=0.67  P>0.05 Insignificant
Group1 Vs Group 3 $t_{47}$ d.f=0.50  P>0.05 Insignificant
Group1 Vs Group 4 $t_{26}$ d.f=0.11  P>0.05 Insignificant
Group2 Vs Group 3 $t_{70}$ d.f=0.14  P>0.05 Insignificant
Group2 Vs Group 4 $t_{49}$ d.f=0.28  P>0.05 Insignificant
Group3 Vs Group 4 $t_{37}$ d.f=0.20  P>0.05 Insignificant

**Table 4:** Distribution of Teachers according to their knowledge about six killer diseases

<table>
<thead>
<tr>
<th>Six killer Diseases</th>
<th>Know</th>
<th>Don’t know</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No.</td>
<td>%</td>
<td>No.</td>
<td>%</td>
</tr>
<tr>
<td>Tuberculosis</td>
<td>53</td>
<td>53</td>
<td>47</td>
</tr>
<tr>
<td>Diphtheria</td>
<td>61</td>
<td>61</td>
<td>39</td>
</tr>
<tr>
<td>Pertusis</td>
<td>61</td>
<td>61</td>
<td>39</td>
</tr>
<tr>
<td>Tetanus</td>
<td>61</td>
<td>61</td>
<td>39</td>
</tr>
<tr>
<td>Polio Myelitis</td>
<td>51</td>
<td>51</td>
<td>49</td>
</tr>
<tr>
<td>Measles</td>
<td>32</td>
<td>32</td>
<td>68</td>
</tr>
</tbody>
</table>

χ²5 d.f=25.43 P<0.001 highly significant

**Table 5:** Distribution of Teachers according to their knowledge regarding common diseases

<table>
<thead>
<tr>
<th>Common Diseases</th>
<th>Yes</th>
<th>No</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No.</td>
<td>%</td>
<td>No.</td>
<td>%</td>
</tr>
<tr>
<td>Disease due to Bacteria</td>
<td>16</td>
<td>16</td>
<td>84</td>
</tr>
<tr>
<td>Communicable Diseases</td>
<td>61</td>
<td>61</td>
<td>39</td>
</tr>
<tr>
<td>Respiratory Diseases</td>
<td>72</td>
<td>72</td>
<td>28</td>
</tr>
<tr>
<td>Nutritional disorders</td>
<td>48</td>
<td>48</td>
<td>52</td>
</tr>
<tr>
<td>Water borne diseases</td>
<td>47</td>
<td>47</td>
<td>53</td>
</tr>
<tr>
<td>Dental diseases</td>
<td>66</td>
<td>66</td>
<td>34</td>
</tr>
</tbody>
</table>
Table 6: Distribution of Teachers according to their knowledge regarding nutritive foods for children

<table>
<thead>
<tr>
<th>Advice nutritive food for children</th>
<th>Male</th>
<th>Female</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Yes</td>
<td>%</td>
<td>Don’t Know</td>
</tr>
<tr>
<td>Milk</td>
<td>16</td>
<td>16</td>
<td>23</td>
</tr>
<tr>
<td>Fruits</td>
<td>19</td>
<td>19</td>
<td>20</td>
</tr>
<tr>
<td>Fresh green leafy vegetables(carrot)</td>
<td>19</td>
<td>19</td>
<td>20</td>
</tr>
<tr>
<td>Egg</td>
<td>17</td>
<td>17</td>
<td>22</td>
</tr>
<tr>
<td>Meat/fish</td>
<td>02</td>
<td>02</td>
<td>37</td>
</tr>
</tbody>
</table>

χ², df=3.73, P>0.05 Insignificant

Table 7: Opinion of Teachers regarding sex education for children

<table>
<thead>
<tr>
<th>Sex education to be given</th>
<th>Male</th>
<th>Female</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No.</td>
<td>%</td>
<td>No.</td>
</tr>
<tr>
<td>Yes</td>
<td>08</td>
<td>21</td>
<td>10</td>
</tr>
<tr>
<td>No</td>
<td>31</td>
<td>79</td>
<td>51</td>
</tr>
<tr>
<td>Total</td>
<td>39</td>
<td>100</td>
<td>61</td>
</tr>
</tbody>
</table>

χ², df=3.73, P=0.05 Insignificant

References
Correlation of Apoptotic Index and Bcl-2 protein with other histological prognostic factors in prostate carcinoma

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Abstract
This study was conducted to evaluate the correlation of Apoptotic Index and Bcl-2 protein expression immuno-histochemically in Prostate Cancers with known histological prognostic factors in prostate Carcinoma including Gleason’s score and Angiogenic index (AgCD-31). The study group consisted of 35 cases of prostate adenocarcinoma obtained from the patients admitted in Dept. of Surgery and from cancer registry of Department of SN medical collage Agra. The specimens were subjected to hematoxylin and eosin staining, bcl-2 immuno histochemical staining, Apoptotic index calculation, Gleason’s grading. Univariate analysis showed no correlation between apoptotic index (A.I.) & Gleason's grade (mean A.I. 2.23±1.72 (intermediate grade) and 1±0.85 (high grade); p>0.05), negative correlation between apoptotic index and angiogenic index (mean angiogenic index =109.82 (intermediate grade) & 247.35 (high grade); p<0.05).Bcl-2 showed a positive correlation with Gleason's grade (intermediate grade 7.69% & high grade 21.82%), but Bcl-2 positive immunoreaction was not correlated with apoptotic index (p>0.05). Besides grading, quantification of Bcl-2 antiapoptotic protein and angiogenic index can be of value to predict prognosis of prostate carcinoma. Bcl-2 immunostain can be used on TURP specimens where often grading may not be representative of the entire specimen.

Keywords
Apoptotic Index, bcl-2, angiogenic index, prostate cancer.

Introduction
The incidence of prostate adenocarcinoma has risen dramatically in the past decade, probably owing to early detection, by employing digital rectal examination, serum Prostate Specific Antigen (PSA) assay and transrectal ultrasonography. Latest estimates show that it has now become the most common form of cancer and second leading cause of cancer death in men1. This change predominantly represents an increase in the number of cancers diagnosed rather than a real increase in the number of cancers in the population. Early detection has also resulted in increased life expectancy2,3.

Molecular prognostic markers in carcinoma prostate are not a new concept. As we know three main factors are required for tumor establishment- 1) Doubling time of tumor cells, 2) Fraction of tumor cells that are in replicative pool and 3) Rate at which cells are shed and lost in the growing lesion. One of the important pathways is inhibition of apoptosis or programmed cell death which may be involved in the pathogenesis of cancer by prolonging cell life and facilitating retention of deleterious mutation. Among the most important regulators of apoptosis and programmed cell death are that of B- cell lymphoma proteins(Bcl-2) and its related proteins. The protein encoded by this gene is a potent blocker of apoptosis. Complex interaction among the three subgroups within the Bcl-2 family controls the signaling events of apoptosis. Over-expression of the Bcl-2 protein occurs in a wide variety of human cancers and presumably contributes to neoplastic expansion by prolonging cell survival. Bcl-2 proteins at endoplasmic reticulum also regulate autophagy, a survival pathway that limits metabolic stress, genomic instability and tumourigenesis4.

In normal prostate Bcl-2 protein is present in the androgen independent basal cells that line the basement membrane of the prostate gland but not in the differentiated luminal secretory cells that are androgen dependent and undergo apoptosis when deprived of testosterone. In our study we evaluated the correlation of apoptotic index and Bcl-2 protein expression immuno-histocamically in prostate carcinoma with known histological prognostic factors like Gleason’s grading, scoring and angiogenic index.

Material & Methods
Thirty five cases of prostate adenocarcinoma were randomly selected among the Patients registered in Department of Surgery and Pathology of S.N. Medical College, Agra. Random sampling of 35 cases consisted of 23 cases of TURP, 11 cases of Radical prostatectomy and one needle biopsy specimen (table-1). Haematoxylin-eosin stained slides from each specimen were reviewed and the Gleason's grade was assigned. Tumours were classified as high grade when the combined Gleason's score was _8 and intermediate when score was between 5 and 7. Apoptotic index was calculated in histological tumor material. Apoptotic index was calculated by measuring apoptotic cells per 1000 tumor cells per 10high
power fields. Immuno-histochemical staining for Bcl-2 and CD-31 was performed by means of a modified labeled avidin-biotin technique, using monoclonal antibody and staining kit (DAKO LSAB2 System, HRP). The percentage of immunopositive tumour cells were scored as follows: 1=0 to <1%; 2=1 to 25%; 3=26 to 50%; 4=51 to 75% and 5=76 to 100%. Immuno-staining intensity was rated as follows: 0(none); 1(weak); 2(moderate) and 3(intense). Specimens were considered immunopositive when ≥1% of the tumour cells had clear evidence of staining. Microvessels marked with anti-CD31 in highly vascular tumour areas were counted in four adjacent high power fields. Micro vessel density was calculated per square millimeter of tumour tissue.

Statistical analysis: We categorized our sample (total specimens=35) into two groups: intermediate (13) and high grade cases (22). There were no low grade adenocarcinomas (Gleason score _ 4) in any of our specimens. After calculating apoptotic index and angiogenic index and evaluating Bcl-2, the statistical analysis was performed to draw the significance, if any, amongst various grades of prostate adenocarcinoma. Chi-sq. test was applied to find out any correlation between apoptotic index and Bcl-2. By using student’s ‘t’ test correlation was observed between the angiogenic index and apoptotic index.

Results

The study group consisted of 35 specimens of prostate adenocarcinoma which were categorized according to Gleason grading system. Apoptotic index was calculated using light microscopy on H&E stained sections. No statistically significant correlation was found between Gleason’s grade and apoptotic index (p>0.05). (Table-2): (Bar diagram). Bcl-2 immunostaining demonstrated highly variable expression in the malignant epithelium (Fig-1.). On the average, 22.86% of cases were Bcl-2 positive. Bcl-2 positive percentage increased with increasing Gleason’s grade. Strong expression of Bcl-2 protein was related to high tumour category, metastatic disease, poor histological differentiation, weak infiltration of the tumour by lymphocytes. Statistically Bcl-2 and apoptotic index were not found to be significantly correlated (p=0.08). There were cases that showed apoptotic bodies and were Bcl-2 positive (Fig-2). These cases were also subjected to CD-31 staining to find out angiogenic index. It was observed that angiogenic index increased with increasing grade(Fig-3). It showed statistically significant correlation between angiogenic index and Gleason’s grade (p<0.05).

Table 1: Case distribution of randomly sampled prostatic adenocarcinoma.

<table>
<thead>
<tr>
<th>S. NO</th>
<th>CASE OBTAINED</th>
<th>NUMBER</th>
<th>PERCENTAGE</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>TRUP</td>
<td>23</td>
<td>65.71</td>
</tr>
<tr>
<td>2</td>
<td>RADICAL PROSTATECTOMY</td>
<td>11</td>
<td>31.43</td>
</tr>
<tr>
<td>3</td>
<td>NEEDLE BIOPSY</td>
<td>1</td>
<td>2.86</td>
</tr>
<tr>
<td>TOTAL</td>
<td></td>
<td>35</td>
<td>100.00</td>
</tr>
</tbody>
</table>
Table 2: Correlation of Gleason grade with apoptotic index (A.I.), Bcl2-immunopositive cases and angiogenic index (AgCD-31).

<table>
<thead>
<tr>
<th>S.No</th>
<th>Gleason’s grade</th>
<th>Cases</th>
<th>Bcl2 positive%</th>
<th>AgCD31</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Intermediate grade (5-7)</td>
<td>13</td>
<td>0.6(2.23±1.72)</td>
<td>1(7.69)</td>
</tr>
<tr>
<td>2</td>
<td>High grade (8-10)</td>
<td>22</td>
<td>0.3(1.00±0.85)</td>
<td>7(21.82)</td>
</tr>
<tr>
<td>TOTAL</td>
<td></td>
<td>35</td>
<td>0.8(22.86)</td>
<td>8(57.17)</td>
</tr>
</tbody>
</table>

Discussion

Most prostate cancers, at the time of clinical diagnosis, present as mixture of androgen dependent and androgen independent cells. A vast majority of prostate cancers respond initially to androgen ablation since the population of androgen dependent cells undergoes rapid apoptosis upon androgen withdrawal. However, androgen ablation rarely cures patients, most of whom will experience recurrence due to takeover of the tumour mass by androgen independent tumour cells as well as emergence of apoptosis-resistant clones as a result of further genetic alterations such as Bcl-2 amplification. Proteins coded by the Bcl-2 family of genes are important regulators of programmed cell death and apoptosis. Bcl-2 is a potent antiapoptotic protein which help in development of neoplasm, by maintaining the longevity of cells via suppressing their programmed death. Its expression is more common in tumour with invasive behavior, high proliferation rate and abnormal cell differentiation. We observed a significant correlation in the Bcl-2 protein expression between the different tumour grades (Table-2). The expression of Bcl-2 family in prostate cancer (gene bax, bcl-xl and mcl-1) positively correlates with Gleason’s grade. Our results are in full agreement with those of Hering et al.10, who found positive correlation in frequency of Bcl-2 positive expression in prostate adenocarcinoma with low and high Gleason score. They concluded that over expression of Bcl-2 is significantly higher in patients with an initially elevated Gleason score8,10. The Bcl-2 expression probably exerts a role in the longevity of the cells submitted to hormonal therapy11. Khor et al.200712 observed that combination of negative Bcl-2 / normal Bax expression seemed more significantly related to reduced biochemical or any other treatment failure. In a previous analysis in prostate cancer it was suggested that the expression of Bcl-2 seems to be related to androgen independence, while in breast cancer expression of Bcl-2 is sex steroid dependent13. Accordingly, the results suggest that, along with increasing genetic instability (i.e. increased G-grade); the abnormal expression of Bcl-2 protein becomes more common in prostatic cancer. We observed that Bcl-2 positive immunoreaction did not correlate with apoptotic index (p=0.08). The relationship between inhibited apoptosis and expression of Bcl-2 protein is not clear cut; since Bcl-2 positive tumour also showed apoptosis. However, in these tumours the apoptosis rate may be reduced in relation to mitosis rate and other regulatory mechanisms, such as androgen and other growth factors may be involved. Expression of Bcl-2 protein is not a critical factor that determines the degree of apoptosis14.

In our study, we quantified apoptosis by calculating apoptotic index (A.I.) but did not find any correlation of A.I. with Gleason’s grade (p >0.05). Findings of apoptotic index in prostate cancer were in accordance with those of Amirghofran et al.15, who did not observe any correlation between apoptosis and different grades of carcinoma. They found a significant correlation between bax expression and stage of carcinoma, but not with apoptotic index, suggesting the presence of non functional bax protein or the role of other proapoptotic molecules in oncogenesis. Similarly Brown et al. (1996)16 found that the pathologic stage did not correlate with cell proliferation, apoptosis or overall daily growth. Their study concluded that cell proliferation and apoptosis do not correlate with pathological stage in clinically organ confined cancer. Similarly Matushima et al. (1998)17 showed A.I. does not correlate with Gleason score nor was there any significant correlation between AI and pathologic stage. Tu et al. (1996)18 found that localized prostate cancer cells exhibited a relatively low rate of apoptosis, which was significantly lower than the apoptotic index of normal prostate glandular epithelial cells. Metastatic prostate tumor cells, however, exhibited a significantly higher apoptotic index compared with localized prostate cancer cells.

In our study we observed that angiogenic index increases with Gleason’s grade and was inversely related to apoptotic index (p<.05) Spontaneous apoptosis is related inversely to angiogenesis and positively related to proliferative activity. Quantitative micro vessel density has been shown to provide important staging and prognostic information in prostatic carcinoma. Correlation between angiogenic index and apoptotic index has been shown to be altered significantly by androgen ablation17. The increase in micro vessels from low to high grade was in accordance with the finding of Bettencourt et al.19. They found that micro vessel density count in the tumour are significantly increased with increasing Gleason’s sum and nuclear grade but did not increase significantly in adjacent benign prostate.

Thus beside grading, scoring, lymphovascular invasion, perineural invasion; quantification of bcl-2 antiapoptotic protein and angiogenic index can be of value to predict prognosis of prostate carcinoma. Bcl-2 immunostain can be used on TURP specimens where often grading and scoring may not be what is given, as radical prostatectomy is not done. Similar large scale studies are required to derive a cut-off value of bcl-2 immuno positivity which may help in predicting progression of patients and their better management. It is also seen that high Bcl-2 positivity is present at the spreading edge of...
the tumour. The reason of this interesting finding could be decreased apoptosis at the spreading edge ensuring survival of cells to achieve invasiveness or it could be that malignant neoplasm has different cell kinetics in different areas according to survival/invasive need.

Of the many new molecular prognostic markers being evaluated in carcinoma prostate Bcl-2 and angiogenic index appear to have promise. There is an increase in Bcl-2 positive percentage with increasing Gleason's grade and a significant inverse correlation (p<0.05) exists between angiogenic index and apoptotic index. These biomarkers appear useful in Radical Prostatectomy as well as TURP (Trans urethral resection of prostate) specimens.

References
Comparative cytomorphological profile of paediatric and adult lymphadenopathy with emphasis on smear characteristics in tuberculous lymphadenitis

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¹Assistant Professor, Pathology Department, Santosh Medical College, Ghaziabad, ²Scientist E, Division of Cytopathology, Institute of Cytology & Preventive Oncology (ICMR), New Delhi

Abstract

Aims
1. To compare the cytological spectrum of pediatric and adult lymphadenopathy.
2. To analyze the differences between and cytomorphological appearances of the aspirates from tuberculous lymph nodes in the two age groups.

Material and Methods
Fine needle aspirate (FNA) smears from 500 consecutive patients with peripheral lymphadenopathy, each in paediatric (≤ 14 years) and adult (> 14 years) age groups, were retrospectively analyzed. Smear patterns in tuberculous lymphadenitis were studied in detail. The smears were air dried and stained by May Grunwald Giemsa (MGG) stain. Ziehl Neilson (ZN) staining to demonstrate acid fast bacilli (AFB) was performed where necessary.

Observations
The most frequent diagnosis on FNA was tuberculosis in both age groups (children 57%, adults 82 %), followed by reactive hyperplasia in children (37.8 %) and metastasis in adults (7.9%). Frequency of tuberculosis was significantly more in adults (p <0.001) while that of reactive process was significantly more in children (p<0.001). Cervical lymph nodes were involved most frequently by tubercular process (children 87%, adults 84 %). Caseous material or pus was aspirated in almost 2/3rd patients. The most frequent microscopic pattern was presence of granulomas against a lymphoid background, followed by necrosis only and the highest AFB positivity was found in necrotic smears in both age groups. Acute/ mixed inflammatory response, devoid of granulomas, was observed significantly more often in children as compared to adults. (p< 0.001).

Conclusions
FNAC is a useful, minimally invasive first line investigation to evaluate lymphadenopathy. An early diagnosis on FNA allows clinicians to plan and institute appropriate treatment in most cases.

Key Words
Lymph node, Fine needle aspiration cytology, tuberculous lymphadenitis

Introduction
FNAC is a well accepted diagnostic tool in adult patients. However, paediatricians and cytologists were initially hesitant in employing FNAC as a diagnostic method in children for a variety of reasons.

Cytopathologists are adapt to the procedure in adults but find it difficult in crying and squirming children. They also believed that cytomorphological features differ in paediatric lesions from those of adults and interpretation of FNAC could be inconclusive.

However, now FNAC is increasingly being applied to paediatric lesions as it permits rapid diagnosis with minimal intervention. This study evaluates the role of FNAC in paediatric lymphadenopathy with a comparative cytomorphological evaluation of lymphadenopathy in adults and children. Emphasis has been given on the comparative smear characteristics in tubercular lymphadenitis.

Superficial lymph nodes in 500 patients each of paediatric and adult age group are aspirated and subjected to microscopic evaluation regarding the diagnosis and cytomorphological assessment thereafter subjecting to comparison.

Material and Methods
Fine needle aspirate smears from 500 consecutive patients with peripheral lymphadenopathy, each in paediatric (≤ 14 years) and adult (> 14 years) age groups, referred to Santosh Medical College and Institute of Cytology and Preventive Oncology for routine diagnostic purpose, were retrospectively analyzed. The smears were air dried and stained by May Grunwald Giemsa (MGG) stain. Ziehl Neilson (ZN) staining was performed initially in all cases with grossly necrotic, purulent or blood mixed aspirates. In cases where granulomas were found without evidence of necrosis on MGG, ZN staining was performed later if an extra smear was available, or else after destaining an MGG stained smear. In patients with clinical suspicion of malignancy, a smear was also wet fixed for Papanicolaou staining.
Demonstration of AFB was considered mandatory for diagnosis of tuberculosis. Criteria to diagnose tuberculosis included a) presence of epithelioid cell granulomas with/without necrosis and AFB +ve or b) presence of necrosis/mixed inflammatory infiltrate and AFB +ve. Cases with granulomas and/or necrosis or inflammation but without AFB positivity were labelled as ‘suggestive of tuberculosis’ and advised further investigations for confirmation.

Observations

The age of patients in paediatric age group ranged from 1 month to 14 years (mean age 7.5 years) while in adults it ranged from 15 to 78 years (mean age 52 years). The maximum number of paediatric patients (40%) were in 10-14 years age group while in adults, the maximum cases (35%) were referred in 21-30 year age group. Clinically, all patients presented with peripheral lymphadenopathy, mostly painless, present since a duration of one week to one year. The most common lymph nodes aspirated were cervical, followed by axillary in both age groups. Submandibular, supraclavicular, submental and inguinal lymph nodes were occasionally involved. Bilateral involvement occurred in 27%(135/500) of paediatric cases and 14% (71/500) of adults cases while the right and the left sided distribution was almost equal. The lymph nodes were discrete in 342/500 (68.4%) children vs. 385/500 (77%) in adults while they were matted in 158/500 (31.6%) children and in 115/500 (23%) adults.

Out of 500 aspirates in paediatric age group, 15(3%) were unsatisfactory for evaluation while 10/500 (2%) aspirates from adults were unsatisfactory. The most frequent diagnosis on FNA was tuberculosis in both age groups (57% in children, 82% in adults), followed by reactive hyperplasia in children (37.8%) and metastasis in adults (7.9%) [Table 1]. NHL was diagnosed in 10 (2.4%) adults as compared to 3 (0.63%) children. (P <0.005).

Acute suppurative lymphadenitis was more frequent in paediatric age group (2.6%) as compared to the adults (1.2%). 3 cases of ‘Sinus histiocytosis with massive lymphadenopathy’ were diagnosed, all in children. Frequency of Hodgkin’s Lymphoma was similar in the two age groups.

Cervical lymph nodes were the most frequent site for metastasis, being involved in 29/39 cases.

The most common malignancy to metastasize to cervical lymph nodes was squamous cell carcinoma (25). [Primary site oropharynx in 15, larynx in 3 and esophagus in 3; primary not known in 4]. Metastatic adenocarcinoma was diagnosed in 4 cases (primary: lower end of esophagus and colon in 2 each). Axillary lymph nodes were the next common nodes to be involved in metastasis (8/39); the primary was breast carcinoma in all. In 2 cases, metastasis from squamous cell carcinoma of penis was diagnosed in inguinal lymph nodes.

Comparative evaluation of tuberculous lymphadenitis in paediatric and adult age groups.

The M: F ratio was 1:1 in both age groups with diagnosis of tuberculous lymphadenitis. Almost 50% of children presented in 10-14 year age group while most common age groups to be involved in adults was 21-30 years (48%) and 60-70 years (30%). Cervical lymph nodes were involved most frequently in both age groups (87% in children and 84% in adults); next in frequency were axillary lymph nodes (8% in children and 11% in adults). Inguinal lymph nodes were uncommon sites for tuberculosis. In children, bilateral cervical lymph node involvement was seen in 23% cases while unilateral left and right sided lymph nodes were involved 35% and 36% respectively. 5% cases showed only midline (submental) lymphoid swellings. In adults, 12% cases showed bilateral involvement, 31% left sided, 36% right sided and in 3% submental lymph nodes were involved. 54% paediatric cases presented with a solitary lymphadenopathy while 46% cases showed multiple lymph node involvement. In adults, 65% cases had solitary while 35% had multiple lymph nodes.

Gross examination of the aspirates revealed caseous material in 33%, pus in 30%, blood mixed material in 25% and turbid fluid in 12% paediatric cases while the corresponding figures were 37%, 31%, 28% and 4% respectively for adults. On microscopic examination, the most frequent pattern in paediatric tuberculous lymph nodes was presence of granulomas against a lymphoid background (30%); followed by and presence of necrosis.

| Table 1: Frequency of lesions on FNAC in paediatric and adult lymph nodes |
|---------------------------------|-----------------|---------------|------------------|
| Lesion                          | Paediatric (n=485) | Adult (n=490) | P value          |
| Tuberculosis or ‘suggestive of tuberculosis’ | 277 (57.1%)   | 402 (82%)     | <0.001           |
| Reactive hyperplasia            | 183 (37.8%)    | 29 (5.9%)     | <0.001           |
| Acute suppurative               | 13 (2.6%)      | 5 (1.2%)      | >0.05            |
| Sinus histiocytosis with        |                |               |                  |
| massive lymphadenopathy         | 3 (0.6%)       | 0             | >0.05            |
| Hodgkin’s Lymphoma              | 6 (1.2%)       | 5 (1%)        | >0.05            |
| Non Hodgkin’s Lymphoma          | 3 (0.6%)       | 10 (2.4%)     | <0.005           |
| Metastasis                      | 0              | 39 (7.9%)     | <0.001           |
only without granulomas (27%) and presence of acute/mixed inflammatory infiltrate without granulomas or necrosis (23%). In adults, the commonest smear pattern was presence of granulomas in a lymphoid background (30%), followed by necrosis without granulomas (27%).

The highest yield of AFB positivity was found in smears with necrosis only (devoid of granulomas) in all age groups (97% in paediatric and 93% in adult age groups). AFB positivity was also high in smears showing granulomas with necrosis or purulent background as well as in smears showing mixed inflammatory infiltrate without granulomas. In smear pattern showing granulomas without necrosis, AFB positivity was comparatively low in both age groups (57% in children vs. 64% in adults) [Table2]

Other cytomorphological features in the smears from tuberculous lymph nodes:

139/277 (50%) aspirates from children showed epithelioid cells, 131 of these had associated well defined granulomas while in 8 cases epithelioid cells were seen lying scattered singly or ill defined collections. In adults 209/402 (52%) aspirates showed epithelioid cells; 98 (49.4%) had associated granulomas while 11 had scattered / ill formed collections of epithelioid cells only. Langhan’s giant cells were observed n 53/277 (19.1%) paediatric cases and in 96/402 (23.9%) adult FNAs. Foreign body giant cells were found in 17/277 (6.1%) paediatric and 24/402 (5.9%) adult cases and were associated with granulomas in all the cases. Non epithelioid histiocytes were present in 107/277 (38.6%) paediatric and 133/402 (33.1%) adult FNAs. These were also seen in association with granulomas in all cases (except 2 paediatric FNAs). Foamy macrophages were present in 44/277 (15.9%) paediatric cases; 28 of these were associated with granulomas. In adult FNAs 48/402 (11.9%) cases revealed foamy macrophages, 29 of these were associated with granulomas. However, there were no significant differences between the two age groups for the presence of any of these associated features.

Discussion

The adequacy of paediatric lymph node aspirates was found to be 97% as compared to 98% in adults. Unsatisfactory aspirates resulted due to either lymph nodes being too small in size or due to excessive dilution with peripheral blood. Also, the children were at times very uncooperative during the FNA procedure leading to unsatisfactory aspirates. An almost similar inadequacy rate has been reported by Annam et al1 (3.7%) while it is 4.6% in the study of U Handa et al2, and 6% by M Jain et al6.

In some other studies it is higher (6-12%). This might be due to use of improper aspiration techniques or unsatisfactory handling of aspirated material in these studies, while in the present study, the pathologists themselves aspirated the lymphnodes. In the present study, the cervical lymph nodes were the most commonly aspirated nodes in both children (91.1%) as well as adults (83%). This closely corresponds to the figure of 85% in a study of 100 paediatric cases by Reddy et al3 and 84.3% in the study of Handa et al2. This is obvious as the cervical lymph nodes are most frequently involved during the course of tuberculosis as well as in acute upper respiratory tract infections. Also, the cervical lymph nodes are more easily visualised as compared to axillary and inguinal lymph nodes and therefore are more often reported by the patients.

Tuberculosis was the most frequent diagnosis in both paediatric and adult patients. However, when compared, tuberculosis was significantly less frequent (57% vs. 82%) while reactive hyperplasia significantly more frequent (38% vs. 6%) in children as compared to the adults. Reactive hyperplasia is fairly common in children, especially after pharyngitis or tonsillitis and skin infections. Also, the lymphoid tissue reacts briskly and for a longer period in children. The higher frequency of tuberculous lymphadenopathy (57%) in children in the present study contrasts with the studies of Annam et al1 and Handa et al2 and M Jain et al6 in which reactive hyperplasia was the most frequent diagnosis (58% and 63% respectively). This is probably because Annam et al and Handa et al have conducted their studies in Karnataka and Chandigarh respectively where due to lesser poverty and higher socio-economic status there is lesser prevalence of tuberculosis. In contrast, the patients in our study belonged to poor socio-economic status.

M Jain et al6 have worked in military hospital where the socioeconomic status and living conditions of attending patients are definitely better than of the subjects attending the civil hospital as in our study.

**Table 2:** Smear patterns correlated with AFB staining on aspirate smears from tuberculous lymph nodes in paediatric and adult age groups

<table>
<thead>
<tr>
<th>Smear pattern</th>
<th>Paediatric (n=277)</th>
<th>Adult (n=402)</th>
<th>P value (smear pattern; paediatric vs. adult)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Frequency AFB +ve</td>
<td>Frequency AFB +ve</td>
<td></td>
</tr>
<tr>
<td>Granulomas with lymphoid background (without necrosis)</td>
<td>82 (29.6%) 57.3%</td>
<td>121 (30.1%) 63.6%</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Granulomas with necrosis</td>
<td>39 (14.1%) 94.9%</td>
<td>68 (16.9%) 92.6%</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Granulomas with purulent background</td>
<td>18 (6.5%) 94.4%</td>
<td>20 (4.9%) 90%</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Acute/mixed inflammatory infiltrate without granulomas</td>
<td>64 (23.1%) 96.5%</td>
<td>81 (20.1%) 91.3%</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Necrosis without granulomas</td>
<td>74 (26.7%) 97.2%</td>
<td>112 (27.8%) 92.8%</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>
The children in present study showed a higher incidence of multiple lymph node involvement (45%) as compared to adults (35%). This is probably because of greater lymphoid tissue response and late detection of the painless swellings in children. Also, primary tuberculosis which is commoner in childhood presents as multiple lymph nodes rather than as solitary involvement. Ajali Das Gupta et al showed solitary lymph node involvement in only 19% of cases in their study done in 180 cases of lymphadenopathy in all age groups with special reference to tuberculosis. None of the other workers commented on this aspect.

The bilaterality in paediatric age group (27%) was seen in higher number of cases than in adults (14%) probably due to higher incidence of multiple lymph node involvement in children.

The right and left sided involvement was found to be almost same in all age groups in our study. Annam et al reported a higher incidence of left sided lymph node involvement (54%) than right sided (39.5%).

The aspirates were caseous in 33% paediatric tuberculous cases as compared to 36% in non paediatric cases. The frequency of purulent aspirates in paediatric and non paediatric cases was 30% and 31% respectively. Handa et al obtained caseous aspirates in 67.8% cases while Annam et al reported caseous aspirates in 71% cases. These authors did not categorise the purulent aspirates separately while we kept them in a separate category. If we combine the two, a figure of 63% is obtained which is consistent with that reported by other authors. M Jain et al reported caseous aspirates to be 80%.

Here too, the purulent smears were not categorised separately and thus clubbed together.

In the present study, AFB positivity was kept mandatory before labelling any case as tuberculous.

Cases which had necrosis but were AFB negative were labelled as ‘suggestive of tuberculosis’ and advised other investigations to establish tuberculous etiology. The highest yield of AFB positivity was found in smears with necrosis only (devoid of granulomas) in all age groups (98%). Variable AFB positivity has been reported in other studies. Handa et al found AFB positivity of 63.1% in cases showing necrosis only. Jain et al reported this figure to be 80% while 83.3% of the caseous aspirates showed AFB positivity in the study by Das Gupta et al who worked on the lymphadenopathy in all age groups. Annam et al reported AFB positivity to be 94% in paediatric tuberculous cases. The higher AFB positivity in our study might be due to diligent screening of smears by the pathologists themselves. AFB positivity was lowest across all age groups in the smears showing granulomas but no necrosis (paediatric 57.3%; adults 63.6%). There were no significant differences between the two age groups when the smear patterns were correlated with the AFB positivity. High AFB positivity in purulent aspirates observed in our study reiterates that all cases in which pus is obtained should be subjected to AFB staining and screening in order to pick up tuberculous lesions. Well formed epithelioid cell granulomas were found in 50% and 52% of paediatric and non-paediatric aspirates respectively. There were no significant differences between the two age groups for other associated features such as the presence of Langhan’s giant cells, foreign body giant cells, non epithelioid histiocytes and foamy macrophages.

References
Direct observation pattern of DOTS (Directly Observed Treatment Short Course) by Alternate DOTS providers for patients treated under RNTCP in a Tertiary Care Hospital

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Abstract

Direct Observation of Treatment Short course (DOTS) is the proven effective in controlling TB on a mass basis around the world. Direct observation is the central and key element for the success of the DOTS strategy. In India, Multi-purpose health workers play a major role in treatment observation; where they are not available, treatment observation is done by community volunteers including anganwadi workers, traditional dais, and community and religious leaders. The choice of DOTS provider should be based on access, patient preference and availability of the DOTS providers. This study was done to find the pattern of the direct observation component of the DOTS strategy by the professional and community DOTS providers and to find out their acceptance among the patients.

Material and Method

164 new sputum positive patients treated at rural and urban TB clinics of CMC Vellore from January 2001 to December 2004 were followed up during November 2005 along with their respective DOTS providers using separate structured questionnaires. The data entry and statistical analysis was done using SPSS data analysis package version 12.0. Chi2 test and t-test were used to test the significance of the data.

Result

Intensive Phase (IP) treatment was given alternate days by 94.5% of the community providers compared to 90.1% of professional providers. In Continuation Phase (CP) treatment, it was not expected by the RNTCP guidelines to give alternate days. But 34.2% of community providers had given on alternate days as in IP compared to only 4.4% of professional providers. The difference is statistically significant (p value <0.001). Patients observed by professional providers had to travel the mean distance of 0.91 Km compared to only 0.31 km with community providers (p value <0.001). The average time spent each time to get drugs from their providers in the professional arm is 27.42 minutes compared to only 14.86 minutes with community providers (p value <0.001). In the professional arm the patients are cumulated in the working hours of the professional providers and they are evenly distributed regarding time in the community arm.

Conclusion

The DOTS being provided at home with less consumption of time and lesser distance to travel are significantly different in the community arm of DOTS providers as compared to the professional DOTS providers.

Key Words

DOTS providers, Community providers.

Introduction

M. tuberculosis kills more people than any other single infectious agent after HIV. About 8 million people develop active tuberculosis (TB) every year, and about 1.8 million die¹. Every year 2 million Indians develop tuberculosis and half a million die, though the effective drugs for tuberculosis have been available since the 1940s². One of the principal barriers to the elimination of tuberculosis is non-compliance. To prevent this, WHO has stressed the need for the Direct Observation of therapy, which, as implied by the name, is the central feature of the widely advocated directly observed therapy short course (DOTS) strategy³. DOTS is the only strategy which has proven effective in controlling TB on a mass basis. The DOTS strategy is in practice in more than 100 countries⁴. Early WHO documents emphasized the importance of direct observation by health workers⁵. But later, from the experience gained in DOTS programmes around the world, it was demonstrated that trained lay people were at least as effective in observing treatment. They included community health volunteers (in Bangladesh)⁶, storekeepers (in South Africa)⁷, religious leaders, lay health workers, and community volunteers⁸.

In India, Multi-purpose health workers play a major role in treatment observation; where they are not available, treatment observation is done by community volunteers including anganwadi workers, traditional dais, and community and religious leaders. Observation by a family member is not acceptable in the Revised National Tuberculosis Control Programme⁹.
Directly observed treatment (DOT) is a strategy in which a person observes as patients take their medicine. The choice of DOTS provider should be based on access, patient preference and availability of the DOTS providers. The increase in cure and completion rates of treatment may be explained in terms of the benefit of effective implementation of the diagnosis by microscopy, adequate supply of SCC drugs, accountability of the health system and the political commitment components of the DOTS strategy.

But WHO strongly believes that direct observation is a key element for the success of the DOTS strategy. Directly observed treatment means that every dose is administered under direct observation, and convenience to the patient is essential part for success of the DOTS strategy.

Objective
This study was done to find the pattern of the direct observation component of the 'DOTS strategy' by the professional and community DOTS providers and to find out their acceptance among the patients.

Material and Method
The list of all the new sputum positive pulmonary cases treated under 2 DOTS clinics (CMC- Urban & CHAD Hospital- Rural) covered by Christian Medical College, Vellore from January 2001 to December 2004 enumerated. Those who are residing in Kaniyambadi Block, Vellore town and other areas within the radius of 12 kms and also available during the follow up in the month of November 2005 were interviewed using a structured questionnaire along with their respective DOTS providers.

In CHAD Microscopy centre, the treatment supervision is implemented by a variety of professional and community DOTS providers. Professional DOTS Providers are Doctors, Nurses, Paramedical staff of the Health system and all the RNTCP staff whether they are at private or public health system. Community DOTS Providers are any providers other than the above professional category, who are members of the community and voluntary workers who had minimal training in health care.

Statistical Analysis
The data entry and statistical analysis was done using SPSS data analysis package version 12.0. All the data were cross-tabulated and chi2 test and t- test were used to test the significance of the data.

Result
There were 164 patients followed up. The total number of their respective DOTS providers came to 86. Table 1 shows the occupation of the DOTS providers. In the professional arm, RNTCP Staff and private practitioners have observed more cases. In the community arm, the occupation rages from housewives to politicians.

### Table 1: Occupation of the Dots Providers

<table>
<thead>
<tr>
<th>Professional DOTS providers</th>
<th>Occupation</th>
<th>Number</th>
<th>Patients observed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Private practitioner</td>
<td>8</td>
<td>21</td>
<td></td>
</tr>
<tr>
<td>Doctor in Municipal Health Centre (MHC)</td>
<td>1</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>RNTCP staff</td>
<td>3</td>
<td>23</td>
<td></td>
</tr>
<tr>
<td>Maternity asst. (ANM)</td>
<td>8</td>
<td>17</td>
<td></td>
</tr>
<tr>
<td>Field nurse (MHC)</td>
<td>9</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Nurse in CMC</td>
<td>4</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>Pharmacists</td>
<td>2</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>35</td>
<td>91</td>
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</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Community DOTS providers</th>
<th>Occupation</th>
<th>Number</th>
<th>Patients observed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attenders - CMC/CHAD</td>
<td>4</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Balwadi teacher</td>
<td>2</td>
<td>2</td>
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<tr>
<td>Health aides (CHAD)</td>
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<td>Part time Community Health Workers (CHAD)</td>
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<td>Extension worker</td>
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<td>1</td>
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<td>Other NGOs</td>
<td>3</td>
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<td>Community nutrition worker</td>
<td>1</td>
<td>2</td>
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<td>Maternity Ayah</td>
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<td>Cooley</td>
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<td>Beedi worker</td>
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<td></td>
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<td>House wife</td>
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<td>Milk man</td>
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<td>1</td>
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<tr>
<td>Photographer</td>
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</tr>
<tr>
<td>Local politician</td>
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</tr>
<tr>
<td>Teacher</td>
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</tr>
<tr>
<td>Tea shop</td>
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<td>1</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>51</td>
<td>73</td>
<td></td>
</tr>
</tbody>
</table>

Table 2 describes the direct observation pattern of DOTS by the professional and community providers. Intensive Phase (IP) treatment was given alternate days by 94.5% of the community providers compared to 90.1% of professional providers. In Continuation Phase (CP) treatment, it was not expected by the RNTCP guidelines to give alternate days. But 34.2% of community providers had given on alternate days as in IP compared to only 4.4% of professional providers. The difference is statistically significant (p value <0.001). Regarding observation more than 85% of the patients agreed that they have been fully observed during the intensive phase of DOTS. It was 85.7% in the professional arm and 87.7%
in the community arm. The difference is not statistically significant (p-value = 0.71).

Table 2: Direct Observation of the Treatment

<table>
<thead>
<tr>
<th>Variables</th>
<th>Professional (n=91)</th>
<th>Community (n=73)</th>
<th>Chi2</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IP given alternate days</td>
<td>82 (90.1%)</td>
<td>69 (94.5%)</td>
<td>1.08</td>
<td>0.22</td>
</tr>
<tr>
<td>CP given alternate days</td>
<td>4 (4.4%)</td>
<td>25 (34.2%)</td>
<td>24.79</td>
<td>0.00*</td>
</tr>
<tr>
<td>IP fully observed</td>
<td>78 (85.7%)</td>
<td>64 (87.7%)</td>
<td>0.13</td>
<td>0.71</td>
</tr>
<tr>
<td>CP fully observed</td>
<td>52 (82.4%)</td>
<td>65 (89%)</td>
<td>1.42</td>
<td>0.23</td>
</tr>
</tbody>
</table>

Table 3 shows the distance traveled and time spent by the patients to receive drugs from their providers. Patients observed by professional providers had to travel the mean distance of 0.91 km compared to only 0.31 km with community providers. The difference is statistically significant (p value <0.001). The average time spent each time to get drugs from their providers in the professional arm is 27.42 minutes compared to only 14.86 minutes with community providers. The difference is statistically significant (p value <0.001).

Table 3: Distance Travelled and Time Spent by the Patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Mean</th>
<th>S.D.</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Distance travelled (Km)</td>
<td>Professional 0.91</td>
<td>1.21</td>
<td>0.00*</td>
</tr>
<tr>
<td></td>
<td>Community 0.31</td>
<td>0.54</td>
<td></td>
</tr>
<tr>
<td>Time Spent (Minutes)</td>
<td>Professional 27.42</td>
<td>15.8</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Community 14.86</td>
<td>17.8</td>
<td></td>
</tr>
</tbody>
</table>

Table 4 illustrates the place of DOTS being given. In the professional arm the main place was clinics/hospitals (90.1%) and in the community arm mainly either it was patients’ house (452%) or providers’ house (41.1%).

Table 4: Place of Dots for Patients Interviewed

<table>
<thead>
<tr>
<th>Place of DOTS</th>
<th>Professional</th>
<th>Community</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients’ house</td>
<td>3 (3.3%)</td>
<td>33 (45.2%)</td>
</tr>
<tr>
<td>Providers’ house</td>
<td>6 (6.6%)</td>
<td>30 (41.1%)</td>
</tr>
<tr>
<td>DOTS clinics/ hospital</td>
<td>82 (90.1%)</td>
<td>8 (11%)</td>
</tr>
<tr>
<td>Common place</td>
<td>0</td>
<td>2 (2.7%)</td>
</tr>
<tr>
<td>Total</td>
<td>91</td>
<td>73</td>
</tr>
</tbody>
</table>

Table 5 shows the timing of the provision of DOTS by the providers. In the professional arm, the patients are cumulated in the working hours of the professional providers and they are evenly distributed regarding time in the community arm.

Table 5: Timing of Dots for the Patients Interviewed (According to DOTS providers)

<table>
<thead>
<tr>
<th>Timing of DOTS</th>
<th>Professional</th>
<th>Community</th>
</tr>
</thead>
<tbody>
<tr>
<td>Morning</td>
<td>17 (48.6%)</td>
<td>22 (43.1%)</td>
</tr>
<tr>
<td>Noon</td>
<td>12 (34.3%)</td>
<td>6 (11.8%)</td>
</tr>
<tr>
<td>Evening</td>
<td>4 (11.4%)</td>
<td>13 (25.5%)</td>
</tr>
<tr>
<td>Night</td>
<td>2 (5.7%)</td>
<td>10 (19.6%)</td>
</tr>
<tr>
<td>Total</td>
<td>35 (100%)</td>
<td>51 (100%)</td>
</tr>
</tbody>
</table>

Chi2: 10.11 Df: 3  p-value: 0.017

Discussion

In the professional arm while most of them are the professionals working in the public or the private health sector, 8 out of 35 (22.9%) of them are the private practitioners. This shows better awareness and acceptance of DOTS by the private practitioners. The occupation in the community arm ranges from housewives to local politician. Whatever may be the profession, the accessibility and acceptability by the patients and the accountability to the health system are the guiding principles to choose a person as a DOTS provider.

According to the RNTCP treatment guidelines the treatment provider should observe each dose in the intensive phase (IP) which is given every alternate days for 2 months and at least the first dose in the continuation phase (CP) which is given once a week. In this study irrespective of the intensive and continuation phase, significant numbers of the providers in the community arm continue to give DOTS on alternate days in the continuation phase. They are doing it with enthusiasm since they felt DOTS will be effective if it continued alternate days in the continuation phase also.

The distance between the patients and the healthcare workers is a formidable obstacle, making regular observation of doses, even three times weekly, difficult to achieve. The table shows that there is the significant difference in the average distance travelled by the patients and the average time spent for each dose of DOTS including time taken for travel. The patients in the community arm are spending significantly less time for travel and less over all time for getting DOTS (p value < 0.001). But for each patient arranging a community provider is the big challenge and burden to the health system/ service provider. In practical situation, where the health facility is not available or inconvenient to the patient, the community providers are arranged.

Convenience to the patient is essential part for success of the DOTS strategy. For the success of the tuberculosis treatment, the DOTS providers must be organized based on patient’s convenience rather than the convenience of the treatment providers. In this study, in the professional arm 90% of the patients have gone to MHCs / DOTS clinics in CHAD/CMC or to the clinics of the private practitioners. In the community arm 33 patients (45.2%) had received the treatment at their homes since the community providers are willing to go patients place and administer the drugs.

But opinion among the DOTS providers regarding the ideal place of DOTS varies. The place is whether the patient’s house or the provider’s house/Clinics, both have its own advantages and disadvantages. DOTS at patient’s house ensures the compliance to the schedule and is convenient for the patients. But then the responsibility of the regular intake of drugs by the patient is on the DOTS providers. As expressed by some of the DOTS
providers, the regular visits to the patients’ house make
the patients stigmatized in the community. Some of the
DOTS providers prefer the patients’ house as the ideal
place to avoid the inconvenience to schedule their time
for the patients.

The timing of the DOTS observation in the professional
arm is cumulated during the morning and noon because
of the working hours of the professional providers. In
the community arm it is evenly distributed because of
their availability to the patients’ convenient time. This
difference is statistically significant (p value 0.017).

Conclusion

The characteristics of the DOTS providers showed that
the community DOTS providers included the different
categories including house wives, carpenter, milkman,
photographer, and teacher and so on. The DOTS being
provided at home with less consumption of time and
lesser distance to travel are significantly different in the
community arm of DOTS providers as compared to the
professional DOTS providers. This involves the challenge
to the health system/voluntary agencies to train, organise
and supervise the community DOTS providers to make
the DOTS convenient to the patients. The community
DOTS providers have certain advantages like making
DOTS more accessible at home level for the patients.

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Tomans’ tuberculosis: Case detection, treatment and
monitoring – questions and answers. Second edition, Geneva:
Oral plasmablastic lymphoma in an HIV negative patient – A case report

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Abstract

Plasmablastic lymphoma (PBL) is a rare variant of a diffuse B-cell lymphoma that is most commonly found in the oral cavity affecting the mucosa, and so named for its blastoid morphology and immunophenotype. Oral PBL has been treated heterogeneously and well defined treatment guidelines are still lacking. It includes chemotherapy, radiotherapy, combination of both or local excision followed by radiation. Prognosis is usually poor regardless of the site of origin. The clinical course is very aggressive with most of the patients dying in the first year after diagnosis. Here we present a case who was treated with high grade lymphoma protocol, CODOX-M alternating with IVAC followed by radiotherapy & Patient showing a good clinical response.

Keywords

Oral Plasmablastic lymphoma, HIV- Negative.
Abbreviations: CODOX-M, C-cyclophosphamide, O-vincristine, DOX-doxorubicin, M-methotrexate.
LCA –leucocyte common antigen.

Introduction

Plasmablastic lymphoma (PBL) is a rare variant of a diffuse B-cell lymphoma that is most commonly found in the oral cavity affecting the mucosa, and so named for its blastoid morphology and immunophenotype1,2. These tumors are found in patients with human immunodeficiency virus (HIV), affecting approximately 3% of all HIV patients3,4. It is not atypical to find non-Hodgkin's lymphomas (NHL) associated with acquired immunodeficiency syndrome (AIDS) and in some cases its manifestation delineates an HIV-positive patient as having AIDS. Male to female ratio is 7:1 for the disease. As a distinct subset, PBLs are noted to be rapidly progressive with a high mitotic index5. Although prognosis is usually poor, with an average survival time being approximately 6 months1,2,6. Report of a case: A 30 year old male patient presented with a growth and pain in the right upper jaw since 2 months. Patient was a known tobacco chewer, since 10 years. Intraoral examination revealed, ulcerative lesion approximately 4 X 3 cm in diameter at right hard palate & posterior alveolus region(Fig.a). Anteriorly posteriorly lesion extended from second premolar to the right tuberosity region. Neck examination revealed right level I b sub centimeter node, tender, soft to firm & mobile in nature. CT scan revealed a well defined irregular heterogeneous soft tissue attenuation of right upper alveolus region measuring in 4.6 X 2.5cm suggested of malignant neoplasm of right maxilla with multiple sub centimeter size node at level Ib & level II.(Fig.b) Complete blood count and metabolic profile presented no abnormalities. FNAC done, was inconclusive. An incisional biopsy of area was performed & submitted for examination, the report suggestive of high grade plasmablastic lymphoma. Immunohistochemistry studies showed that tumor cells were positive for LCA, CD20. CD138 focally positive & CD3 negative. Kappa and lambda stain were not very satisfactory.

Differential diagnosis of oral PBL includes poorly differentiated carcinoma, amelanotic melanoma & other type of lymphoproliferative disease. AIDS related lymphomas that are included in differential diagnosis of PBL are DLBCL (Diffuse large B cell lymphoma), plasmacytoma & small noncleaved (Burkitt-like) lymphoma7,8,9,10. Plasmacytoma and PBL cannot be reliably distinguished by immunohistochemistry alone as they exhibit similar immunoprofile.

Treatment

Oral PBL patients have been treated heterogeneously and well defined treatment guidelines are still lacking. Treatment may consist of chemotherapy, using prednisolone, cyclophosphamide, adriamycin and/ or vincristine, or local excision followed by radiation. The final course of therapy is considered on a case-by-case basis depending on the stage of the disease, the presence of systemic symptoms or the association with HIV infection11. Local radiotherapy has proven successful when only gingival lesions are present. Multi-drug chemotherapy with or without radiation treatment is generally recommended for the disseminated disease. In this case patient was started on high grade lymphoma protocol. CODOX-M alternating with IVAC, but after
second cycle patient developed cytosine arabinoside induced CNS toxicity in the form of altered sensorium and seizures. However CT scan brain revealed no abnormalities. After the recovery, patient was planned for Radiotherapy (40 Gy / 20 fractions) following which a good clinical response was seen (Fig.c).

Discussion

Oral PBL is an uncommon, recently described B-cell derived lymphoma most commonly seen in patients with HIV infection. The median age at presentation is around 50 years, with a broad distribution, but mainly affecting adults. Immunodeficiency caused in the majority of cases by HIV, predisposes to the development of PBL. Other causes of immunodeficiency such as iatrogenic immunosupresion for autoimmune disease or prevention of post transplant therapy allograft rejection may also be implicated. It is characterized by a diagnostic triad of predilection for gingivo- buccal complex mucosa, classical plasmablastic morphology with the lack of neoplastic plasma cells and a limited immunohistochemical panel consisting of CD20 negativity, LCA (+/−), CD138/ VS38c diffuse positivity, light chain restriction. PBL presents most frequently as a mass in the oral cavity, but it is also encountered in other extra nodal sites – particularly mucosal sites including the sinonasal cavity, orbit skin, bone, soft tissues and gastrointestinal tract. Computed tomography and positron emission tomography may show disseminated bone involvement. Cytosplasmic immunoglobulins are expressed in 50-70% of cases, most frequently IgG and either kappa or lambda light chain. The expression of CD56 should raise suspicion for underlying plasma cell myeloma. Oral PBL has been treated heterogeneously and well defined treatment guidelines are still lacking. It includes chemotherapy, radiotherapy, combination of both or local excision followed by radiation. Prognosis is usually poor regardless of the site of origin.

The clinical course is very aggressive with most of the patients dying in the first year after diagnosis.

References

Lymphangiomatous macroglossia: A case report

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Abstract
Lymphangioma is a benign malformation of the lymphatic channels. The anterior two-thirds on the dorsal surface of the tongue is the most common site for intra oral lymphangioma leading to macroglossia. We present a case of lymphangiomatous macroglossia in a 25 year old male who underwent surgical excision of the lesion followed by a primary closure. On follow-up after, there was an improved tongue appearance and good tongue function.

Key Word
Lymphangioma, Macroglossia.

Introduction
The first accurate description of lymphangioma was given by Virchow in 1854. Lymphangiomas are benign hamartomatous tumors of the lymphatic channels1.

They present as developmental malformations arising from sequestration of lymphatic tissue that do not communicate with the rest of the lymphatic channels1. They can also occur in association with hemangioma. Lymphangiomas have a marked predilection for the head and neck region, which accounts for about 75% of all cases and about 50% of these lesions are noted at birth and around 90% develop by 2 years of age2. They are known to be associated with Turner’s syndrome, Noonan’s syndrome, trisomies, cardiac anomalies, fetal hydrops, fetal alcohol syndrome, and Familial pterygium coli2.

Oral lymphangiomas may occur at various sites but they form most frequently on the anterior two-thirds of the tongue, which often result in macroglossia. It can also present in the palate, buccalmucosa, gingiva, and lip3,4.

Case Report
A 25 year old male patient presented to oral surgery department with enlargement of tongue since 2 years (FIGURE 1). Oral examination revealed a 7x5 cm mass at the anterior two thirds of the tongue. History revealed the mass was initially small in size and later it increased up to the present size. The patient could hold his tongue in the oral cavity with no difficulty. The mass had a soft elastic consistency, and its surface was covered by epithelium and small lymphangiomatous vesicles producing an irregular granular appearance. There was no associated pain, dysphagia, dysgeusia, or neurosensory disturbance. No other mass was detected in the head and neck region. laboratory parameters were all within normal limits. FNAC done was inconclusive, later an incisional biopsy was performed of the tongue lesion. A diagnosis of lymphangioma was established. Lymphangioma being a benign neoplasm, a wide excision of the lesion was planned.

Subsequently the patient underwent surgical resection under general anesthesia. An inverted v shaped incision was outlined on the dorsal and lingual aspects of the tongue with methylene blue. (FIGURE 2)

After removal of the lesion, the remaining part of the tongue enclosed in a layered manner, leading to a
relatively normal size and shaped tongue. (FIGURE3,4)
The patient was discharged after a week with a normal looking tongue and marked improvement in speech. The post surgical histopathological report indicated cavernous lymphangioma. At present patient is in regular follow up with an improved tongue appearance and function. (FIGURE 5)

Discussion
According to Sabin5, the lymphatic system arises from five primitive buds that develop from the venous system. This is completed by the second month of gestation. From these buds, paired jugular sacs, paired posterior sacs in proximity to the sciatic veins, and a single retroperitoneal sac are formed. Lymphangioma is generally acknowledged to be a disease of childhood during which time there is active lymphatic growth. Lymphangiomas can occur from birth up to the first 2 years of life.

The predominant site of lymphangiomas is in the head and neck region, and this may be related to the anatomic location of the embryologic sacs6.

Lymphangiomas are classified into three subtypes7: (1) lymphangioma simplex, composed of capillary-sized, thin walled lymphatic channels; (2) cavernous lymphangioma, composed of dilated lymphatic channels, often with fibrous adventitial coats; and (3) cystic lymphangioma or hygroma, composed of cysts varying from a few millimeters to several centimeters in diameter. Clinically, the lymphangiomas that arise in the tongue show macroGLOSSIA with irregular translucent, grape-like vesicles of lymphatic origin. Most lingual lymphangiomas are simple or cavernous lymphangiomas, and affect the dorsum of the tongue.

The anterior two-third on the dorsal surface of the tongue is the most common site for intra oral lymphangioma leading to macroGLOSSIA3,4.

The various treatment modalities for lymphangioma are surgical excision, radiation therapy, cryotherapy, electrocautery, sclerotherapy, steroid administration, embolization and ligation8,9, laser surgery with ND-YAG10-12,CO213,14,and radio-frequency tissue ablation technique15.

The method used in this case is best suited for lymphangiomatosus lesions in the anterior-middle portion of the tongue has the following advantages: 1) After healing, the shape resembles a normal tongue and the suture line resembles the median furrow; 2) The movements of the tongue are not impaired and so articulation is facilitated; and 3) The inverted V-shaped incision leaves normal tissue on the lateral margins of the tongue, which contains taste buds.

References
A Study of disabilities among children (0-14 years) in rural field practice area of Rural Medical College, Loni, Maharashtra

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¹Associate Professor, Department of Community Medicine, ²Statistician, Dr. Pinnamaneni Siddhartha Institute of Medical Sciences & Research Foundation, Chinaoutpalli, A.P.

Abstract

Objectives
1. To find out the prevalence of multidisciplinary disabilities. 2. To assess the relationship between the disabilities & socio economic and cultural factors.

Methodology
The study area comprises of 8 villages under rural field practice area of Rural Medical College, Loni. 7300 children of the age group of 0-14 yrs from the total population of 20,533 were studied by Community based Cross Sectional study. House to house survey was conducted to identify disabled children using a pre tested questionnaire by interview technique.

Results
The overall prevalence rate of disabilities in children was 2.25%. The prevalence rate for Hearing & Speech impairment was more(1.34%) followed by locomotor disability(0.32%), Visual impairment(0.23%), Mental retardation(0.19%) and Multiple & Miscellaneous disability(0.17%). The disability was more in boys, in higher age groups, among Muslims, children of laborers, family size of four and above and in the lower socio economic class.

Key Words
Disabilities, Prevalence, Socio-economic status.

Introduction
Disability has been defined as “any restriction or lack of ability to perform an activity in the manner or within the range considered normal for a human being”¹. According to World Health Organisation’s estimates, approximately 10% of a given population suffer from disability of one kind or other. There were no comprehensive surveys to know the exact incidence of disability in India.

Government of India Census 2001 has included disability, a separate question and revealed that 80 per cent of the disabled children were in rural areas. The major preventable causes of disabilities are malnutrition, communicable diseases, early childhood infections and accidents at home and work place. Early detection of impairment, combined with early and effective curative care can make a significant impact in minimizing or compensating for impairment and its consequences².

Persons with disabilities who belong to poor families are marginalized and disadvantaged by variety of factors such as lack of access to productive resources & to opportunities, and lack of information and skills which enable participation in social, economic and political process. Some groups such as women and girls are more vulnerable to disabilities. It was estimated that only 2 to 3% of disabled in the need of rehabilitation have access to the services.

Persons with disabilities frequently live in deplorable conditions, facing barriers that prevent their integration and meaningful participation in mainstream society. The basic human rights to freedom of movement, access to education and health care are often ignored. Because they suffer the additive difficulties of their disability, marginalization and invisibility, their health, especially their mental health may deteriorate even further³.

The persons with disabilities and their family members are socially, economically and emotionally affected. The negative attitudes of the abled persons in the family and in the community are the greatest obstacles to full participation and equalization of opportunities.

Society’s understanding and approach to the issues of the disabled has been fast changing for the past 30 years. Newer advances in technology, new civil rights movements, greater number of disabled people making their marks in different social, political, economic and other sectors have helped in mainstreaming of the disabled citizens⁴.

The establishment of Rehabilitation Council of India has been a major move for quality assurance in the education, training and management of persons with disabilities. Persons with Disabilities (Equal opportunities, Protection of rights and Full participation) Act,1995 fixes the responsibilities on central and state governments to provide services, create facilities and give up support to the people with disabilities in order to enable them to have an equal opportunity in participating as well as
productive and contributing citizens of the country to their fullest extent. A new strategy termed Community Based Rehabilitation (CBR) was evolved and found extremely useful to rehabilitate persons with disabilities in the community setting and with community participation.

Prevention, early identification, intervention, rehabilitation, integration and inclusion of all persons with disabilities are the concept of today, where by such people also have a right to their family and to a natural environment.

Material & Methods

Study area: The study area comprises of 8 villages under rural field practice area of Rural Medical College, Loni (Maharashtra). Study population: 7,300 children in the age group of 0-14 yrs from the total population of 20,533. Study design: The data was collected through a well designed community based cross sectional study. Sample size determination: The prevalence of disability among children in rural area was considered as 5% (P) for computation of the sample size. Keeping the confidence level as 95% and the relative result of the survey results as 10% of P i.e. 0.5%, the sample size was calculated by using the formula \( n = \frac{Z^2 \times (1 - \alpha/2) \times P \times (1 - P)}{\Sigma^2} \) where \( Z = 1.96 \) (C. L=95%) \( P = 5\% \) \( \Sigma = 0.5\% \) (10% of P = 5). The sample size has been arrived at 7229 rounded to 7300. Selection of study population / Sample survey methods: House to house survey was conducted to identify disabled children using a pre tested questionnaire. The data was collected through interview technique. Respondent was the head of the family or parent or close relative of the children in the house. Quality assurance of the data: Daily checking of the 10% of the filled questionnaire by the senior colleague in the department. Results were discussed with senior colleagues and summarised. Statistical analysis and interpretation of data: Data collected has been presented through frequency distribution tables, cross tables and graphs. Interpretation of the results were done using percentages, proportions and tests of significance – Chi square test.

Results

Table 1: Distribution of children with disabilities by the Type & Prevalence of disability

<table>
<thead>
<tr>
<th>Type of disability</th>
<th>Number</th>
<th>Prevalence rate per 100</th>
</tr>
</thead>
<tbody>
<tr>
<td>Locomotor</td>
<td>23(14.02%)</td>
<td>0.32</td>
</tr>
<tr>
<td>Mental retardation</td>
<td>14(08.54%)</td>
<td>0.19</td>
</tr>
<tr>
<td>Visual impairment</td>
<td>17(10.36%)</td>
<td>0.23</td>
</tr>
<tr>
<td>Hearing &amp; Speech</td>
<td>98(59.76%)</td>
<td>1.34</td>
</tr>
<tr>
<td>Multiple &amp; Miscellaneous</td>
<td>12(07.32%)</td>
<td>0.17</td>
</tr>
<tr>
<td>Total</td>
<td>164(100%)</td>
<td>2.25</td>
</tr>
</tbody>
</table>

Table 2: Distribution of children by Age & Sex

<table>
<thead>
<tr>
<th>Sex</th>
<th>0-4 yrs</th>
<th>5-9 yrs</th>
<th>10-14 yrs</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>16</td>
<td>49</td>
<td>45</td>
<td>110(67%)</td>
</tr>
<tr>
<td>Female</td>
<td>05</td>
<td>19</td>
<td>30</td>
<td>54(33%)</td>
</tr>
<tr>
<td>Total</td>
<td>21(12.81%)</td>
<td>68(41.46%)</td>
<td>75(45.73%)</td>
<td>164(100%)</td>
</tr>
</tbody>
</table>

Chi square = 3.25  D.F.= 2  P > 0.05

Table 3: Distribution of children by Religion

<table>
<thead>
<tr>
<th>Religion</th>
<th>Children 0-14yrs</th>
<th>Children with disabilities</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hindu</td>
<td>626 (85.84%)</td>
<td>136 (82.92%)</td>
</tr>
<tr>
<td>Christian</td>
<td>122 (01.67%)</td>
<td>01 (0.61%)</td>
</tr>
<tr>
<td>Muslim</td>
<td>912 (12.49%)</td>
<td>27 (16.47%)</td>
</tr>
<tr>
<td>Total</td>
<td>7300 (100%)</td>
<td>164 (100%)</td>
</tr>
</tbody>
</table>

Chi square = 3.413  D.F.= 2  P value : 0.1815

Table 4: Distribution of children by Parent's Occupation

<table>
<thead>
<tr>
<th>Occupation of head of household</th>
<th>Total children</th>
<th>Children with disabilities</th>
</tr>
</thead>
<tbody>
<tr>
<td>Farmer</td>
<td>3073(42.10%)</td>
<td>43(26.22%)</td>
</tr>
<tr>
<td>Laborer</td>
<td>2748(37.64%)</td>
<td>81(49.40%)</td>
</tr>
<tr>
<td>Business</td>
<td>557(07.63%)</td>
<td>20(12.19%)</td>
</tr>
<tr>
<td>Service</td>
<td>922(12.63%)</td>
<td>20(12.19%)</td>
</tr>
<tr>
<td>Total</td>
<td>7300(100%)</td>
<td>164(100%)</td>
</tr>
</tbody>
</table>

Chi square = 20.80  D.F.= 3  P value : 0.0001

Table 5: Distribution of children by Family Size

<table>
<thead>
<tr>
<th>No. of living children</th>
<th>Children (0-14yrs)</th>
<th>Children with disabilities</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>112(01.54%)</td>
<td>01(00.61%)</td>
</tr>
<tr>
<td>2</td>
<td>812(11.12%)</td>
<td>28(17.07%)</td>
</tr>
<tr>
<td>3</td>
<td>1211(16.59%)</td>
<td>37(22.56%)</td>
</tr>
<tr>
<td>4+</td>
<td>516(57.75%)</td>
<td>98(59.76%)</td>
</tr>
<tr>
<td>Total</td>
<td>7300(100%)</td>
<td>164(100%)</td>
</tr>
</tbody>
</table>

Chi square = 12.75  D.F. = 3  P value : 0.0052

Table 6: Distribution of children by Socio–economic status *

<table>
<thead>
<tr>
<th>Socio-economic status</th>
<th>Children (0-14yrs)</th>
<th>Children with disabilities</th>
</tr>
</thead>
<tbody>
<tr>
<td>Upper</td>
<td>179 (0.23%)</td>
<td>---</td>
</tr>
<tr>
<td>Upper middle</td>
<td>197(02.70%)</td>
<td>02 (01.22%)</td>
</tr>
<tr>
<td>Lower middle</td>
<td>454(06.22%)</td>
<td>14 (08.54%)</td>
</tr>
<tr>
<td>Upper lower</td>
<td>1940(26.58%)</td>
<td>43 (26.22%)</td>
</tr>
<tr>
<td>Lower</td>
<td>4692(64.27%)</td>
<td>105 (64.02%)</td>
</tr>
<tr>
<td>Total</td>
<td>7300(100%)</td>
<td>164(100%)</td>
</tr>
</tbody>
</table>

*Modified B.G.Prasad’s Socio-economic status classification

Chi square = 2.811  D.F.=3  P value : 0.4217

Discussion

The prevalence rate of disability is 2.25%. Major disability is Hearing and Speech impairment followed by Locomotor disability, Visual impairment, Mental retardation and less is Multiple disabilities. The prevalence rate of
disabilities was less than the estimated figure of 10% of world population by WHO1, but was falling within the range of 2-5% of the Indian population as estimated by Rehabilitation Council of India5. The disability was more than double in boys as compared to girls as reported by Joshi & Sharma(1971)7, but reported equally by Mathur et al.8. The disability was low in lower age group than those in higher age group of children except in Cerebral Palsy which was more in lower age group.

The prevalence of disability was more among Muslim children and less in Christians. It may be because of the common practice of consanguineous marriages and large family size as reported by Vidyabhushan and Sachdeva DR(2003)9. The highest disabilities were observed among children of laborers followed by farmers and lowest among children of businessmen and service people. This may be because of low socio economic status of laborers and farmers with poor nutritional status, poor living conditions and poor utilization of health services.

The disabilities were more with the family size of four and above and less with the size of one, may be due to poor maternal factors and poor utilization of health services. The disabilities were proportionally decreasing with the increasing socio economic status clearly indicating the inverse relationship as was observed by, Mathur et al (1985)8.

Conclusion

The rural community should be made aware of consanguineous marriages and such marriages to be discouraged. The adolescents to be educated about the age at marriage as early pregnancy increases the disability risks. Ensure provision and utilization of Reproductive & Child Health services, especially safe delivery and family planning practices as the disabilities were observed more in home deliveries conducted by untrained personnel, large family size and increased birth order. Complete immunization of all the children is to be ensured. Nutritional education to be imparted. Safety education must be incorporated in the school curriculum.

Acknowledgements

The authors are grateful to Dr.R.C.Goyal, Ex. Professor and Head, Department of Community Medicine, Rural Medical college, Loni for his constant inspiration, encouragement, expert guidance and total involvement in the study. The teaching faculty of Community Medicine and the respondents of the study are greatly acknowledged for their cooperation.

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Left unilateral super numerary renal artery- A case report

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Abstract
Variations of renal arteries are considered critical issues that surgeons should have through envision and appreciation of the condition. Variations of these vessels may influences urological, renal transplantation and laparoscopic surgeries. We present a case of left side unilateral accessory renal artery with a striking pre- hilar branching pattern noticed by routine dissection. The left kidney received two renal arteries from the aorta as a 02 separate trunks to hilar region. However the right side renal artery originated from the aorta as a single trunk and then followed by two divisions supplying the right kidney. Hear we discuss embryological origin and clinical aspects of accessory renal artery.

Key Words
Anatomical variations, Renal artery, supernumerary, aberrant renal artery.

Introduction
Accessory renal arteries are present 30% of the individuals, they supply upper or lower pole of the kidney either passing through the hilum or their poles. The variation renal arteries are among the most critical arterial variations that should be meticulously addressed by vascular and transplants surgeries.
Renal arteries branch laterally from the aorta just below the origin of the superior mesenteric artery. Both crosses the corresponding crus of the diaphragm at right angle to the aorta. The right renal artery is longer and often higher, passing posterior to the inferior vena cava, right renal vein and head of pancreas. A single renal artery to each kidney is present in 70% of individuals, the artery vary in their level of origin, their Calibri, obliquity and precise. In most individuals single renal artery arising from the aorta at the level of L2 vertebral body supplies each kidney. These usually arise from the aorta or iliac arteries any where at the level of T11 to L4 or rarely from the lumbar, suprarenal, celiac trunk. Superior accessory artery is a separate apical artery and an inferior accessory artery is a separate lower segmental artery. Accessory arteries to the Polar Regions are usually smaller than the accessory hilar arteries.

Case Report
During the routine dissection classes for I MBBS students in the department of Anatomy at RIMS Medical College, Kadapa, AP, and India. We noticed an accessory renal artery at the left side of aorta in a 40 years Male cadaver (Fig: 01.) It was located at lateral side of aorta originated directly from the aorta running parallel to the main trunk of renal artery and reaching the hilum. The length, Calibri is equal to the main trunk then it divides into 2 segmental arteries as a pre hilar pattern and reached the hilum. The left gonadal artery is originated from the left main trunk of renal artery.

Discussion
Regarding the increasing the number of renal transplantations, vascular re construction and urological surgery the Knowledge of the variations in renal artery supply is of significance. Renal arteries are terminal vessels which necessitate the removal of that segment of the kidney supplied by accessory arteries which its ligation. Accessory renal arteries tend tom be longer and narrower than main renal arteries resulting in lower perfusion pressure and higher resistance across artery. Accessory renal arteries most often pass anterior to the ureter so that pressure causes obstruction and hydro nephrosis. Renal blood supply undergoes successive changes during embryonic development that is concomitant with migration of metanephros.

Fig. 1: Arrow showing the left side supernumerary Renal Artery
References


Trichobezoar – A case report and literature overview
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Abstract
A sixteen year old girl was admitted in the hospital with a history of continuous pain and vomiting since five days. She has similar complaints intermittantly since five years. Other medical history, serum biochemistry and mental status were normal. On physical examination, a nontender mass was found in the epigastrium. Abdominal Ultrasound showed a dilated stomach filled with large ill defined mass with mixed echogenicity. Endoscopy showed a mass with entangled hair and stucked food particles. At laparotomy a large hair ball completely filling the stomach and extending to first part of duodenum is removed. The patient had a satisfactory post operative convalescence.

Keywords
Vomitting, Abdominal pain, Epigastric mass, Trichobezoar.

Introduction
Trichobezoar is rare but an interesting clinical entity. Some children develop the habit of swallowing hair from their dolls, brushes or blankets.¹ The word bezoar probably originated from the arab word 'bedzehr', the Persian word Padzahr or the Herbian word 'beluzzer', all meaning antidote.² Bezoars are concretions of human or vegetable fibres that accumulate in the gastrointestinal tract of which the most common type is trichobezoar (made of hair), while the other types are made up of vegetable or fruit fibre (phytobezoars), milkcurd (lactobezoars), or any indigestible material³. Most commonly present in adolescents and during second decade of life. We present a case of trichobezoar in a 16 year old female with normal mental and physical behavior.

Case Report
A 16 year old female girl was admitted with complaints of continuous vomiting and pain abdomen since 5 days. History revealed same complaints on and off previously since five years. Examination revealed an epigastric mass extending upto left hypochondrium which was hard and nontender. Routine blood counts, serum biochemistry and anthropometric measurements were normal. No hair loss was evident. X ray of abdomen showed a dilated stomach and Ultrasound showed dilated stomach filled with a mass of mixed echogenicity. Endoscopy showed a mass with hair and food particles stuck over it. Psychiatric examination was absolutely normal. Complementary history, gathered retrospectively revealed trichophagia. As breaking and dislodging of the mass during endoscopy failed, laparotomy was undertaken. At laparotomy a large hair ball, J shaped, filling the stomach, extending into the first part of duodenum (measuring 22/8/5 cms) was removed. It is black in colour, with stucked up food material and foul smelling odour. The post operative recovery was uneventful.

Discussion
For centuries, bezoars have been known to occur in the form of undigested masses found in the stomach of animals and humans⁴. There are several different form of presentations. It may be limited to esophagus, stomach or extend upto duodenum, jejunum or to Ileum. They may also occur as complications in clinical conditions that causes stasis within the gastrointestinal tract such as motility disorders and mechanical obstruction⁵. They may occur in persons with normal mental status with out any behavourial abnormality or may be associated with early childhood neglect or abuse or psychiatric disorders⁶. They grow slowly over many years and form a cast in the shape of the stomach, some times extending to the small bowel⁷. The engulfed hair which escape the peristaltic propulsion, get entrapped within the gastric mucosal folds and with increased accumulation, the hair getted matted up resulting in a ball formation due to churning action of stomach⁸. Halitosis may be present due to decomposition and fermentation of fats trapped in the interstices of the bezoar⁹. The acidic contents of the stomach denature the hair protein and give the bezoar it’s black colour¹⁰. Rapunzel Syndrome is a rare form of trichobezoar with various descriptions in the literature. Some define it as a gastric bezoar with tail extending upto jejunum or beyond to ileocaecal junction; some define it as a bezoar of any size which can cause intestinal obstruction¹¹,¹². Most of the cases are reported in the countries where women traditionally have long hair. The patients remain asymptomatic for many years. Symptoms develop as the bezoar increase in size to the point of obstruction. The common presenting features are abdominal pain, mass and vomiting. Other manifestations include upper digestive tract bleeding,
anemia and bowel intussusception. Though ten percent of patients have ulceration; perforating in about 30% of them, death is rare. Traditionally, a gastric trichobezoars were removed by gastrotomy through an upper midline laparatomy. Since the advent of minimally invasive surgery, surgeons now use laparoscopic techniques for removal of small to moderate sized bezoars.

Conclusion

Even though rarely seen in digestive tract diseases, the probability of bezoar should always be remembered in patients with vomiting and abdominal mass. After the removal of bezoar, precautions should be taken against recurrence by behavioural treatment.

References

A pilot study comparing Propofol-Thiopentone Admixture with Propofol-Ketamine Admixture for ambulatory anaesthesia at SIMS, Ghaziabad, Uttar Pradesh

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Abstract

In the present study, we aimed to compare the efficacy, tolerability, cardiovascular stability and recovery profile of propofol-thiopentone admixture with propofol-ketamine admixture in patient undergoing short surgical procedures. Sixty ASA grade I and II patients, age group 20-50 years of either sex scheduled for elective ambulatory surgical procedures were included. Patients were randomly allocated in two groups. Patients in group PT(n=30) were induced with admixture containing 10 ml of 1% propofol and 10 ml of 2.5% thiopentone and in group PK(n=30) were induced with admixture containing 10 ml of 1% propofol and 10 ml of 0.5% ketamine till loss of verbal response. Top up doses for maintenance were given as 2ml aliquots of the study drug. Induction dose and total maintenance dose of the drug, pain on injection, episode of apnoea, incidence of laryngospasm, bronchospasm, cough, hiccup and emergence phenomenon were recorded. Patients were monitored for pulse rate, non-invasive systolic and diastolic blood pressure throughout the procedure. Recovery and discharge time was recorded. Induction and total maintenance dose of propofol was significantly high in group PT as compare to group PK. Patients in group PK have more stable haemodynamic profile as compare to patients of group PT. Incidence of side effects like apnoea and pain on injection were more in patients of group PT. The recovery and discharge time of the patients of both groups were comparable so it was concluded that combination of propofol and ketamine gives better efficacy and tolerability in terms of dose requirement, cardiovascular stability, less frequent side effects without affecting discharge profile as compare to propofol and thiopentone admixture.

Keywords
Anaesthetics, Intravenous, Propofol, Thiopentone, Ketamine, Ambulatory anaesthesia.

Introduction
One of the most dramatic transformations in health care delivery during the past three decades is general anaesthesia for a very short duration, which can be achieved by short acting intravenous anaesthetic agents. This allows early ambulation and thus, is less disruptive to patients personal life; reducing separation from family and work, reduction of nosocomial infection and patient anxiety.

Propofol, a short acting intravenous inducing agent has emerged as ‘gold standard’ for day care anaesthesia but its tendency to cause apnoea, pain on injection and cardiovascular instability offsets its use. Various agents have been used in combination with propofol to make an ideal anaesthetic regime for ambulatory anaesthesia. Previously propofol has been combined with thiopentone¹ or ketamine². In this study we aimed to compare the efficacy, tolerability, cardiovascular stability and recovery profile of propofol-thiopentone admixture with propofol-ketamine admixture.

Material and Method
After institutional board approval 60 adult patients of age group 20-50 years, body weight between 35-80 Kg of either sex, ASA grade I and II who were scheduled for elective ambulatory surgical procedures were included in this study. A thorough pre-anæsthetic evaluation including complete history, clinical examination and investigations were done. Patients with history of hypertension, ischemic heart disease, psychiatric illness, on medication like psychotropic drugs or opioids, history of allergy to study drug were excluded. All patients were instructed for fasting of 8 hours preoperatively and each patient was thoroughly explained about the procedure and written consent was obtained. After taking patient in operation theatre base line pulse rate, blood pressure, respiratory rate and SpO₂ was recorded and continued throughout the surgical procedure. A good intravenous line established. Patients were premedicated with injection ondansetron 4mg five minutes before induction and were randomized in following two groups.

Group PT- Admixture containing 10 ml of 1% propofol and 10 ml of 2.5% thiopentone.

Group PK- Admixture containing 10 ml of 1% propofol and 10 ml of 0.5% ketamine.

The admixtures were indistinguishable from propofol visually. The study drugs were prepared freshly in the operation theatre by the anaesthesiologist who was not involved in the observations. Patients and observer were
kept blind about the drug administered. Patients were induced with the admixture according to their group at a rate of 4ml per 10 second till loss of verbal response and total induction dose was noted. Patients were maintained on oxygen and nitrous oxide (1:1) by Bain circuit throughout the procedure. Top up doses for maintenance of anaesthesia were given as 2ml aliquots of the study drug as and when required, which was judged by lighter plane of anaesthesia (increased heart rate, tearing and movement of limbs). Total amount of drug given for maintenance was also recorded and total duration of surgery in both groups was recorded.

Pain on injection, episode of apnoea, incidence of laryngospasm, bronchospasm was noted and lungs were manually ventilated by bain circuit if episode of apnoea occurred or Spo₂ decreased to less than 90%. Recovery time was calculated from last dose of drug till patient followed verbal command. All patients were shifted to recovery room and later on discharged when they met the standard guideline for safe discharge which includes stable vital signs, return to baseline orientation, ambulation without dizziness, minimal pain and PONV and minimal bleeding at surgical site. Data is presented as number of patients or mean±SD. Statistical analysis between groups was done by using unpaired ‘t’ test.

Result

Demographic data of patients of both groups were comparable (table-1). Preinduction baseline haemodynamic parameters were also comparable. The mean value of induction agent in group PT is significantly high (14.9±2.17ml) as compared to PK group (9.8±1.03ml). Similarly total maintenance dose of drug in group PT is significantly high (13.96±1.98ml) as compared to PK group (11.8±3.67ml) as shown in table 2. On comparison of recovery and discharge time between both groups, it was found that recovery and discharge time of patients in both groups were statistically not significant (table 3). Analysis of haemodynamic data during the procedure showed that there was highly significant fall in systolic blood pressure after induction in PT group, which recovered at 30 minutes after induction. In PK group, systolic blood pressure showed significant fall at 5 minutes after induction and which become non significant at 10 minutes (fig. 1). In group PT fall in diastolic blood pressure at 5, 10, 15, and 20 minutes was highly significant and recovered to non significant level at 30 minutes. Similarly, fall in diastolic blood pressure in group PK at 5, 10, 15, and 20 minutes was highly significant and recovered at 25 minutes (fig.2). In group PT pulse rate showed highly significant fall at 5 minutes after induction which showed a similar pattern through out the procedure and did not return back to baseline value till the completion of surgical procedure. In group PK pulse rate decreased at 5, 10, 15, 20 minutes which was highly significant statistically but, became non significant at recovery of patient.

Incidence of pain on injection of propofol was present in eight patients in group PT as compare to no pain on injection in group PK. Episodes of apnoea was present in four patients in group PT while there was no incidence of apnoea in group PK. Incidence of cough and hiccups was found in one patient of each group whereas emergence phenomenon occurred in one patient of group PK (table-4).

Discussion

Ambulatory surgeries offers a number of advantages like patient preference, lack of dependence on availability of hospital bed, greater flexibility in scheduling operations, low morbidity and mortality, lower incidence of infection, lower incidence of respiratory complications, greater

Table 1: Demographic profile

<table>
<thead>
<tr>
<th></th>
<th>Group PT</th>
<th>Group B</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mean±sd)</td>
<td>30.16±9.26</td>
<td>31.20±8.76</td>
<td>NS</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>14/16</td>
<td>18/12</td>
<td></td>
</tr>
<tr>
<td>Weight(kg)</td>
<td>53.16±9.52</td>
<td>51.24±10.4</td>
<td>NS</td>
</tr>
<tr>
<td>Duration of surgery(min.) mean±sd</td>
<td>22.24±3.82</td>
<td>21.46±4.31</td>
<td>NS</td>
</tr>
</tbody>
</table>

Table 2: Induction and total maintenance dose of admixtures

<table>
<thead>
<tr>
<th></th>
<th>Group PT</th>
<th>Group PK</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bolus dose(ml) (induction dose)</td>
<td>14.9±2.17</td>
<td>9.8±1.03</td>
<td>Significant</td>
</tr>
<tr>
<td>Total top up dose(ml)</td>
<td>13.96±1.98</td>
<td>11.8±3.67</td>
<td>Significant</td>
</tr>
</tbody>
</table>
efficiency, short surgical waiting lists and low procedure cost. Because of this ambulatory surgeries are gaining popularity now a day. The rapid growth of ambulatory surgeries would have not been possible without the changing role of anaesthesiologist and development of more highly titrable anaesthetic drugs. The availability of rapid shorter acting anaesthetic, analgesic and muscle relaxant drugs has facilitated the recovery process, allowing more procedures to be performed under day care anaesthesia. Propofol, a newer short acting intravenous inducing agent, is preferred because of its fast recovery and minimum post operative complications. But, propofol has some potential disadvantages which can be offset by combining it with other inducing agents like thiopentone, ketamine and opioids. Admixture of thiopentone and propofol is compatible and stable. This admixture does not support the growth of microorganisms despite the presence of nutrients in the admixture. Admixture of thiopentone and propofol has been previously used by Naguib M and et al. Both authors said admixture of thiopentone and propofol results in additive hypnotic effect, a reduction in pain of injection and reduced hypotensive response while Chilvers. M et al. said that addition of thiopentone to propofol does not delay recovery from anaesthesia and even does not increase post operative analgesic or antiemetic requirement. W.H.Wong also studied hypnotic effect of propofol and thiopentone and concluded that effect is additive at dose ranging for induction of anaesthesia.

Addition of ketamine to propofol in sub anaesthetic doses has gained attention in ambulatory anaesthesia because of its analgesic properties. Psychotomimetic effects of ketamine at analgesic doses (sub hypnotic) have also been reported. These psychotomimetic side effects can be minimised by administration of propofol.

Sympathomimetic effect of ketamine can be attenuated when administered in combination with propofol. This haemodynamic stability of propofol and ketamine combination makes it more suitable for use during ambulatory anaesthesia. Respiratory depression with small doses of ketamine is very minimal.

In this study, we did a prospective randomised study of 60 patients comparing propofol-thiopentone admixture with propofol-ketamine admixture in patients undergoing short surgical procedures. In our study dose of propofol required for induction as well as total dose of propofol for maintenance was less in PK group. The hypnotic and anaesthetic action of ketamine-propofol were found to be additive. Because of lesser degree of propofol required along with ketamine, duration of recovery and discharge of the patients was not prolonged as compared to propofol-thiopentone group.

Per operative haemodynamic variable were found to be more stable in group PK, this may be because of cardio stimulant effects of ketamine counterbalanced by depressive effects of propofol which is in accordance to T.W.Hue and Kaushik Saha. The fall in blood pressure in group PT following induction appears to be due to vasodilatation and myocardial depression done by both drugs.

Pain of injection of propofol is attenuated by ketamine due to its central analgesic as well as peripheral action.

In our study, there was no incidence of pain on injection of propofol in group PK as compare to eight patients in group PT. Incidence of apnoea was absent in group PK as compared to 4 patients in PT group. Emergence phenomenon occur in only one patient of group PK while there was no significant difference in discharge time of both group patients which is more important in ambulatory anaesthesia.

In conclusion, combination of propofol and ketamine gives better efficacy and tolerability in terms of dose requirement, cardiovascular stability, less frequent side effects without affecting discharge profile as compare to propofol-thiopentone admixture.

Interest of conflict- None

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Comparative analysis of variation in intraocular tension with vecuronium bromide and atracurium besylate: A prospective study

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Abstract

Intraocular pressure is the pressure exerted by the contents of the eye against its containing wall. During general anaesthesia, the intraocular pressure can increase, rendering problems in ophthalmic surgery especially surgery on the eye bulb. Pre-treatment with non-depolarising muscle relaxant prevents fasciculation and so the intraocular pressure does not rise. This is a prospective randomized study of 100 patients divided into two equal groups, where we have compared the effect of vecuronium bromide and atracurium besylate on intraocular pressure. The intraocular pressure was measured before surgery, at the time of intubation and at every 10 minutes interval during surgery. All the data were analysed statistically using unpaired student t test. We concluded that vecuronium bromide has better control over intraocular pressure as compared to atracurium besylate.

Key Words

Vecuronium bromide, Atracurium besylate, Intraocular pressure.

Introduction

Raised intraocular pressure during ophthalmic surgery may damage the contents of orbit, so it is mandatory to control the intraocular pressure during surgery. The main problems of general anaesthesia are the potential risk of coughing and straining post-operatively. There is also incidence of nausea and vomiting. These multifactorial factors may precipitate rise in I.O.P. Gold Smith reported that there is marked rise in I.O.P. following intubation with gallamine as muscle relaxant¹. According to Hollyoway, the general anaesthesia for intraocular ophthalmic procedures should provide stable intraocular pressure². It has been observed that administration of vecuronium during steady state anaesthesia was associated with significant decrease in I.O.P. during normal sequence of induction following significant reduction after thiopentone³. The effect of atracurium drug on I.O.P. was studied by Al-Albarek and Samuel and it was observed that the use of neuromuscular blocking drug without cardiovascular side effects have some advantages⁴; although James et al concluded that atracurium when administered during steady state anaesthesia had no significant effect on intraocular pressure⁵. The present work has been done to compare the effect of vecuronium and atracurium on intraocular pressure, so that the better drug could be use during opthalmic surgery.

Patients & Method

This study had been conducted in Saraswathi Institute of Medical Sciences, Hapur after obtaining the permission from the institutional ethical committee. The study had been conducted on patients between the age group of 18-50 years of ASA grade I admitted for various routine surgical procedures from September ‘09 to September ‘10. The patients with raised intraocular tension were excluded from this study. This is a prospective analysis of 100 patients divided randomly in to two equal groups. All the patients underwent a thorough pre-anesthetic assessment and ASA grade I fit patients were explained about the procedure and informed written consent taken. All the patients were given tablet diazepam 10mg orally the night before surgery and were pre-medicated with intramuscular glycopyrrolate 0.2 mg/kg and pethidine hydrochloride 1.5 mg/kg about 45 minutes before surgery. The baseline intra ocular pressure was recorded in both eyes with Schiotz tonometer on the operation table, before the induction of anaesthesia. The pressure was measured by Schiotz tonometer with 5.5 gm plunger weight after anaesthetizing each eye of the patients with 4% xylocaine 1-2 drops in both eyes 3-4 times. The patients were pre-oxygenated for 3-5 minutes with 100% oxygen and then induced with intramuscular glycopyrrolate 0.2 mg/kg and pethidine hydrochloride 1.5 mg/kg about 45 minutes before surgery. The baseline intra ocular pressure was recorded in both eyes with Schiotz tonometer on the operation table, before the induction of anaesthesia. The pressure was measured by Schiotz tonometer with 5.5 gm plunger weight after anaesthetizing each eye of the patients with 4% xylocaine 1-2 drops in both eyes 3-4 times. The patients were pre-oxygenated for 3-5 minutes with 100% oxygen and then induced with 2.5% thiopentone sodium 4-5 mg/kg intravenously after the administration of intubating dose of muscle relaxant. The group I patients received vecuronium bromide, whereas group II received atracurium besylate as muscle relaxant for intubation. The dose of vecuronium bromide was 0.1 mg/kg whereas for atracurium besylate 0.5mg/kg. The intraocular pressure was measured just after induction with thiopentone sodium, immediately after intubation, at every 10 minutes interval after intubation and after reversal. After the surgery was completed, all the patients were reversed with 0.4mg of glycopyrrolate followed by 2.5 mg neostigmine sulphate. After extubation, the patients were ventilated with 100% oxygen with the help of mask, so that increase in intraocular pressure could be prevented. All the data were analysed statistically with the help of unpaired Student t test. The aim of this study
was to compare the effect of these two non-depolarising types of muscle relaxants on intraocular pressure.

**Observation**

Out of 50 patients receiving vecuronium bromide, 20(40%) were male and 30(60%) were female. Maximum number of the patients i.e. 14(28%) were female in the age group of 20-30 years. In case of atracurium besylate group, the male were 26(52%) and female were 24(48%) and the maximum number of patients 12(24%) were female in the age group of 31-40 years (Table-1).

<table>
<thead>
<tr>
<th>Table 1: Showing Age &amp; Sex Distribution</th>
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<tbody>
<tr>
<td>Age (Yrs)</td>
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<td></td>
</tr>
<tr>
<td>20-30</td>
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<tr>
<td>31-40</td>
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<tr>
<td>41-50</td>
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</tbody>
</table>

The maximum number of 12(24%) female patients were in the weight group of 30-45 and 46-50 kgs had received vecuronium bromide, where as the higher figure (16%) in male patients was encountered in 46-50 kgs age group. In case of atracurium besylate group, the maximum female patients (24%) were in weight group of 30-45 kgs and male patients (22%) were in weight group of 51-60 kgs (Table-2).

The changes in intraocular pressure were highly significant in 5 minutes after intubation in patients who had received either vecuronium bromide or atracurium besylate. Although significant values were seen after 20 minutes of intubation in vecuronium bromide group, but there was no significance in different phases of anaesthesia in both groups (Table-3).

<table>
<thead>
<tr>
<th>Table 2: Showing Weight Wise Distribution</th>
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<tbody>
<tr>
<td>Weight (Kg)</td>
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<td></td>
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<tr>
<td>30-45</td>
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<tr>
<td>46-50</td>
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<td>51-60</td>
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<td>61-70</td>
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**Discussion**

An increase in intraocular pressure during general anaesthesia has been observed since long and this rise in intraocular pressure has always been a matter of anxiety for both surgeons and anesthesiologists during ophthalmic surgery. The results of ophthalmic surgery, especially when the globe is opened depend on good control of intraocular pressure. This is usually achieved with controlled ventilation and by the use of non-depolarising muscle relaxants. The patient's age, sex and body weight had no significant effects on intraocular pressure.

The patients who were intubated with vecuronium bromide, the pre-induction mean intraocular pressure was 15.91±0.22 mm of Hg, and 5 minutes after intubation it became 18.9±0.297 which is statistically highly significant. After 10 minutes of intubation there was insignificant fall in I.O.P but after 20 minutes of intubation there was significant fall. These findings were in accordance with Mirkhur RK et al3 and Jantzaj JP et al7. In the study conducted by Mirakhur RL et al3 the effect of vecuronium on I.O.P were investigated in doses of 0.15mg/kg as a part of rapid sequence induction with vercuronium administered prior to thiopentone sodium and intubation was done 30 seconds after induction. The I.O.P was measured after thiopentone sodium administration, immediately after intubation 2 minutes after when the study was terminated.

<table>
<thead>
<tr>
<th>Table 3: Showing Changes in Variations of Intraocular Pressure in Different Intervals</th>
</tr>
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<tbody>
<tr>
<td>GROUP</td>
</tr>
<tr>
<td>------</td>
</tr>
<tr>
<td>VECURONIUM BROMIDE</td>
</tr>
<tr>
<td>VECURONIUM BROMIDE</td>
</tr>
<tr>
<td>ATRACURIUM BESYLATE</td>
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<tr>
<td>VECURONIUM BROMIDE</td>
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</table>
Thiopentone sodium was associated with significant decrease in intraocular pressure; tracheal intubation was associated with increase I.O.P., over the next 2 minutes I.O.P. decreased significantly below the control level. Jantzaj et al. postulated that I.O.P. decreased by 22.6% in association with neuromuscular blockade produced by vecuronium 0.1 mg/kg. This appeared to be the result of an indirect action via an effect of central venous pressure. Vecuronium would be suitable for patients undergoing eye surgery where an increase I.O.P. would be undesirable. The patients who were intubated with atracurium besylate, the pre induction mean I.O.P. was 15.97±0.22 mm of Hg and there was a rise in mean I.O.P. of 18.90±0.297 after 5 minutes of intubation, which is statistically highly significant. Similar findings were observed by Payne et al. According to their study laryngoscopy and intubation can cause a significant rise in intraocular pressure. The intraocular pressure was decreased continuously in 10 and 20 minutes after intubation, but it was statistically insignificant. These findings were similar to the findings of Maharaj RJ et al. and Murphy DF et al. According to Maharaj RJ et al., the effect of atracurium on I.O.P. after pre-medication of patients with intramuscular morphine and oral diazepam 1 hour before surgery is compared with pancuronium. The anaesthesia was induced with thiopentone sodium and intubated without prior use of muscle relaxant. The I.O.P. was measured 5, 10, 15 and 20 minutes after induction, it was found that mean I.O.P. decreased 20 minutes after induction of anaesthesia and atracurium did not produce any deleterious effect on I.O.P. According to the study conducted by Murphy DF, atracurium has significant effect on I.O.P. after intubation. He concluded that atracurium provides an acceptable choice for ophthalmic surgery.

So with vecuronium bromide, there was highly significant rise in intraocular pressure in 5 minutes after intubation. Although there was fall in I.O.P. after 10 minutes of intubation insignificantly, but after 20 minutes of intubation, the pressure was below the pre-induction level. In case of atracurium, after 5 minutes of intubation, there was highly significant rise in I.O.P. The fall in pressure was insignificant after 10 and 20 minutes of intubation. After reversal the rise in pressure was insignificant in both groups.

**Conclusion**

This study reveals that both non-depolarising muscle relaxants had significant rise in intraocular pressure after intubation which gradually decreases at or below the preinduction value. We observed that vecuronium had a better control over intraocular pressure than atracurium.

**References**


A comparison of post operative analgesia and adverse effects produced by neostigmine and morphine when given intrathecally

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¹Assistant Professor, ²Associate Professor, Department of Anaesthesia, Saraswathi Institute of Medical Sciences, Hapur, Ghaziabad
245304

Abstract

Various modalities had been tried for optimum duration of post operative analgesia with minimal side effects. The present study was undertaken to evaluate the duration and quality of post operative analgesia by intrathecal bupivacaine alone, intrathecal bupivacaine12.5mg (0.5% heavy) with neostigmine (50µg) or intrathecal bupivacaine12.5mg (0.5% heavy) with morphine (100µg) in spinal block. 150 patients of ASA physical status 1&2, age 20-70 years of either sex, body weight 50-100Kg were randomly assigned in three study groups. Group I - 0.5% bupivacaine (2.5ml bupivacaine+0.5ml normal saline), group II - 0.5% bupivacaine with 50µg neostigmine (2.5ml bupivacaine+0.5ml neostigmine) and group III - 0.5% bupivacaine with morphine100µg (2.5 ml bupivacaine+0.5 ml morphine). Onset of sensory and motor block of intrathecal neostigmine and intrathecal morphine group was much faster as compare to control group. Duration of post operative analgesia was also more in intrathecal neostigmine and intrathecal morphine group as compare to control group. Intrathecal neostigmine and intrathecal morphine groups were comparable in terms of onset of sensory and motor block and level of highest block. Nausea, vomiting and bradycardia were more in neostigmine group as compared to control group. Patients in intrathecal morphine group had more incidence of pruritus. We concluded that intrathecal morphine 100µg is a better choice than intrathecal neostigmine during spinal block.

Key Words

Anaesthesia, Spinal anaesthesia, Drugs, Bupivacaine, Neostigmine, Morphine.

Introduction

Post operative pain control is essential for optimal care of surgical patients. If post operative pain is not controlled effectively it results in respiratory, cardiac, gastrointestinal, urinary and psychological adverse effects. Management of post operative pain remains a point of concern with spinal anaesthesia.

Intrathecal neostigmine inhibits the breakdown of endogenous spinal neurotransmitter acetylcholine which augments analgesic duration of bupivacaine. But higher doses of intrathecal neostigmine is associated with increased incidence of nausea, vomiting and bradycardia. Intrathecal morphine also increases the duration of analgesia of bupivacaine but is associated with post operative respiratory depression, pruritus and urinary retention.

The present study was undertaken to evaluate the duration and quality of post operative analgesia by intrathecal bupivacaine alone, intrathecal bupivacaine12.5mg (0.5% heavy) with neostigmine (50µg) or intrathecal bupivacaine12.5mg (0.5% heavy) with morphine (100µg) in spinal block.

Material and Methods

After approval with institutional review board, 150 patients of ASA physical status 1&2, age 20-70 years of either sex, body weight 50-100Kg, scheduled for lower abdominal & lower limb surgeries were enrolled in the study. Informed consent was taken from each and every patient. Patients were randomly assigned using coded envelop to one of the three study group receiving either 0.5% bupivacaine (2.5ml bupivacaine+0.5ml normal saline) group I, 0.5% bupivacaine with 50µg neostigmine (2.5ml bupivacaine+0.5ml neostigmine) group II or 0.5% bupivacaine with morphine100µg (2.5 ml bupivacaine+0.5 ml morphine) group III in a double blind fashion. Patients with known contraindications of spinal anaesthesia, weight more than 100Kg and known allergy to study drug were excluded from the study.

Pre anaesthetic check up of all patients was done one day prior to surgery and patients were kept fasting for 8 hours prior to surgery. All patients were given tablet alprazolam 0.5 mg night before surgery. Pulse rate, arterial oxygen saturation, Non invasive Blood Pressure, E.C.G and respiratory rate were monitored with B.P.L. excelleo monitor on arrival in O.T. and continued till end of surgery. A good intravenous line established and patients were preloaded with 10ml per Kg of ringer lactate solution. Patients were placed in sitting position and under all aseptic precaution spinal anaesthesia was administered with 25G Quincke’s needle in L₂-L₃ or L₃-L₄ interspace and after obtaining free flow of C.S.F, test drug was given according to patient’s assigned group. Patient was immediately turned supine and oxygen was given by
face mask at 4 litre per minute.

Level of sensory block was recorded by using pin prick method and motor block was assessed by modified bromage scale.

Time of onset of block was recorded as time taken from intrathecal drug administration to loss of pin prick at T₁₀ and highest level of block was noted.

Duration of surgical analgesia was defined as the period between intrathecal injection and recovery from block, both sensory and motor. In post op period pain score was assessed by VAS every 30 minute till 4 hours and than hourly for up to the period of 12 hours. When pain score was more than 4 analgesia was supplemented by intramuscular diclofenac sodium in dose of 75mg and time of drug given was recorded.

Side effects like hypotension, bradycardia, nausea, vomiting, sweating, sedation, pruritus, increased salivation, urinary incontinence and nystagmus were recorded and treated accordingly. If the systolic B.P. decreased more than 25% from base line value or less than 90mm of Hg the injection mephentermine 6mg intravenously was given. Bradycardia (H.R. less than 55 per minute) was treated with injection atropine sulphate 0.3 to 0.6 mg intravenously. Nausea and vomiting was treated with injection ondansetron 4mg intravenously.

All data were analysed by student t test.

Results

All the groups were statistically comparable regarding demographic profile (Table I). Surgeries done were appendicectomy, herniorrhaphy, hernioplasty, prostectomy, vaginal hysterectomy, and all lower limb orthopaedic procedure. Mean onset time of sensory as well as motor block was faster in group II and group III as compared to group I (table 2). Onset time of sensory and motor block of both groups (group II and group III) was statistically highly significant as compared to group I. On comparison between group II and group III, onset time of sensory and motor block was not significant.

Level of highest block of all 150 patients is shown in table 3. All patients were able to complete their surgical procedures well. Patients who received intrathecal morphine or intrathecal neostigmine had highly significant prolonged duration of analgesia as compared to control group. Time in hours from spinal anaesthesia to first analgesic requirement (VAS > 4) of all three groups is shown in table 4. Duration of analgesia of group II and group III was comparable.

In the present study there was fall in systolic and diastolic blood pressure during the study period from their base line values. Some patients required injection mephentermine also, but on comparison between the three groups the decrease in systolic and diastolic blood pressure was found to be insignificant. Pulse rate also varied in different point of time but was statistically insignificant on comparison between three groups (Table 5).

In our study side effects including nausea, vomiting and bradycardia were more in neostigmine group in comparison to control group. Incidence of nausea and vomiting in morphine group was comparable to control group. 10 patients in morphine group had pruritis, while no patient in control group and neostigmine group had pruritis. Other side effects like sweating, sedation, respiratory depression, increased salivation, urinary incontinence and nystagmus were not found in any patient of three groups (Table 6).

Table 1: Demographic profile.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>40±16</td>
<td>42±16</td>
<td>43±18</td>
</tr>
<tr>
<td>Sex</td>
<td>22/28</td>
<td>20/30</td>
<td>23/27</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>77±13</td>
<td>72±11</td>
<td>75±14</td>
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</tbody>
</table>

Table 2: Duration of onset of sensory and motor block (Mean ± SD).

<table>
<thead>
<tr>
<th>Group</th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensory block (min.)</td>
<td>5.64±.5834</td>
<td>3.45±.7223</td>
<td>3.24±.5850</td>
</tr>
<tr>
<td>p value (as compared to Gp I)</td>
<td>&lt; .001</td>
<td>&lt; .001</td>
<td></td>
</tr>
<tr>
<td>Motor block (min.)</td>
<td>6.5±.5744</td>
<td>4.05±.7697</td>
<td>3.82±.6615</td>
</tr>
<tr>
<td>p value (as compared to Gp I)</td>
<td>&lt; .001</td>
<td>&lt; .001</td>
<td></td>
</tr>
</tbody>
</table>

Table 3: Level of highest sensory block in all 150 patients.

<table>
<thead>
<tr>
<th></th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
</tr>
</thead>
<tbody>
<tr>
<td>T₆</td>
<td>7</td>
<td>8</td>
<td>8</td>
</tr>
<tr>
<td>T₈</td>
<td>17</td>
<td>20</td>
<td>19</td>
</tr>
<tr>
<td>T₁₀</td>
<td>26</td>
<td>22</td>
<td>23</td>
</tr>
<tr>
<td>T₁₂</td>
<td>0</td>
<td>0</td>
<td>0</td>
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</table>

Table 4: Mean duration of analgesia (Time in hours).

<table>
<thead>
<tr>
<th></th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
</tr>
</thead>
<tbody>
<tr>
<td>When VAS &gt; 4 (Mean± SD)</td>
<td>3.46 ± 0.3852</td>
<td>6.66 ±0.6724</td>
<td>6.68±0.5179</td>
</tr>
<tr>
<td>p value (as compared to Gp I)</td>
<td>&lt; .001</td>
<td>&lt; .001</td>
<td></td>
</tr>
</tbody>
</table>

Table 6: Side effects (%) in all three groups

<table>
<thead>
<tr>
<th>Side effects</th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
</tr>
</thead>
<tbody>
<tr>
<td>NO.</td>
<td>%</td>
<td>NO.</td>
<td>%</td>
</tr>
<tr>
<td>Bradycardia</td>
<td>3</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>Nausea</td>
<td>4</td>
<td>8</td>
<td>22</td>
</tr>
<tr>
<td>Vomiting</td>
<td>2</td>
<td>4</td>
<td>11</td>
</tr>
<tr>
<td>Pruritis</td>
<td>0</td>
<td>0</td>
<td>0</td>
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</table>
following the surgery. Morphine that is needed during the first 24 hours and significantly decrease in the amount of intravenous ranging from 0.1 to 0.3 mg, provide adequate analgesia postoperative pain control after knee and hip arthroplasty. Rathnell and his team did a dose ranging study for 156 Indian Journal of Public Health Research & Development. April - June, 2012, Vol. 3, No. 2.

Intrathecal morphine has been studied by various investigators for pain control after caesarean delivery. Intrathecal neostigmine and intrathecal morphine was compared to control group. Onset time of sensory and motor block of intrathecal neostigmine and intrathecal morphine was comparable.

Results of our study demonstrated that onset of sensory and motor block is enhanced by both intrathecal neostigmine and intrathecal morphine as compared to control group. Onset time of sensory and motor block of intrathecal neostigmine and intrathecal morphine was comparable.

Level of sensory block achieved in our study groups was similar in all three groups suggesting that spread of subarachnoid bupivacaine was not affected by adjuvants. All patients in our study were able to complete their procedures well.

Mean duration of analgesia in intrathecal neostigmine group was 6.66 hours and in intrathecal morphine group 6.68 hours as compared to 3.46 hours in control group, which is highly significant. Chung et al and Lauretti et al also observed statistically significant lower VAS score in the doses ranging from 25-75µg neostigmine group compared to saline group.

P-H Tan et al compared intrathecal morphine (300µg) and intrathecal neostigmine (50µg) and found that intrathecal neostigmine and intrathecal morphine prolonged the effect of spinal anaesthesia but the time of first requirement of analgesia in neostigmine group

Discussion

Spinal anaesthesia using intrathecal bupivacaine is a very safe technique of anaesthesia for lower abdominal and lower limb procedures lasting for one to two hours. Spinal anaesthesia over general anaesthesia is preferred by anaesthesiologist and surgical staff, because spinal anaesthesia is associated with less complications as compared to general anaesthesia.

Management of post –operative pain remains a point of concern with spinal anaesthesia, optimum pain relief with minimal side effects can be achieved by adding various adjuvants like opioids, clonidine, beclofen, ketamine, midazolam and neostigmine.

The present study was undertaken to assess the efficacy of intrathecal neostigmine and intrathecal morphine with heavy bupivacaine with aim of maximising analgesic duration while minimising the possibilities of potential side effects.

Intrathecal morphine has been studied by various investigators for pain control after caesarean delivery, various orthopaedics procedures, spinal surgeries, cardiac surgeries and other surgical procedures. Optimal dose of intrathecal morphine for pain control seems to be 100 µg with minimal side effects. Doses above 100 µg is often associated with nausea, vomiting, pruritus, urinary retention and respiratory depression.

Rathnell and his team did a dose ranging study for postoperative pain control after knee and hip arthroplasty and concluded that low doses of intrathecal morphine, ranging from 0.1 to 0.3 mg, provide adequate analgesia and significantly decrease in the amount of intravenous morphine that is needed during the first 24 hours following the surgery.

Motamed and colleagues suggested that a low dose of spinal morphine combined with a low dose of bupivacaine can be a simple and trustworthy component of a multimodal approach to surgical anesthesia.

Hooq DD et al in 1995 used intrathecal neostigmine in healthy human beings. Neostigmine, a cholinesterase inhibitor inhibits the metabolism of spinally released acetylcholine and thus enhances analgesia. Previous studies had used intrathecal neostigmine in dose ranges of 10 to 200µg. Spinal cord toxicity resulting from intrathecal neostigmine has not been reported.

Table 5: Showing haemodynamic variables : Pulse rate, Systolic blood pressure and diastolic blood pressure (mean ± S.D.)

<table>
<thead>
<tr>
<th></th>
<th>Preop</th>
<th>5 min</th>
<th>10 min</th>
<th>15 min</th>
<th>20 min</th>
<th>30 min</th>
<th>45 min</th>
<th>60 min</th>
<th>90 min</th>
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<td><strong>Pulse Rate</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Gp I</td>
<td>85.0±</td>
<td>86.6±</td>
<td>87.5±</td>
<td>87.7±</td>
<td>88.4±</td>
<td>87.9±</td>
<td>87.2±</td>
<td>86.6±</td>
<td>86.3±</td>
</tr>
<tr>
<td>Gp II</td>
<td>84.5±</td>
<td>85.4±</td>
<td>85.0±</td>
<td>86.1±</td>
<td>87.5±</td>
<td>86.1±</td>
<td>86.3±</td>
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<td>85.9±</td>
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<td>7.73±</td>
<td>8.10±</td>
<td>8.34±</td>
<td>8.50±</td>
<td>7.81±</td>
<td>7.92±</td>
<td>8.24±</td>
<td>8.25±</td>
<td>7.58±</td>
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<tr>
<td>Gp III</td>
<td>86.1±</td>
<td>86.18±</td>
<td>87.68±</td>
<td>87.82±</td>
<td>88.12±</td>
<td>87.54±</td>
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<td>7.03±</td>
<td>7.41±</td>
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<td>6.39±</td>
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<td><strong>Systolic B.P.</strong></td>
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<tr>
<td>Gp I</td>
<td>135.0±</td>
<td>126.4±</td>
<td>119.2±</td>
<td>117.3±</td>
<td>122.9±</td>
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<tr>
<td>10.21±</td>
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<tr>
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<td>122.9±</td>
<td>121.5±</td>
<td>120.5±</td>
<td>125.4±</td>
<td>126.9±</td>
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<tr>
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<td>130.4±</td>
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<tr>
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<td>11.28±</td>
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<td>8.38±</td>
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<td>7.79±</td>
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<td>8.39±</td>
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<tr>
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</tr>
<tr>
<td>Gp I</td>
<td>84.0±</td>
<td>82.8±</td>
<td>82.6±</td>
<td>81.6±</td>
<td>81.5±</td>
<td>81.9±</td>
<td>81.8±</td>
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<td>5.74±</td>
<td>5.36±</td>
<td>5.89±</td>
<td>5.34±</td>
<td>5.29±</td>
<td>5.13±</td>
<td>6.13±</td>
<td>6.4±</td>
<td>4.73±</td>
<td></td>
</tr>
<tr>
<td>Gp II</td>
<td>82.9±</td>
<td>82.0±</td>
<td>81.4±</td>
<td>80.3±</td>
<td>80.7±</td>
<td>81.0±</td>
<td>81.0±</td>
<td>81.0±</td>
<td>81.6±</td>
</tr>
<tr>
<td>5.73±</td>
<td>5.64±</td>
<td>5.68±</td>
<td>5.48±</td>
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<tr>
<td>Gp III</td>
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<td>82.0±</td>
<td>81.0±</td>
<td>80.6±</td>
<td>81.0±</td>
<td>81.6±</td>
<td>81.6±</td>
<td>82.5±</td>
<td>82.6±</td>
</tr>
<tr>
<td>6.44±</td>
<td>6.30±</td>
<td>6.06±</td>
<td>5.56±</td>
<td>5.28±</td>
<td>5.62±</td>
<td>5.96±</td>
<td>5.64±</td>
<td>6.02±</td>
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</table>
was shorter than the morphine group\textsuperscript{17}. The results are in accordance to our study except for prolonged analgesic effect in post operative period in intrathecal morphine group, which may be because of higher doses of morphine (300µg) used.

Almeida et al also concluded that intravenous requirement of ketoprofen during first postoperative 24 hours was less in intrathecal morphine group as compared to intrathecal bupivacaine only\textsuperscript{9}.

Chung et al and Muhammad ishaque et al also reported that 25µg of intrathecal neostigmine and 100µg of intrathecal morphine produce analgesia of almost similar duration\textsuperscript{13,18}. These results are in accordance to our study. Study done by Klamt et al is also in accordance to our study\textsuperscript{19}.

Intrathecal neostigmine produce nausea and vomiting in dose dependent manner, due to cephalad migration of neostigmine to brain stem and accumulation at CTZ induces vomiting. Klamt et al, Hood et al and Tan et al reported that droperidol and metoclopropamide were not effective in controlling nausea and vomiting\textsuperscript{10,11,17}. Nausea and vomiting resolved with time. In our study 44% of patients complained of nausea and 22% of patients complained of vomiting which was effectively controlled by injection ondansetron(4mg) and injection dexamethasone(8mg) simultaneously. Nausea and vomiting in intrathecal morphine group was comparable to control group. Tan et al reported that there was no significant difference between intrathecal neostigmine and intrathecal morphine groups in the incidence of nausea and vomiting which may be due to high doses of morphine (300µg)\textsuperscript{17}.

Pruruitis was observed in 20 % of patients in intrathecal morphine group which was not distressing to patients and was relieved by intravenous injection of chlorpheniramine.

The serious side effect of respiratory depression was not found in any of patients in our study.

In conclusion, addition of intrathecal neostigmine or intrathecal morphine to heavy bupivacaine for spinal anaesthesia doubled the duration of requirement of rescue analgesia. Both drugs increased the duration of analgesia in post operative period on the expense of increased incidence of side effects. These side effects were effectively controlled by intravenous medications. Because of more incidence of side effects in intrathecal neostigmine group (nausea and vomiting) as compared to intrathecal morphine group (pruritis), we concluded that morphine 100 µg is a better choice than neostigmine 50µg when combined with bupivacaine (0.5%) during spinal block.

Interest of conflict- None

References

Nasopalatal Cyst – A case report

Shilpa B.J A¹, Kodhandarama G.S B², Nagarajappa D C³, Tanveer Ahmed D¹, Shivshankar C E⁴
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Abstract

Nasopalatal cyst is rare but most common among the developmental, non-odontogenic cysts of the jaws. It is located in the incisive canal or in the anterior palate. It arises from the embryonic ducts of scrapa and stenson. It accounts for about 5-10% of the jaw cysts and 1% of the population. Majority of the cases occur between fourth to sixth decades of life. Males are more commonly affected than the females. We report case of Nasopalatal cyst in 50 year old female.

Keywords

Nasopalatal duct cyst: Incisive canal cyst; Non-odontogenic cyst; Cysts of the hard palate.

Introduction

Nasopalatal cyst (NPC) also called as incisive canal cyst, Nasopalatal canal/duct cyst, median palatal cyst or median anterior maxillary cyst were regarded as fissural cyst in the past¹. At present, according to the classification of the WHO, these lesions are developmental, epithelial, non-odontogenic cysts of the maxilla along with nasolabial cysts². NPC was first described by Mayer in 1914³. NPC are rare but most commonest among non-odontogenic developmental cysts of the oral cavity occurring about 1% of population. The cyst is believed to arise from the remnants of the Nasopalatine duct, an embryonic structure connecting the oral and nasal cavities in the area of the incisive canal⁴. It is one of many pathologic processes that may occur within the jawbones, but it is unique in that it develops in only a single location, in the midline anterior maxilla.

Case Report

A 50 year old female patient reported to the out patient department of Oral Medicine and radiology, S.J.M. Dental College and Hospital, Chitradurga with the chief complaint of swelling in the anterior region of palate since 4 months. Patient noticed the swelling 4 months back which has gradually increased to present size. The swelling associated with discomfort in sipping the fruits, chocolate and ice candy, sometime associated with the numbness over the anterior palate. Extra-orally there was no detectable abnormality or lymphadenopathy.

Intra-oral examination revealed a solitary, well defined, oval shaped swelling measuring about 2x1.5cms in the anterior palate in the midline. Overlying mucosa was normal. The swelling was hard, tender and non mobile (Figure-1). On the basis of history, clinical findings a provisional diagnosis of nasopalatine cyst was made and a differential diagnosis of radicular cyst was considered. Maxillary anterior teeth were vital on heat vitality test which excluded the differential diagnosis of radicular cyst in relation to maxillary anterior teeth. Fine needle aspiration cytology (F.N.A.C) was done, straw colored fluid aspirated. The cytology revealed chronic inflammatory cells with few epithelial cells and RBC's. Maxillary occlusal radiograph revealed a solitary, well defined, ovoid radiolucency measuring about 2x1.5cms located in midline of the anterior maxilla below the roots of maxillary central incisors (Figure-2). Computed tomography (C.T) confirmed the presence of well defined, midline cystic lesion in the anterior portion of the hard palate measuring about 1.4x1.2 cms with bony expansion breaching the cortex and bulging into the oral cavity (Figure- 3 and 4). Deviated nasal septum to left side also noted

Figure 1: Intra oral photograph showing ovoid swelling over the hard palate.

Figure 2: Maxillary occlusal radiograph showing solitary, well defined, ovoid radiolucency located in midline of the anterior maxilla.
in the C.T. On the basis of the clinical and radiographic findings, a diagnosis of Nasopalatine cyst was made. All preliminary investigations were done and results were within normal limits. Then the cyst was enucleated under general anesthesia and the specimen was subjected for histopathological examination. Post operation session was uneventful. On microscopic examination, the cystic lining showing transition from columnar to cuboidal epithelium and underlying connective tissue showing chronic inflammatory cell infiltrate like lymphocytes and plasma cells (Figure- 5). The histological features were suggestive of nasopalatal cyst.

**Discussion**

Nasopalatal cysts are the most common among the developmental, epithelial, non-odontogenic cysts of the oral cavity, representing up to 1% of all maxillary cysts. NPCs develop in the bony nasopalatine channel located at the anterior tip of the suture line of the palatal apophyses of maxillary bones. Lower mammals have inside this conduit a permeable “nasopalatine duct” that acts as an ancillary olfactory organ (vomeronasal or Jacobson’s organ). Although NPCs were initially thought to originate from fissures, they are now believed to derive from incompletely developed epithelial remnants in the embryonic nasopalatine duct. The stimulus for cyst formation from the epithelial remnants of the nasopalatine canal is uncertain, although trauma, bacterial infection and mucin retention within the mucous gland lining are thought to have a role. Genetic factors have also been suggested.

Nasopalatal cyst has intra-osseous and extra-osseous variants. The extra-osseous cyst is termed as cyst of the incisive papilla. The incisive canal cyst includes median palatal cyst and anterior median maxillary cyst, although these once were thought to represent separate entities. NPCs may develop at any age but is most common in the fourth to sixth decades of life. Men are affected more often than women with ratio 3:1. Most of NPCs are asymptomatic or cause such minor symptoms that they are tolerated for very long time. The most common complains are palatal swelling, upper anterior teeth displacement, sublabial swelling, and low grade pain. The swelling usually fluctuant and blue if the cyst is near the surface. The deeper NPC is covered by normal appearing mucosa unless it is ulcerated from mastication. If cyst expands, it may penetrate the labial plate and produce swelling below the maxillary labial frenulum or to one side. The lesion also may bulge into nasal cavity and distort the nasal septum. The swelling usually fluctuant and blue if the cyst is near the surface. The deeper NPC is covered by normal appearing mucosa unless it is ulcerated from mastication. Various combinations of swelling, discharge and pain may occur. Discharge may be mucoid in which case the patient describes a salty taste or it may be purulent and patient may complain of a foul taste.

The definitive diagnosis of NPC is more easily made on conventional radiographs; other advanced imaging modalities are being used to differentiate this entity from other lesions. NPCs appear as a bilateral, symmetric, well defined, ovoid, round or inverted, heart shaped radiolucency with or without sclerotic borders located.
in the midline of the maxilla, which is inter-radicular and apical to the roots of the maxillary central incisors. It may cause root displacement and root divergence. Occasionally it causes root resorption, expansion of labial/palatal cortex and perforation of nasal floor. NPCs should be differentiated from large incisive foramen and periapical cyst/granuloma associated with maxillary incisors. Radiolucencies of the incisive canal measuring less than 0.6 cm in diameter should not be considered cystic in the absence of other symptoms. The periapical cyst/granuloma associated with maxillary incisors can be differentiated with the presence or absence of the lamina dura and enlargement of the periodontal ligament space around the root apex of the incisors and with the help of vitality tests. NPCs are usually treated by enucleation, in case of large cysts, marsupialization may be considered before definitive enucleation. Recurrence rate ranges from 0% to 11%.

**Conclusion**

Nasopalatal cysts are rare but commonest among the developmental, epithelial, non-odontogenic cysts of the jaws. The lesion may be asymptomatic or manifests as swelling, pain, and discharge from the hard palate. The clinician should differentiate it from large incisive foramen and periapical pathology of maxillary incisors and should include NPC in the differential diagnosis for swelling of hard palate.

**References**

Feeding practices and early childhood caries - A review

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Abstract
Early childhood caries (ECC) is defined as the presence of 1 or more decayed (noncavitated or cavitated lesions), missing (due to caries), or filled tooth surfaces in any primary tooth in a child 71 months of age or younger. It has a debilitating effect on the development, speech, general health, and self-esteem of infants. Predisposing factors for ECC is considered to be multifactorial and is still unclear. The association between feeding practices and caries remains controversial with some studies finding no correlation, while others showing relation. ECC is a complex entity in which feeding practices alone does not play a prime role but is influenced by a myriad of cultural, behavioral, biological and microbiological factors. So, the area has conflicting reports which needs further research.

Key Words
Feeding practices, early childhood caries, breast feeding, bottle feeding, pacifiers.

Introduction
Caries remains the principal dental disease affecting preschool children. Despite the success of preventive dentistry, there is growing number of reports showing decline in the quality of very young children's teeth. Early childhood caries (ECC), a disease characterized by rampant dental decay in the primary dentition of infants and young children, continues to pose a serious threat to child welfare.

The American Academy of Pediatric Dentistry (AAPD) recognizes ECC as a significant public health problem. It is particularly a virulent form of caries, beginning soon after dental eruption, developing on smooth surfaces, progressing rapidly and having a lasting detrimental impact on the dentition. Not only does ECC affect teeth, but consequences of this disease may lead to more widespread health issues. It can manifest itself in severe pain, infection, abscesses, chewing difficulty, malnutrition and gastrointestinal disorders. It leads to malocclusion by adversely affecting the correct guidance of the permanent dentition, cause poor speech articulation and low self-esteem.

ECC is a complex, multifactorial infectious disease. Feeding patterns are said to be of prime importance in the etiology of ECC. Infant feeding practice associated with the development of caries in the primary dentition is poorly understood and somewhat controversial. Feeding behaviors most commonly implicated are the prolonged and at-will breast feeding, falling asleep while feeding, prolonged and nocturnal bottle feeding habits, nursing beyond the recommended age for weaning, bottle feeding with sweetened beverages and the use of sweetened comforters. However, recent studies call into question the association between feeding practices and ECC2,3,4.

To fully understand the problem, it is important to investigate factors that have an influence on the quality of the teeth. Thus, an attempt is made to review the various feeding practices and study the effect of breast-feeding, bottle-feeding, comforters / pacifiers on ECC.

Definition
Early childhood caries (ECC) is defined as the presence of 1 or more decayed (noncavitated or cavitated lesions), missing (due to caries), or filled tooth surfaces in any primary tooth in a child 71 months of age or younger2.

In children younger than 3 years of age, any sign of smooth-surface caries is indicative of severe early childhood caries (S-ECC). From ages 3 through5,1 or more cavitated, missing (due to caries), or filled smooth surfaces in primary maxillary anterior teeth, or a decayed, missing, or filled score of ≥4 (age 3), ≥5 (age 4), or ≥6 (age 5) surfaces, constitutes S-ECC.

Explanation of the Caries Pattern6
A potentially cariogenic oral habit that begins soon after the child is born will affect the primary teeth as soon as they erupt. The maxillary incisors, which are the first to erupt, will be the first to experience the cariogenic challenge and for longer duration. Continuation of the habit will ultimately involve all the other teeth in the sequence as they erupt if the habit is not discontinued before age2.

During sucking, the natural or artificial nipple rests against the palate, while the tongue lies over the lower incisors. If the liquid is consumed frequently and for prolonged periods during the day or night, milk cannot be eliminated from the oral cavity and it pools around
the tooth surfaces. If the liquid contains a fermentable carbohydrate, in this stagnant acid environment, lesions can develop quickly. Thus the upper incisors are most ravaged. While the lower incisors, which are protected by the tongue and also by saliva from the mandibular salivary glands, remain unscathed. As the child alternately sleeps and sucks, a pool of stagnant liquid collects around the necks of the upper anterior teeth and the decalcification process continues. Moreover, the swallowing reflex is absent during sleep and the diminished salivary flow encourages the damage to the teeth. Continuation of the sucking habit on a daily or usually nightly basis for several months leads to the clinical picture.

**Feeding Practices and ECC**

All carious lesions, including those associated with ECC, result from the interaction among 3 variables: pathogenic microorganisms, fermentable carbohydrates that the microorganisms metabolize to organic acids, susceptible tooth and host. In order for the lesions to progress and to be clinically diagnosed, these 3 variables must interact over a suitable period of time.

**Fermentable Carbohydrates**

Sucrose or common table sugar is considered to be the major cariogenic food in the human diet. Other common sugars include the monosaccharides glucose and fructose, which are present in fruit and honey. A common practice is the use of a pacifier (also called a “dummy” or “comforter”), which has been dipped in a sugar solution or honey, a mixture of glucose and fructose. Another practice is the use of syrupy, sweet vitamin preparations added to nursing bottles or infant feeders.

**A. Milk and caries:** The only carbohydrate contained in milk is lactose. Several in vitro studies using milk suggest that this nutrient is not cariogenic. However, Vianna7, using an artificial mouth, showed that plain bovine milk produced the least decalcification, followed in order by milk formula, human milk, and milk and honey. His study strongly suggests that lactose plays a role in the decalciﬁying potential of milk.

1. **Human milk:** Compared to bovine milk, human milk contains a higher concentration of the carbohydrate lactose. There are now suggestions from case reports that prolonged and excessive breast-feeding is associated with caries prevalence. In spite of these reports, the issue of human milk being linked to ECC is complex. The calcium and phosphorus content of milk would contribute to the remineralization of enamel. Milk contains a number of proteins including casein, which could provide a protective organic coating on the enamel surface. Moreover, cariogenic bacteria may not be able to utilize lactose as an energy source as readily as sucrose.

2. **Bovine milk:** Lactose, a disaccharide sugar, is present in bovine milk at a concentration of around 4% and in human milk at around 7%. Although the potential demineralization from fermentation of lactose has been shown in some laboratory studies, these results may have only limited clinical significance as many of these tests were done under extreme conditions, or used very high concentrations of lactose. In summary, from the literature, there is no evidence to suggest that bovine milk is cariogenic. On the contrary, there is much evidence that milk is cariostatic. This suggests that, under usual dietary conditions, milk is not very cariogenic and may be caries protective. This conclusion does not contradict the clinical evidence indicating milk in cases of ECC, because in these children exposure to the milk is frequent and prolonged, resulting in conducive environment.

3. **Milk formulas:** The cariogenicity of milk formula has not been well investigated. The concentrations of the constituents in baby formula are similar to that of human milk, including the lactose content. In recent investigations, Erickson and co-workers10, using in vitro methods, reported that some common milk formulas have as much cariogenic potential as sucrose. In contrast, the results of Bowen et al using desalivated rats11, suggested that all milk formulas tested caused less caries compared with sucrose.

**B. Fruit juices, carbonated beverages and other sweet liquids:** Many reports have indicated that the content of the nursing bottle was frequently not just plain milk, but, in fact, the milk had been sweetened with other sugars such as table sugar or honey or sweet, syrupy vitamin formulations. Fruit juices and carbonated beverages contain a sugar (fructose) and are intrinsically acidic; erosion may be the primary enamel change preceding the rampant caries.

**C. Sweetened pacifiers:** Nonnutritive sucking can be just as damaging to the teeth as nutritive sucking when a pacifier is sweetened with a decay-promoting agent.

**Frequency of Consumption**

There are now many studies which suggest that children with ECC have a high frequency of sugar consumption in the nursing bottle and also sweetened solid foods13,14. Increased frequency of eating sucrose increases the acidity of plaque, and enhances the establishment and dominance of the aciduric mutants streptococci15. The increased time sugar is in the mouth, there is inadequate time for remineralization by saliva, with the result that demineralization becomes the predominant mechanism.

**Oral Clearance of Carbohydrates**

In infants with ECC, the sleep-time consumption of sugar is another common characteristic. The salivary flow during sleep decreases oral clearance of the sugars, slowest on the labial surfaces of the maxillary incisors.
and the buccal surfaces of the mandibular molars which increase the length of contact time between plaque and substrates, thus increasing the cariogenicity of the substrate significantly.

**Time**

Time is important in ECC in relation to the frequency and amount of exposure of the offending liquid. Most reports of ECC stress the duration of the habit, be it bottle, breast, or sweetened pacifier, beyond the normal weaning period of about 14.2 months, 23.4 months with as high as 54 months.

**Feeding Practices**

Infant feeding practices associated with the development of caries in the primary dentition is poorly understood and somewhat controversial. To a major extent, the severity of caries attack lies with the feeding habit and not the type of feeding. Inappropriate nursing habits involving either the breast or the bottle, breast-feeding prolonged beyond the normal age of weaning, the regular use of baby bottles filled with a liquid with sugar added at bedtime and/or during the day, falling asleep with a pacifier covered with honey or jam may have been associated with ECC.

Breastfeeding versus ECC: reasons and counterarguments

Dr. Palmer suggests, it would be “evolutionary suicide for human milk to cause decay.”

According to Gardner et al, the lactose content of human milk alone can be cariogenic, if the milk is allowed to stagnate on the teeth for long periods especially at night. Most authors argue that ECC is associated with breastfeeding when the consumption pattern has certain characteristics such as ad libitum feeding, large number of breastfeedings a day, prolonged breastfeeding and, mainly, frequent breastfeedings during the night, resulting in accumulation of milk in the teeth, which, combined with reduced salivary flow and lack of oral hygiene, may produce tooth decay.

Alaluusua et al reported that breast feeding alone cannot be connected to increased or decreased caries prevalence. Furthermore, it has been stated that “population based studies do not support a definitive link between prolonged breast feeding and caries.”

The same was observed by Valaitis et al in a systematic review of 151 articles. Recent studies of Jose et al came to a conclusion that further investigations are needed to determine the prevalence of ECC in exclusively breast fed children.

Breast milk contains certain protective elements and continues to confer an advantage to prevent caries occurrence. Prolonged breast feeding beyond 2 years of age is associated with increased risk of developing ECC and could be due to the gradual depletion of the protective elements in human breast milk after prolonged lactation. Alternatively, mothers who breast feed beyond 12 months may be more likely to sleep with the child and breast feed ad lib (on demand) during night which is considered to be an important contributing factor for higher caries prevalence in these children.

There are also reports showing the relationship between early weaning from the breast and increased caries prevalence. It seems reasonable to support that earlier weaning increases the likelihood and duration of bottle feeding, which, in turn, is related to the caries prevalence.

The fact that either never being breast fed or breast fed for shorter duration or having been breast fed for longer periods were associated with caries development. This support the theory that it is not the breast feeding per se that causes dental caries but that the breast feeding habit may have an association both with the child’s dietary habits and with the rearing practice of the family which may serve as confounding variables. Various studies have investigated the association between breastfeeding and ECC. The most important limitation of these articles is that they do not employ the internationally adopted definitions of breastfeeding or ECC, while others use multiple definitions.

Therefore, the presumable cariogenicity of breast milk is an issue of paramount importance because, along with its substitutes, it is the major nutritional source in the first years of life.

**Bottle feeding and ECC**

Prolonged or inappropriate bottle feeding has been shown to be implicated in ECC. Many investigations have found that prolonged and inappropriate bottle feeding affects caries rate. Inappropriate feeding behaviors, such as nocturnal and prolonged use of a baby bottle during the day, the manner in which the child fell asleep were found to be associated with rampant caries.

They also reported that the severity of ECC increases as the number of feedings per day increases. But the reports of Wendt et al showed that neither the frequency of use of a feeding bottle nor the habit of nocturnal bottle feeding containing formula seems to influence the ECC prevalence.

**Pacifiers/ Comforters and ECC**

The positive correlation between caries and the use of sweetened comforter bottles and other types of sweetened comforters and pacifiers was found to be apparent by many studies. Although the habit of dipping the pacifier in sugar is associated with early colonization by Streptococcus mutans in predentate infants, a systematic review did not find any consistent correlation between the use of pacifiers and the development of ECC, regardless of the length of use of pacifiers and of the introduction of sweeteners or not.
Conclusion

Thus, there is no scientific evidence that confirms breastmilk is associated with ECC development. This relationship is complex and contains several other confounding variables, which turns a usually noncariogenic and potentially caries protective infant food staple into a cariogenic source. These variables need to be identified. Inappropriate feeding behaviors, such as nocturnal and prolonged use of a baby bottle were found to be associated with ECC. Hence, the association between bottle feeding and ECC needs to be strengthened by further research. Also, the practice of adding sugar or sweetening agents to milk has been clearly recognized as a risk factor for caries.

Thus, it is evident throughout the literature that ECC is a complex entity in which feeding practices alone does not play a prime role but is influenced by a myriad of cultural, behavioral, biological and microbiological factors. So, the area has conflicting reports which needs further research.

References

Gamma Glutamyl Transpeptidase and total cholesterol levels in smokers and congestive cardiac failure patients

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Abstract

Background and Objectives

The level of gamma glutamye transeptidase (GGTP or GGT) is specifically increased in alcoholic liver diseases with biliary obstruction. There are many other diseases where alterations in the level of GGTP are noted. Hence, the study was aimed to estimate the level of GGTP in smokers and congestive cardiac failure (CCF) patients along with total cholesterol (TC).

Material and Methods

35 each clinically diagnosed CCF patients, smokers and normal healthy male subjects were included in the study. GGTP was estimated by enzymatic colorimetric method and total cholesterol by cholesterol oxidase peroxidase method using P-800 chemistry autoanalyser. Statistical analysis was done using ANOVA.

Results

The GGTP and TC levels were significantly (p<0.001) increased in both CCF patients and smokers but increase is more significant in CCF patients.

Conclusion

GGTP can be used as a potential biomarker along with total cholesterol in cardiovascular diseases especially CCF because of their direct relationship with the indication of atherogenesis.

Key Words

GGTP, TC, CCF, Smokers.

Introduction

GGTP (or GGT) catalyses the transfer of a gamma – glutamyl group to an aminoacid or peptide to another substrate molecule or to water. It is an enzyme primarily present in kidney, Pancreas, Brain, Heart, Spleen, Liver and Prostate. It is stated that GGTP participate in the transport of aminoacids into cells which accounts for its membrane bound structure. This enzyme is universally distributed in tissue but with considerable variance in contest program of tissue. Renal tissue has highest level of GGTP, the enzyme present in Serum appears to Originate primarily from hepatobiliary system and GGTP activity is elevated in any and all form of liver disease. It is maximally increased in intra and post heaptic biliary obstruction; especially related to alcoholic liver disease. Apart from alcoholic liver diseases, GGTP elevation occurs in renal failure, fatty liver, primary or secondary (metastatic) neoplasms, especially pancreas (if associated with hepatobiliary obstruction), drug intake like barbiturates, Phenytoin etc. Reports on the levels of GGTP and TC in cardiovascular diseases and smoking are scanty. So, the present project was undertaken to estimate the levels of GGTP and TC in CCF patients and smokers.

Material and Methods

The test group included 35 male smokers who smoke about 8-10 cigarettes per day with filter (belongs to ITC company) since 10 years, 35 Clinically diagnosed CCF patients who were admitted in ICU were selected with 35 normal healthy control subjects; all subjects of test and control group belongs to 30-60 years of age.

Exclusion Criteria

Patients suffering from acute or chronic Pancreatitis, chronic infection, chronic granulomatous disease, acute of chronic renal failure, any liver disease, organic brain diseases, cardiomyopathies, disorders of spleen and any primary or secondary malignancies were excluded.

2ml of blood was collected from median cubital vein under strict aseptic precautions and with appropriate processing, subjected to P-800 chemistry autoanalyser for estimation of GGTP levels by enzymatic colorimetric method and total cholesterol by cholesterol oxidase peroxidase method.

Statistical Analysis

Done using ANOVA and comparison by Turkey HSD.

Results

• The levels of GGTP and TC increased significantly (p<0.001) in both CCF patients and smokers but
more significant in CCF patients.
• No significant Correlation is noted in any of the parameters between the groups.

Discussion
GGTP plays a very important role in amino acid transport and transfer of gamma-glutamyl group from peptides such as glutathione to other aminoaacids. In liver diseases GGTP correlates with alkaline phosphatases levels and it is the most sensitive indicator of alcoholic liver disease. It is noted that GGTP elevations may be associated with pancreatic, renal, cardiac and pulmonary disorders7.

In present study, the levels of both total cholesterol (TC) and GGTP are increased in both smokers and CCF patients but the increase is more noted in CCF patients. GGTP is involved in glutathione metabolism transferring glutamyl moiety to variety ofacceptor molecule including water and peptides having cysteine product to preserve intracellular homeostasis of oxidative stress8. In smoking, free radical damage the cellular antioxidant defense system causing oxidative stress. GGTP in vivo seems to recover cysteine from extracellular glutathione to preserve the cellular homeostasis of oxidative stress. Increase in intracellular oxidative stress may induce the enzyme GGTP. Glutathione being the important endogenous antioxidant (which is linked with GGTP in glutathione cycle), who’s homeostasis is disturbed; which might be the cause of increased levels of GGTP in smokers. Our study points the increase of GGTP in smokers which is supported by Yokoyama et al9, they report the development of hypertension, hyperlipidemia and diabetes mellitus were associated with increased GGTP levels.

There are evidences from recent studies that GGTP likely to be associated with cardiovascular diseases (CVD)10,11. In regard to coronary heart disease, it was observed that serum GGTP levels were associated with increased risk of myocardial infarction, cardiac death and failure12,13. GGTP is having direct involvement in atherosclerotic plaque formation and its level is increased in myocardial infarction and stroke patients14,15,16. The possible role of GGTP in atherogenic process is evidenced by increased level of oxidative stress17,18. GGTP was found to play a role in ethiopathogenesis because it was detected in atheromatous plaques of carotid and coronary arteries triggering oxidation of lipids; which is also precipitated by hypercholesteremia, resulting in complication of atherosclerosis as myocardial infarction, stroke, CCF etc19,20.

The link between serum GGTP levels and atherosclerotic plaque formation is based on oxidant action of glutathione catabolites in the extracellular space21. Our reports of increased cholesterol and GGTP levels in CCF patients are supported by Nakaniishi N et al22 and Lee DH et al23.

To conclude, the level of GGTP with lipid profile parameters can be considered as a potential biomarker for the indication of assessment of risk of complication of atherosclerosis like myocardial infarction, stroke and CCF patients.

Table 1: Comparison of GGTP and TC levels among smokers and CCF patients compared to controls.

<table>
<thead>
<tr>
<th>Sl. No</th>
<th>Parameter</th>
<th>Normal (Mean± SD)</th>
<th>Smoking (Mean± SD)</th>
<th>CCF Patients (Mean± SD)</th>
<th>p-value</th>
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<td>No of Sample</td>
<td>-</td>
<td>35</td>
<td>35</td>
<td>35</td>
<td>-</td>
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<tr>
<td>1</td>
<td>Total Cholesterol(mg/dL)</td>
<td>96 +20.147</td>
<td>218.50+11.4240</td>
<td>223.21+14.992</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>2</td>
<td>GGTP(U/L)</td>
<td>23 +8.990</td>
<td>74.474+8.5849</td>
<td>171.778+46.4207</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

References


Clinical audit in dentistry – The third eye of oral care

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Abstract
Producing good quality goods should be the primary responsibility of the manufacturer, and ensuring good quality services is the responsibility of the government. Utilizing good quality services is the basic human right of each and every individual. Provision of efficient and effective quality health care to the public is one of the ethical principles. Assessment of quality of health care provided to the public can be done through clinical audit system. Clinical audit is a quality improvement process that seeks to improve patient care and outcomes through systematic review of care against explicit criteria, and the implementation of change. Clinical audit results in improved clinical practice, increased efficiency, better clinical outcomes or more cost-effective service. In India, there is hardly any audit system in dentistry though the concept has existed since a long time. This review is an insight on methodology of clinical audit and the need for implementation of such a system for dental practice in Indian scenario.

Key Words
Clinical audit, dentistry, quality assurance, dental records.

Introduction
Dentistry is both an art and science, demanding high standard of skill and competency of the operator, along with quality armamentarium for providing the best possible oral care. This distinguishes the profession of dentistry uniquely from other health professions. Professional ethics mandate that the dentist stay current with new techniques, therapeutics, and materials, and eliminate outmoded ones in the pursuit of providing the best care for his/her patient. Providing just health care is not at all important, rather providing quality care is all that matters. Quality assessment of oral care can be done through systematic and organized clinical auditing system. Clinical audit results in improved clinical practice, increased efficiency, better clinical outcomes or more cost-effective service, all of which are a part of clinical governance.

Despite India having 290 dental colleges and approximately 80,000 dentists, catering to the needs of approximately 1.15 billion people, there is hardly any clinical audit system for quality assessment of oral care. Adequate infrastructure and resources in the form of manpower, money and materials are essential for conducting dental audit. This highlights the need of a third party system - government and/or non-governmental organization(s) and/or professional bodies to take proactive measures to carry out quality assessment in all the dental clinics and dental institutions. Sound methodology and supportive environment with minimal investment of resources is the need of the hour to carry out dental audit.

Defining Clinical Audit
Clinical audit is a quality improvement process that seeks to improve patient care and outcomes through systematic review of care against explicit criteria and the implementation of change. Aspects of the structure, processes, and outcomes of care are selected and systematically evaluated against explicit criteria. Where indicated, changes are implemented at an individual, team, or service level and further monitoring is used to confirm improvement in healthcare delivery. This definition is endorsed by the National Institute for Clinical Excellence.

Types of Audit
Formal and informal audit
Formal audit is often published. Informal audit may involve regular meetings to present and discuss selected cases.

Internal and external audit
Audit may be internal where the practitioner reviews his/her own work or external where the review of work is by an outside body separated by distance, experience and values.

Broad based and focused audit:
Audit can be broad or narrow in focus. Under the category of broad based audits, two types of audit exist – procedural and analytical audit.

Procedural audit is an assessment of adequacy of record-keeping. It answers the following questions:

a. Is the documentation of the patient care process adequate?
b. Does it meet the standards of practice?
c. Are there serious procedural omissions or errors that affect the quality of care?

**Analytical audit** is employed to determine the level of quality of care, by answering these questions:

- a. Was the diagnostic information (data base) adequate to identify all significant clinical problems?
- b. Was the diagnoses correct based on the information available?
- c. Was the treatment plan well-founded based on the diagnostic information and statement of problems?
- d. Were there avoidable errors, omissions or inadequacies in the execution of treatment, resulting in complications, compromised results, or failure?
- e. Was the overall care of the patient appropriate and adequate?

**Focused audits** are topic-specific, focus more narrowly on some aspect of patient care. For instance, any diagnostic test or procedure or any therapeutic modality could be the subject of this type of audit. It focuses on the following questions:

- a. Were the indications and contraindications for a particular therapy properly considered?
- b. Was the treatment technically adequate?
- c. Were there complications? Were they avoidable?
- d. Were the short term and long term results satisfactory based on the preoperative condition?

**Stages in Clinical Audit**

The methodology of clinical audit and the environment in which it operates are inter-related. If the environment is supportive but clinical audit methods are not used appropriately, there may be less improvement than expected, or no evidence that improvements have been made. Similarly, if clinical audit methods are used well but in an environment that is not supportive, the result may also be a failure to improve care and frustration among those involved.

**Stages in Clinical Audit**

**Stage one: Preparing for audit:**
Good preparation is crucial to the success of an audit project. The following issues are considered in preparation stage:

- a. Is the topic concerned of high cost, volume or risk to staff or users?
- b. Is there evidence of a serious quality problem, for example, patient complaints or high complication rates?
- c. Is good evidence available to inform standards, systematic reviews or national clinical guidelines?
- d. Is the problem concerned amenable to change?
- e. Is the topic pertinent to national policy initiatives?

**Defining the purpose**
A project without clear objectives cannot achieve anything. A sense of purpose must be established before appropriate methods for audit can be considered. The following series of verbs may be useful in determining the aims of an audit:

- a. To improve
- b. To enhance
- c. To ensure
- d. To change

**Providing a structure**
To enhance the benefits of audit, an organization needs a structured audit programme and a team of well
Audit in general and the objectives of the project

Everyone who becomes involved in an audit project needs to involve the right people with the right skills from the outset. To be successful, a clinical audit project needs an understanding of what is expected of the team and objectives of the project. The focus of any audit project must be those receiving care. The most common method of involving users in clinical audit is the satisfaction survey.

Identifying and developing skills for audit projects

Lack of training and audit skills are highlighted in the review as barriers to successful audit. It is important that the team includes members from all the relevant groups involved in care delivery, and not just those with clinical experience. All the team members should have a basic understanding of clinical audit, an understanding of and commitment to the plans and objectives of the project, and an understanding of what is expected of the project team. The focus of any audit project must be those receiving care. The most common method of involving users in clinical audit is the satisfaction survey.

Stage two: Selecting criteria

Within clinical audit, criteria are used to assess the quality of care provided by an individual, a team or an organization. These criteria are explicit statements that define what is being measured and represent elements of care that can be measured objectively. Criteria can be classified into those concerned with:

- Structure (What you need)
- Process (What you do)
- Outcome of care (What you expect)

Structure criteria: Refer to resources required. It includes the number of staffs, organizational arrangements, the provision of equipment and physical space.

Process criteria: Refer to actions and decisions taken by practitioners together with users. These actions may include communication, assessment, education, investigations, prescribing, surgical and other therapeutic interventions, evaluation and documentation.

Outcome criteria: These are typically measures of the physical or behavioral response to an intervention, reported health status, and level of knowledge and satisfaction. Sometimes surrogate, proxy or intermediate outcome criteria are used instead.

For criteria to be valid and lead to improvements in care, they need to be based on evidence, related to important aspects of care, and above all, measurable. Developing such criteria can be time-consuming and requires considerable expertise. Therefore, in some situations, implicit criteria have been used. This means that the review of care is undertaken by senior clinicians who rely on their own experience in judging care. Because of the difficulties of ensuring reliability in the interpretation of information about the care that was given, this method should be avoided where possible. Benchmark criteria should be set to avoid setting unnecessarily low or unrealistically high target levels of performance.

Stage three: Measuring level of performance

Data collection is the key step to measure the level of performance. It is always tempting to collect more data than necessary, but only the minimum amount required by the objectives of the audit should be collected.

Dental records

A well designed and properly maintained patient record is an important quality assurance tool because it documents all aspects of patient care. It has high utility because it is an existing part of the practice, and can be used in quality assurance activities without disrupting normal office procedures.

The dental record should adequately document each of its five phases – assessment of patient’s condition, diagnoses arrived at from this assessment, the therapeutic plan which follows from the diagnoses or statement of problems, the provision of treatment, and outcomes resulting from the previous four phases of the processes. Patient care is not a linear process with a beginning and an end. It is actually a dynamic cyclic process that repeats itself for each sequence of treatment. At each recall visit, the dentist evaluates not only the outcome of the previous course of treatment, but also any new or altered conditions of pathology and the results of patient’s self-care.

Quality dental record keeping is recognized by the profession as a requirement of prudent and effective practice management. Universal standards for dental record management need to be established which should include documentation of consultations by and referrals to specialists and medical professionals, authorization of treatment plans and plan modifications by faculty and patients, status and timeliness of care, radiographic services, communications with patients regarding their care, treatment provided, faculty supervision and care provider identification. These standards form the basis of dental record audit. Unfortunately, charts in most dental practices do not meet these requirements, and therefore, are inadequate instruments for quality assessment.

Sampling users

It may not always be practical or feasible to include each and every user. In this case, a representative sample is usually chosen from which inferences about the total population can be made. Various statistical methods can be used for calculating sample sizes, depending upon the type of data. Random sampling should be used whenever possible to minimize the risk of bias. Sometimes, two-stage sampling is done in order to improve efficiency. A small sample is selected first, and if unequivocal
conclusions can be drawn, no more data are collected. If the results are ambiguous, a larger sample is selected.

Confidentiality of data

According to the requirements of Data Protection Act, 1998, these guidelines must be followed to maintain the confidentiality of the data.

a. When information is gathered directly from patients, auditors must explain why the information is needed and what will happen to it, before asking for the patient’s consent.
b. If information is obtained from medical records, either patients must consent to identifiable data being used, or a member of the healthcare team should make the information anonymous before it is used in audit.
c. If neither of these approaches is possible, it may be permissible for someone from outside the team, who is suitably trained and subject to a duty of confidentiality, to collect the data from the records and make it anonymous without seeking patient consent.

Data Analysis

The analysis can range from a simple calculation of percentages to relatively sophisticated statistical techniques. If the results are to simulate change, the analysis must be simple enough for everyone in the care process to understand. If samples have been taken, the most appropriate calculation to perform is confidence intervals.

Stage four: Making improvements

It would be a futile exercise of conducting dental audit without drawing conclusions from the work that has been done. Once conclusions are made, it is easy to make appropriate recommendations. An action strategy has to be designed which should make certain that the required action is planned to take account of what needs to be done, by whom and by when. The implemented change can be subjected further to re-auditing to ensure that the desired change has taken place.

Stage five: Sustaining improvement

While improving performance is the primary goal of audit, sustaining that improvement is also essential. Systematic and continuous monitoring should be carried out to detect declining performance. Errors and adverse incidents can also be used for continued monitoring. Comments from users may be included as sources of information about performance.

Clinical Auditing – Ongoing Process

Clinical auditing should neither be once for all nor short-term process. If this is the case, the results of audit might probably be biased because of Hawthorne effect on the providers of the care. Thus, auditing should be an ongoing process to sustain the quality of health care.

Barriers to Successful Clinical Audit

Success of clinical audit depends on how well the barriers are tackled. Some of the barriers to audit include:

1. Lack of resources
2. Lack of expertise in project design and analysis
3. Lack of overall plan for audit
4. Poor relationships between professional groups or agencies and within teams
5. Organizational problems, such as lack of supportive relationship between clinicians and managers

Feasibility of Implementing Dental Audit in India

The concept of clinical audit for oral health care is a vision for a country like India. Such a vision would turn into reality in the near future, if the following issues are addressed.

1. Do we have adequate resources to implement audit system?
2. Who will comprise the audit team?
3. Who will be the central authority for conducting dental audit – government and/or non-governmental organization(s) and/or professional bodies like Dental Council of India (DCI), Indian Dental Association (IDA)?
4. Will the dental practitioners and concerned authorities of dental institutions extend their cooperation to carry out a systematic audit?
5. What should be the time interval required for conducting audit and how long should the process continue?
6. What is the benchmark level of dental care that needs to be set for making comparisons?

Conclusions

Quality oral health care increases the demand for seeking health care among public which in turn calls for continuous upgradability in the existing quality of health care system. This means that provision of quality health care gives immense satisfaction to the consumers; thereby convert their felt needs in to demands and build a high level of hope and anticipation to seek a much better quality of health care. Lobbying of policy-makers should be done to implement mandatory auditing by a third party for all the dental practitioners and dental institutions. If implemented, it would certainly serve as the third eye of oral care.

References


Aesthetic management of malpositioned implant following extraction of endodontically failed tooth: A case report

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Abstract
Implant stability is attributed to its anchorage in the surrounding alveolar bone, referred to as osseo integration1,2. Proper implant placement in bone is necessary for the success of osseo integration and function3. Implant-supported single crowns have become a valid alternative to conventional fixed dental prosthesis due to their excellent clinical long-term results. Adequate bone base is a prerequisite for functionally and aesthetically optimum reconstruction of soft tissue architecture around a dental implant4. When implants are malpositioned axially, prefabricated or custom made angled abutments may be used for acceptable comfort, function and aesthetics. When implants are apico-coronally malpositioned gingival ceramics can aid in restoration of aesthetics and imitation of emergence profile of the restored tooth. However, along with good survival rates, aesthetic factors are important for success in anterior regions. Gingival harmony is an important element in the aesthetics of the smile. Clinicians need to have the essential knowledge to create an optimal soft-tissue profile around teeth and implant restorations. The following case report describes the treatment of a malpositioned osseo-integrated dental implant with an intruded axially placed implant fixture.

Key Words
Malpositioned implant, Implant aesthetics, Gingival ceramics, Endodontic failures.

Introduction
Titanium dental implants have been used in the treatment of partial or complete edentulism. The height and width of the residual alveolus and surrounding anatomical structures can determine the proper position and path of insertion of dental implants.

An ideal positioning of implant in all 3 dimensions is required for optimal aesthetic results to be secured5,6. These dimensions, i.e. mesio-distal, oro-facial and apico-coronal are described defining “comfort” and “danger” zones for proper positioning in the anterior maxilla7. Aesthetics includes both the white and red aesthetics of which much importance is given to the red aesthetics8,9. The standard parameters for achieving aesthetic implant prosthesis result in one that is in harmony with the perioral facial structures of patient. The aesthetic peri-implant tissues, in their health height volume, colour and contours must be in harmony with the healthy surrounding tissue. The restoration should imitate the natural appearance of the missing dental unit in colour, form, texture, size and optical properties10,11.

Case Report
A 45 year old female patient reported with a mobile segment of maxillary left lateral incisor. Radiovisiograph revealed endodotically restored fractured tooth in an irreparable condition [figure 1]. The tooth was extracted and the granulation tissue cleared. Post extraction examination showed a major labial bone deficiency [figure 2]. A GS 4x11.5mm implant fixture (Osstem) was placed to a desired and determined position considering the position of adjacent teeth and availability of bone

Figure 1

Figure 2
Two of the exposed threads of fixture were covered with bone graft material and GTR membrane. The labial and palatal flaps were properly approximated and sutured. The implant fixture could not be ideally positioned due to the poor quality of bone. It was observed that the implant fixture was significantly malpositioned axially and apically [figure 4].

The implant was restored at 6 months after a second stage surgery. Osstem healing abutment /gingival former was placed to reveal a healthy gingival tissue. An implant level impression was made using a GS Osstem transfer transfer coping and a custom tray with addition polyvinyl siloxane material. Implant analog was positioned in the impression and gingival replica was completed in silicone material. A definitive cast was poured in a type IV die stone.

A 150 angled abutment with a collar height of 2mm was used to correct the labial malpositioning of implant. A PFM crown with the cervical gingival shade ceramic was fabricated for the correction of gingival recession and to mask an elongated crown length.

The abutment was fixed to the implant fixture and crown was tried for the marginal integrity and aesthetics and sent for final finish.

The screw access opening was blocked using low strength temporary cement prior to the final cementation. The patient has been followed regularly for routine hygiene and implants condition and has been found to have continued to report excellent comfort, aesthetics and function.

Discussion
Among various techniques reported to improve soft tissue deficiency are, the use of gingival-coloured acrylic resin facade, flexible silicone-based tissue-coloured material, or removable prostheses. The loss of peri-implant tissue can also be corrected by applying gingival-coloured ceramic on the cervical portion of implant-supported metal ceramic restorations, achieving harmonious mucogingival contours, as described in this clinical report. Another advantage of the described technique is the repositioning of the cement interface between the abutment and the restoration away from the tissues. When the gingival-coloured ceramic is applied to the crown, the cervical extension of porcelain toward gingival embrasure spaces may be limited by the path of insertion of the crown and adjacent teeth. In contrast, the application of gingival-coloured ceramic on a customized abutment allows a gingival embrasure space to be filled where interproximal papillae are missing and allows for the creation of a natural appearance, as well as fewer limitations from the contours of the adjacent teeth. In the situation presented, the aesthetics was also improved by creating the illusion of a tooth emerging from the soft tissue.

Conclusion
Horizontal and vertical bony deficiencies may result in malpositioning of dental implants. The maximum disangulation in relation to the long axis of the implant is recommended not to exceed 300. Severe malpositions of
implant in aesthetic zone may be solved by prosthodontist to some extent. Gingiva-coloured dental porcelain added on the abutment to replace the missing gingival tissue, further improves the aesthetic outcome of the definitive restoration.

Reference
Biomedical waste management practices: A cross-sectional study in an urban setting
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Abstract

Objective
To study the biomedical waste management practices in different health facilities of Noida U.P.

Material and Methods
It is a cross-sectional study including all medical practitioners of urban NOIDA. A modified Biomedical waste auditing questionnaire of Central Pollution Control Board was used to collect data. Data was tabulated on Microsoft Excel sheet and analyzed using Epi-Info version-6 software.

Results
Unqualified medical practitioners which were 39.4% of the study group were not using protective gear; not segregating/labeling/disinfecting the biomedical waste. Among qualified medical practitioners, 32.5% were using mechanical devices, 65% were segregating and labeling biomedical wastes while only 6.9% were disinfecting the sharps. Conclusion: Biomedical waste management is grossly inadequately practiced and there is a need to implement present rules and regulations with renewed vigour.

Key Words
Biomedical Waste, Bio-waste Management.

Introduction
Biomedical waste is the waste generated in the diagnosis, treatment or immunization of human beings or animals, in research or in the production of testing of biological products including all categories of infected and toxic waste that is a potential threat to human beings and the environment1.

It is a major issue of concern not only for Hospitals and Nursing Homes but also equally for the Environmental and Law enforcing agencies, general public and media.

All individuals exposed to hazardous health-care waste are potentially at risk, including those within health-care establishments that generate hazardous waste, and those outside these sources who either handle such waste or are exposed to it as a consequence of careless management. The main groups at risk are Medical doctors, nurses, healthcare auxiliaries, and hospital maintenance personnel; Patients in health-care establishments or receiving home care; Visitors to health-care establishments; Workers in support services allied to health-care establishments, such as laundries, waste handling, and transportation and Workers in waste disposal facilities (such as landfills or incinerators), including scavengers2.

Material and Methods
Present study was conducted in NOIDA city, UP. It is located in the state of Uttar Pradesh at the fringes of Delhi, the national capital. The study was carried out over a period of one year from August 2007 to October 2008. It is a cross-sectional study, institution based study. The universe consist of all the government hospitals, dispensaries and research institutes; all hospitals, nursing homes, clinics run by private sector and charitable; all dental hospitals/clinics; all laboratories and radiological diagnostic centre; all veterinary hospitals and all medical practitioners without any formal qualification (quacks) are also included. A purposive/feasible, non-probable sampling technique was taken for the study. The entire universe was taken for this study. Superintendent/in-charge of the health care facility generating biomedical waste was interviewed to collect data. A modified Biomedical waste auditing questionnaire of Central Pollution Control Board was used to collect data. The questionnaire was pre-tested and administered to collect information about knowledge of biomedical waste disposal, including section for observation, collection, segregation, and disposal of biomedical waste. Every health care facility of Noida, generating biomedical waste was included in the study. Health care facilities which do not cooperate or do not respond to Questionnaire or found closed at the time of collection of data were excluded from the study. Data was tabulated on Microsoft Excel sheet and analyzed using Epi-Info version-6 software.

Observations
Table no. 1 shows distribution of various types of health care facilities generating bio medical waste studied. 56
centers providing Medical care, 13 Diagnostic centers and 1 Research Facility and 1 Veterinary Facility participated in the study. Table no. 1 also shows distribution of qualification among doctors working in the health care facility. Doctor with highest qualification was taken into account. The table no-1 shows that a total of 43 qualified medical practitioners and 28 practitioners without any recognized medical qualification (by MCI) participated in the study. It was observed that the difference in the distribution of use of protective gears worn by health care workers while handling biomedical waste in the health care facilities of qualified medical practitioners and unqualified medical practitioners. It was observed that 65.5% of qualified medical practitioners and none of the unqualified medical practitioners were segregating waste into the designated colored container. Table no. 2 shows the difference in the distribution of labeling of biomedical waste, as practiced, among qualified and unqualified medical practitioners. None of the unqualified medical practitioners were labeling the waste; whereas 34.8% of qualified medical practitioners were labeling the waste. It was observed that the utilization of offsite treatment facility for treatment of biomedical waste by health care facilities of qualified and unqualified medical practitioners under study. 65.2% of qualified medical practitioners and none of the unqualified medical practitioners were utilizing offsite treatment facility for disposal of biomedical waste. It was observed that the classification of waste sharps in the health care facilities of qualified medical practitioners and unqualified medical practitioners. 65.1% of qualified medical practitioners had classification of waste sharps where as none of the unqualified medical practitioners had classification of waste sharps. It was observed that distribution of methods of disinfection of waste classified as waste sharps by health care facilities of qualified medical practitioners and unqualified medical practitioners. 6.9% of health care facilities of qualified medical practitioners were disinfecting sharps before disposal where as none of health care facilities of unqualified medical practitioners were doing so.

**Discussion**

A total 71 health care facilities generating bio medical waste participated in the study. 56 centers Medical care, 13 Diagnostic centers, 1 Research Facility and 1 Veterinary Facility formed the study group. A total of 43 qualified medical practitioners 28 practioners without any recognized medical qualification (by MCI) responded to the questionnaire. Of the 71 health care facilities under study, only 30 health care facilities were registered with State Pollution Board for biomedical waste management and 41 health care facilities were not registered. 69.7% among qualified medical practitioners and none among unqualified medical practitioners.

In the study group, health care workers of only 1.4% health care facilities was using puncture-proof gloves, 4.2% were using apron, 0% were using goggles, 2.8% were using facemask and 7.8% were using shoes, while handling biomedical waste. These findings where very less as compared to the observations of Rao et al, most of the Safai karmacharis were using complete protective equipment like gloves, masks, shoes etc while handling waste.

39.4% of the health care facilities were segregating waste into the prescribed colored container. 65.5% of qualified medical and 0% of unqualified medical practitioners. Hemchandra in his paper recommends Segregation is the essence of waste management and should be done at the source of generation of bio-medical waste e.g. all patient care activity areas, diagnostic services areas, operation theatres, labour rooms, treatment rooms etc. The responsibility of segregation should be with the generator of biomedical waste i.e. doctors, nurses, technicians etc. (medical and paramedical personnel). The biomedical waste should be segregated as per categories mentioned in the rules.

It was observed in the study that 65.2% of qualified medical practitioners and 0% of unqualified medical practitioners were utilizing offsite treatment facility for disposal of biomedical waste. Central pollution Control Board recommends a Common Bio-medical Waste Treatment Facility (CBWTF), where bio-medical waste generated from a number of healthcare units, is imparted necessary treatment to reduce adverse effects that this waste may pose. The treated waste may finally be sent for disposal in a landfill or for recycling purposes. 39.4% of health care facilities were segregating waste

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**Table 1:** Distribution of types of health facility and qualification of health facility in-charge.

<table>
<thead>
<tr>
<th>S. No</th>
<th>Variable</th>
<th>Number</th>
<th>Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Type of health facility</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Research Facility</td>
<td>1</td>
<td>1.4%</td>
</tr>
<tr>
<td></td>
<td>Medical Facility</td>
<td>56</td>
<td>78.8%</td>
</tr>
<tr>
<td></td>
<td>Diagnostic Facility</td>
<td>13</td>
<td>18.3%</td>
</tr>
<tr>
<td></td>
<td>Veterinary Facility</td>
<td>1</td>
<td>1.4%</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>71</td>
<td>100</td>
</tr>
<tr>
<td>2.</td>
<td>Highest Qualification of Doctor</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>DM/MD/MCh</td>
<td>2</td>
<td>2.8%</td>
</tr>
<tr>
<td></td>
<td>MD/MS</td>
<td>21</td>
<td>29.5%</td>
</tr>
<tr>
<td></td>
<td>MBBS/PG Diploma</td>
<td>11</td>
<td>15.4%</td>
</tr>
<tr>
<td></td>
<td>BDS/BUMS/BHMS/BAMS</td>
<td>9</td>
<td>12.6%</td>
</tr>
<tr>
<td></td>
<td>No Medical Qualification</td>
<td>28</td>
<td>39.4%</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>71</td>
<td>100</td>
</tr>
<tr>
<td>3.</td>
<td>Qualification of Doctor</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Qualified Medical practitioner</td>
<td>43</td>
<td>60.5%</td>
</tr>
<tr>
<td></td>
<td>No Medical Qualification</td>
<td>28</td>
<td>39.4%</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>71</td>
<td>100</td>
</tr>
</tbody>
</table>
into waste sharps and 60.6% of health care facilities did not have classification for waste sharps. 65.1% of qualified medical practitioners had classification of waste sharps whereas 0% of unqualified medical practitioners had classification of waste sharps. Hollie Shaner et al in their Eleven Recommendations for Medical waste Management states that of the 10 percent or less portion of the waste stream that is potentially infectious or hazardous, the most immediate threat to human health (patients, workers, public) is the indiscriminate disposal of sharps (needles, syringes, lancets, and other invasive tools). Proper segregation of these materials in rigid, puncture proof containers, which are then monitored, for safe treatment and disposal is the highest priority for any health care institution. If proper sharps management were instituted in all health care facilities most of the risk of disease transmission from medical waste would be solved. This would include proper equipment and containers distributed everywhere that sharps are generated (needle cutters and needle boxes), a secure accounting and collection system for transporting the contaminated sharps for treatment and final disposal, and proper training of all hospital personnel on handling and management of sharps and personal protection.

Conclusions
1.4% health care facilities was using puncture-proof gloves, 4.2% were using apron, none were using goggles, 2.8% were using facemask and 7.8% were using shoes, while handling biomedical waste. None of the staff of unqualified medical practitioners was using any protective gear.

39.4% of the health care facilities were segregating waste into the prescribed colored container. 65.5% of qualified medical practitioners and none of the unqualified medical practitioners were segregating waste into the designated colored containers.

21.1% of all health care facilities were labeling the containers of biomedical waste generated, 39.4% were not labeling the containers and 39.4% were not aware of regulation regarding labeling of biomedical waste containers. None of the unqualified practitioners was labeling the waste; whereas as 34.8% of qualified medical practitioners were labeling the waste.

39.4% of all health care facilities were utilizing offsite treatment facility for treatment of biomedical. 65.2% of qualified medical practitioners and none of the unqualified medical practitioners were utilizing offsite treatment facility for disposal of biomedical waste.

39.4% of health care facilities were segregating waste into waste sharps and 60.6% of health care facilities did not have classification for waste sharps. None of the health care facilities of unqualified medical practitioners had classification of waste sharps 4.2% of Health Care Facilities were disinfecting waste classified as waste sharps before disposal and 95.8% were disposing it without disinfecting it. 6.9% of Health Care Facilities of qualified medical practitioners were disinfecting sharps before disposal whereas as none of the health care facilities of unqualified medical practitioners were doing so.

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A study of cytomorphological patterns and their diagnostic importance in tubercular lymphadenopathy

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Abstract

Tuberculosis [TB] is a specific communicable disease and is one of the most common disease in developing countries like India. There are 20-25 million cases of infectious tuberculosis in the world. In India alone there are 12-15 million cases of primary tuberculosis of which about 2 million are bacillary cases at any given period of time.[PARK & PARK 1986]. Currently, more people die of tuberculosis than from any other infectious disease. Death from tuberculosis comprises 25% of all avoidable deaths in developing countries. Every minute, a death occurs due to tuberculosis in our country. To add to the existing burden, the situation is compounded by the large scale increase of new TB cases associated with increasing HIV infection. India is estimated to have 3.5 million HIV patients, and about 1.8 million of these are co-infected with TB. Tuberculosis usually affects lungs and the next common entity is the tubercular lymphadenopathy [TLN].

Many a time exact diagnosis of TLN poses a great problem to the clinician. This is because of wide spread nature of the disease and consequently symptomatology produced is very vague. In TLN specialised investigation like ELISA test, PCR etc. are also not always diagnostic. In pulmonary cases diagnosis can be confirmed by demonstrating AFB in the sputum but the same is not true for TLN. Inability to demonstrate AFB in most of the patients of TLN has prompted many workers to develop serological tests. However these attempts have not so far resulted in providing a single reliable serological test for either pulmonary or extrapulmonary TB. Open surgical biopsy is a time consuming procedure and also this facility is not available at all centres in developing country like India. Thus in the present scenario, fine needle aspiration biopsy [FNAB] is recommended for evaluation of cytomorphological pattern [CMP] and its utility in establishing the diagnosis of tuberculosis [TB].

Key Words

Cytomorphological patterns, tuberculous lymphadenopathy/ lymph node, fine needle aspiratin biopsy, Tuberculosis.

Aim

A patient of TLN can not be treated unless and until histopathological or bacteriological proof for the disease is available. The first one is time consuming and also risky in debilitated patients of TB. The second one is usually not positive. In view of lack of any other sensitive diagnostic method and considering the importance of early diagnosis as the disease is easily curable in its earlier stages, present study was undertaken to overcome all these shortcomings and to ascertain whether there are any cellular constituents peculiar to TLN, which are useful in making the diagnosis in cases persistently negative on bacteriological examination.

Material and Method

The study was conducted on 52 FNAB material from randomly selected known patients of TLN in which treatment is not yet started. These patients were belonging to different age groups and involving lymphnodes at different sites. Out of these 38 were male and 24 were female patients. Aspiration material obtained in all these were adequate except in 3 patients where repeat aspiration was done. In the present study, technique described by Franzen et al [1968] was employed. At least 5 smears were prepared from each aspiration site. These smears were stained by—

1. Haematoylin and Eosin,
2. Papanicoloau and

After CMP evaluation these smears were divided broadly in to 2 groups.

GROUP A - Includes the smears which showed cytological features of Granulomas.

GROUP B - Includes the smears without any granulomas.

Granulomas were defined as aggregates of epitheloid cells. These are the cells with tapered ends and elongated folded nuclei, having poorly defined finely vacuolated amphophilic or cyananophilic cytoplasm. These cells are

<table>
<thead>
<tr>
<th>Table A: Shows Number of Cases in Each Group</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cytotlogic Findings</td>
</tr>
<tr>
<td>---------------------</td>
</tr>
<tr>
<td>With Granuloma</td>
</tr>
<tr>
<td>Without Granuloma</td>
</tr>
<tr>
<td>TOTAL</td>
</tr>
</tbody>
</table>
Observation

Discussion

The non-availability of a technically simple, reliable, reproducible and rapid test often makes the diagnosis of TLN difficult. The traditional method of mycobacterial culture, though considered as a gold standard, is less sensitive in pauci-bacillary situations like TLN. Moreover, it requires a sample from the site of lesion, which is often difficult. Methods like radiometric detection of bacterial growth, gas chromatography, DNA hybridization and PCR are technically too demanding to be widely available in developing nations. ELISA techniques, using various mycobacterial antigens, have been employed for the serodiagnosis of TB, but these are not always reliable in TLN because of the spectrum of antibodies in serum varies with genetic background of the subject as well as the stage of the infection. In endemic areas antibodies would normally be present in the population, and these antibodies may have been induced in response to tuberculosis/environmental mycobacteria which are frequently cross-reactive.

Present study was carried out at "SARSWATHI INSTITUTE OF MEDICAL SCIENCES" Anwarpur, Hapur. Total 52 diagnosed cases of TLN were selected. Z-N staining was used to reach the definitive diagnosis. After CMP evaluation, the cases were divided into two groups [Theodore et al.]

Group A ---- with granuloma
Group B ---- without any granuloma.

C caseation in a smear showing evidence of granulomatous inflammation is suggestive of TB [Koss, 1979]. Patra et al. [1983] for the first time found 87.1% diagnostic accuracy in TLN by FNAB.

In our study TABLE "A" Shows that granulomas were present in 32 [54%] cases where as there were no granuloma in 20 [46%] cases.

Smears from cases of Group A AND Group B were further analysed according to individual cell findings.

TABLE "B" shows epitheloid cells in 87% cases and Giant cells in 51% cases of Group A. In Group B caseation, fibrous strands and mixed inflammatory cells were the [100%] constant findings. Eosinophilic hyaline to granular material is suggestive of caseated debris.

TABLE "C" shows that Z-N staining was positive in about 4% of the cases and thus does not contribute much in making the diagnosis.

Summary and Conclusion

The diagnostic problem in TLN results from the frequent failure of bacteriological methods of smear-microscopy in establishing the diagnosis. One of the aim of present study was to provide an early diagnosis and there by subjecting the patient to treatment in time for a disease that is practically curable. The results obtained are highly reliable with the sensitivity achieved is 91.3%. The advantages of simplicity, cost-effectiveness and rapidity make FNAB and CMP in TLN an attractive adjunct in diagnosis.

The findings of the present study can be summarized as follows—

1. Epitheloid cell and Giant cell is a characteristic feature of tuberculosis and was a persistent finding.
2. Caseous material and Histiocytes in a smear showing evidence of granulomatous inflammation is suggestive of tuberculosis.

Present study shows that CMP in TLN is very useful in making the diagnosis. The advantages of simplicity, cost-effectiveness and rapidity make CMP in TLN an attractive adjunct in diagnosis. It should be employed routinely to diagnose the TBN. Early diagnosis of tuberculosis and initiating optimal treatment would not only enable a cure of an individual patient but will also curb the transmission of infection and disease to others in the community.

References

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Knowledge of Swine flu among students in a university of western Turkey: A cross-sectional study

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Abstract

Objectives
The aim was to determine the level of knowledge about swine flu among a group of Turkish university students.

Methodology
This cross sectional study was conducted on 1146 students studying at some departments of Eskisehir Osmangazi University, western Turkey. The questionnaire prepared with reference to previous studies consisted of 15 questions and it was evaluated over a total of 19 points.

Results
The average score of knowledge level about swine flu obtained from the students was slightly above the middle level (11.75±2.19). Most respondents reported television/radio was the major sources of their information about swine flu (82.8%). The proportion of those who had knowledge about the fact that the most important and effective means for the prevention of swine flu is vaccination was only 32.0%.

Conclusion
This study concludes that accurate and complete information about the swine flu should be provided for students through the media and education when considered that the level of knowledge about the flu was not enough.

Key Words
Swine flu, students, information source, level of knowledge, vaccination

Introduction
Influenza is a highly contagious, acute illness which has afflicted humans and animals since ancient times1,2. In April 2009, a new influenza virus of swine origin emerged in Mexico and spread rapidly around the world.3 After the case from Mexico, the first two cases of human infection with a novel influenza A (H1N1) virus were reported in the United States (US). The virus was found to be an H1N1 virus that was antigenically and genetically unrelated to human seasonal influenza viruses and genetically related to viruses known to circulate in swine. In the ensuing weeks, the swine-origin influenza virus H1N1 spread worldwide, constituting a pandemic, as defined by the World Health Organization4,5.

The first case of swine flu in Turkey was identified on a tourist coming from the US to Istanbul in May 16, 2009. After this case, in the late May and early June, 3 new cases were confirmed in Istanbul, Izmir and Ankara. With laboratory examinations, since the beginning of pandemic influenza in Turkey, the number of those who died from flu has been 627. Of these people, 41 were pregnant or within the maternity period. As of January 25, 2010, 67 people have been still treating in the specialized hospitals such as state or university; 22 cases are in intensive care services, the lives of 17 cases are maintained depending on the breathing apparatus, being monitored. Thirty five percent of the persons who lost his life were healthy people who did not have an underlying chronic disease6.

In this study, the aim was determine the knowledge level about the swine flu among a group of university students.

Material and Methods
This cross sectional study was conducted on a group of students studying at some departments (Faculty of Arts and Sciences, Health Services Vocational High School, Faculty of Economics and Administrative Sciences, Faculty of Education) of a university in a province of western Turkey, Osmangazi University, Eskisehir, between Nov 1th and Dec., 31th 2009.

Subjects were randomly selected with a stratified sampling method. Osmangazi University is an urban, mid-sized university with six schools, all of which were included in the survey except for medical faculty. At least one class from the first, second, third and fourth years in each department were randomly selected to participate in the survey. In result, total study group consisted of 1146 students (13.8%).

The questionnaire, prepared with reference to previous studies in the literature7-10 included two parts. In the first part of the questionnaire, the students were asked to state their socio-demographic characteristics. The second part of the questionnaire evaluated information sources
and knowledge level about swine flu.

In the process of evaluation the questionnaires, 1 point for each correct answer and 0 point for each wrong answer was given. In the questionnaire for each correct answer to question 1 is, each wrong answer was given 0 points. One point for each of the questions 1., 3., 4., 5., 6., 8., 14., 15., and 6 options of the question 12 and 4 options of the question 13 was given, and the total points of knowledge were estimated over 19 points.

Permission for the study was obtained prior to collection of data, by contacting and receiving approval from the university directorship of the city involved. Participants were assured of the confidentiality of their responses and provided informed consent. All students gave their informed consent prior to their inclusion in the study.

Statistical analyses were made using the student's t test and ANOVA variance analysis. Results were given as numbers and percentages with a 95% confidence interval (CI). A value of p<0.05 was considered statistically significant.

Findings

Most of those constituting the study group were female students (n=740, 64.6%), with 406 male students constituting less (35.4%). The average age of the participants was 20.22±1.82 years (range=17-28 years). The most students were studying at the Faculty of Arts and Sciences (34.8%). The average level of knowledge about swine flu was slightly above the middle level (11.75±2.19; min=0, max=17).

In those studying at the Health Services Vocational High School, the average level of knowledge about swine flu were higher than those at the other schools (F=3.062; p=0.027). There was no significant relationship by gender and age groups in terms of the average levels of knowledge (p>0.05, each one). The distribution of the students’ average level of knowledge about swine flu by their gender, age and schools are presented in Table 1.

All the students reported that they had heard of the swine flu (100.0%). Most respondents indicated that mass media (television/radio) was the major sources of their information about swine flu (82.8%), followed by newspaper (13.4%) and internet (5.6%) (Unshown data).

Table 2 shows the distribution to the answers to some epidemiological characteristics, clinical symptoms, symptoms that require to apply to emergency department, risk groups, protection methods, vaccination and treatment related questions by the students: Over 90% of the students gave correct answers to the item ‘agent of swine flu is a kind of virus’ and nearly 100.0% (97.3%) to the item ‘swine flu can be transmitted by coughing, sneezing, shaking hands and kissing from person to person.’ The most frequently known clinical symptom by the students was fever (98.6%). Nearly all of the students (96.0%) gave correct answer to the item ‘fever lasting for more than three days’ as a clinical symptom that require to be admitted to emergency services. Although 80% students knew that those with any chronic disease were at risk for swine flu, most students (about 50% or less) did not know that who were at risk. The proportion of those who knew that the item ‘the most important and effective way is to be vaccinated’ was rather low (32.0%), where as the other proportions were very high (86.0% and over), especially for the item ‘washing hands frequently with soap is useful’ (98.2%). More than half of the students (52.1% and over) gave false answers to the items related to swine flu vaccination and treatment. Only 47.9% students knew that swine flu vaccine should not be made to those who have severe egg allergy and to those who developed severe allergy previously against the flu vaccine. In similar, their 66.9% did not know that protection of swine flu vaccination begins 2 weeks after it is applied, and their 64.6% were not aware of the fact that swine flu vaccine can be made to pregnant women.

Table 1. The distribution of the students’ average level of knowledge about swine flu by their gender, age and schools

<table>
<thead>
<tr>
<th>Variables</th>
<th>Number (percentage) n (%)</th>
<th>Average level of knowledge (X±SD) 20.09±3.80</th>
<th>Statistical analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Schools</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Faculty of science and literature</td>
<td>399 (34.8)</td>
<td>11.66±2.08</td>
<td></td>
</tr>
<tr>
<td>Health services vocational college</td>
<td>202 (17.6)</td>
<td>12.13±2.30</td>
<td>F=3.062; 0.027</td>
</tr>
<tr>
<td>Faculty of Economics and Administrative Sciences</td>
<td>268 (23.4)</td>
<td>11.56±2.39</td>
<td></td>
</tr>
<tr>
<td>Education faculty</td>
<td>277 (24.2)</td>
<td>11.81±2.02</td>
<td></td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Males</td>
<td>406 (35.4)</td>
<td>11.85±2.37</td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>740 (64.6)</td>
<td>11.70±2.08</td>
<td>t=1.103; 0.270</td>
</tr>
<tr>
<td>Age groups (years)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20 and below</td>
<td>677 (59.1)</td>
<td>11.73±2.18</td>
<td>t=0.520; 0.603</td>
</tr>
<tr>
<td>21 and over</td>
<td>469 (40.9)</td>
<td>11.80±2.21</td>
<td></td>
</tr>
</tbody>
</table>
Table 2: The distribution to the answers to some epidemiological characteristics, clinical symptoms, symptoms that require to apply to emergency department, risk groups, protection methods, vaccination and treatment related questions by the students (n=1146)

<table>
<thead>
<tr>
<th>Variables</th>
<th>True n(%)</th>
<th>False/ No idea n(%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Epidemiological characteristics</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Agent of swine flu is a kind of viruses</td>
<td>1108 (96.7)</td>
<td>38 (3.3)</td>
</tr>
<tr>
<td>Swine flu emerged firstly in Mexico</td>
<td>850 (74.2)</td>
<td>296 (25.8)</td>
</tr>
<tr>
<td>Swine influenza is very contagious but is not lethal.</td>
<td>552 (48.2)</td>
<td>594 (51.8)</td>
</tr>
<tr>
<td>Swine flu can be transmitted by pool, lake, sea and drinking water.</td>
<td>331 (28.9)</td>
<td>815 (71.1)</td>
</tr>
<tr>
<td>Swine flu can be transmitted by coughing, sneezing, shaking hands and smooching/ kissing from lips or cheeks</td>
<td>1115 (97.3)</td>
<td>31 (2.7)</td>
</tr>
<tr>
<td><strong>Clinical symptoms</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fever</td>
<td>1130 (98.6)</td>
<td>16 (1.4)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>1010 (88.1)</td>
<td>136 (11.9)</td>
</tr>
<tr>
<td>Widespread body pain</td>
<td>824 (71.9)</td>
<td>322 (28.1)</td>
</tr>
<tr>
<td>Cough</td>
<td>771 (67.6)</td>
<td>375 (32.4)</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>636 (55.5)</td>
<td>510 (44.5)</td>
</tr>
<tr>
<td>Sore throat</td>
<td>605 (52.8)</td>
<td>541 (47.2)</td>
</tr>
<tr>
<td><strong>Clinical symptoms that require to be admitted to emergency services</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fever lasting for more than three days</td>
<td>1100 (96.0)</td>
<td>46 (4.0)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>534 (46.6)</td>
<td>612 (53.4)</td>
</tr>
<tr>
<td>Difficult breathing</td>
<td>492 (42.9)</td>
<td>654 (57.1)</td>
</tr>
<tr>
<td>Chest pain</td>
<td>203 (17.7)</td>
<td>943 (82.3)</td>
</tr>
<tr>
<td>Headache</td>
<td>179 (15.6)</td>
<td>967 (84.4)</td>
</tr>
<tr>
<td><strong>Risk groups for swine flu</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Those with any chronic disease</td>
<td>917 (80.0)</td>
<td>229 (20.0)</td>
</tr>
<tr>
<td>Pregnants</td>
<td>591 (51.6)</td>
<td>555 (48.4)</td>
</tr>
<tr>
<td>Children under 2 years old</td>
<td>570 (49.7)</td>
<td>576 (50.3)</td>
</tr>
<tr>
<td>Hypertensive patients</td>
<td>276 (24.0)</td>
<td>870 (76.0)</td>
</tr>
<tr>
<td>Obese people</td>
<td>68 (5.9)</td>
<td>1078 (94.1)</td>
</tr>
<tr>
<td><strong>Protection methods</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>The most important and effective way is to be vaccinated.</td>
<td>367 (32.0)</td>
<td>779 (68.0)</td>
</tr>
<tr>
<td>Washing hands frequently with soap is useful.</td>
<td>1125 (98.2)</td>
<td>21 (1.8)</td>
</tr>
<tr>
<td>Patient people should wear masks</td>
<td>1104 (96.3)</td>
<td>42 (3.7)</td>
</tr>
<tr>
<td>When we cough and sneeze we should close our mouth.</td>
<td>1121 (97.8)</td>
<td>25 (2.2)</td>
</tr>
<tr>
<td>Rest of patient people at home is important</td>
<td>1024 (89.4)</td>
<td>122 (10.6)</td>
</tr>
<tr>
<td>Adequate and balanced diet is beneficial to protect people from influenza</td>
<td>985 (86.0)</td>
<td>161 (14.0)</td>
</tr>
<tr>
<td><strong>Vaccination and treatment of swine flu related items</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2 weeks after swine flu vaccine is made its protection begins</td>
<td>379 (33.1)</td>
<td>767 (66.9)</td>
</tr>
<tr>
<td>Swine flu vaccine can be made to pregnant women</td>
<td>408 (35.6)</td>
<td>838 (64.4)</td>
</tr>
<tr>
<td>Normal (seasonal) flu vaccine is protective against swine flu, too.</td>
<td>177 (15.4)</td>
<td>969 (84.6)</td>
</tr>
<tr>
<td>Swine flu vaccine should not be made to those who are severely allergic to eggs and to those who had been previously vaccinated against the flu and developed severely allergic.</td>
<td>549 (47.9)</td>
<td>597 (52.1)</td>
</tr>
</tbody>
</table>

**Conclusion**

When the schools that the students studied in terms of the average scores were compared, in those studying at the school of Health Services Vocational High School, the average levels of knowledge about swine flu were higher than those at the other schools (p<0.05). This is an expected result since the students in this school take education much concerning health related subjects.

In the current study, it was reported that for most respondents, ‘mass media (television/radio) was the major sources of the students’ information about swine flu (n=949, 82.8%). This result indicates radio and
television institutions have important tasks to raise the awareness of public about swine flu. Since swine flu is a viral disease, it has got an outbreak potential, having led to pandemia. Therefore, it is important to know the epidemiological features of this disease. In our survey, 91.6% of the students reported that 'the agent of swine flu is a kind of virus' and that 95.6% reported that this 'disease is transmitted by air'. These results indicate that the students have a quite good level of knowledge on this issue. Knowing that this disease is transmitted through the air will help to understand the reason for influenza pandemia.

In our survey, 96.7% of the students reported that 'the agent of swine flu is a kind of viruses' and 97.3% reported that 'this disease is transmitted by air (cough, sneezing, handshake, kissing from person to person)'. These results indicate that the students have a quite good level of knowledge on this issue. Knowing that this disease is transmitted through the air will help to understand the reason for influenza pandemia.

Knowing the symptoms of the disease are an important situation in terms of early applies of patients to health institutions and making necessary treatment before the clinical symptoms of the disease becomes more severe. The study subjects gave correct answers between 52.8% and 98.6% to clinical symptoms related symptoms. Especially, that about 100 (98.6%) of the students knew that fever is one of the clinic symptoms of swine flu suggests that the students have enough information of swine flu.

The best known risk group by the students participating in our study was those who had chronic diseases (80.0%), and the proportions were lower for the other risk groups. Although these rates are not enough, it is acceptable. More than half the students reported that they did not know that pregnant women and children younger than 2 years of age were in the risk group. This may show that the university students do not deal with these people in terms of social life.

Become aware of the symptoms that require contacting the emergency services will contribute to for patients to apply earlier to doctor, and thus to decrease in morbidity and fatality of the disease. In this study, the study subjects most frequently reported that one of the symptoms that require to applying to emergency services was fever lasting more than 3 days (96.0%). However, the subjects did not know enough the other symptoms that require to apply to emergency services (50% and below). This points out that the subjects' level of knowledge about swine flu was not enough.

In this study, just 18.7% of the subjects reported that the most effective method in protection the disease was vaccination; whereas, most students between 84.6% and 95.6% were aware of the other methods for protection. These results, as in many countries, may be a result of the sensational news in Turkey on pandemic influenza vaccine. Thus, we may say that all training activities against the swine flu have failed. Incomplete, inaccurate and contradictory information given in the Turkish media about the vaccine caused the students to have lower level of information about the vaccination. For example, just 47.9% of the subjects could give correct answers to the item 'swine flu vaccine should not be made to those who are severely allergic to eggs' when compared to the other items.

As a result of incomplete, inaccurate and contradictory information on the swine flu vaccine, the majority of the students (91.8%) expressed that they did not want to be vaccinated and that they had no idea about the vaccination, which causes all the studies conducted on getting under the control the swine flu to be qualified as unsuccessful.

More than 20% of the students (22.9%) did not know that "there is a treatment for the treatment of swine influenza". This indicates that the students have the insufficient information about how to contact health care facilities in case of illness. Considering that there is no treatment for the disease would reduce the frequency of contacting to health care facilities when clinical symptoms occur.

This survey has revealed that public health professionals should develop communication messages closely related to the epidemic situation to target the information needs from the public. When taking into consideration the students' level of knowledge in general, swine flu is still a threat to Turkish students. Thus, comprehensive and effective measures for surveillance and prevention of the disease are needed to control its spread.

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The author wishes to thank the schools' headmasters, teachers, and the research students for helping to collecting the data and also those participating in the study for their valuable efforts and time.

Interest of Conflict

The authors declare that he has no competing interests, and also that there is no funding organizations for this study.

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