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International Journal of Medical Students

The International Journal of Medical Students (IJMS) is a peer-reviewed open-access journal, created to share the scientific production and experiences of medical students worldwide.

Caducus

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All full-text articles are available at:

www.ijms.info

e-ISSN 2076-6327 (Online)
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Training Future Scientists – Enhancing Research Capabilities in Developing Nations

Emmanuel Coronel,* Lorenzo Anez-Bustillos,* Felipe Fregni

"Give a man a fish and you will feed him for a day. Teach a man how to fish and you will feed him for a lifetime"

Chinese proverb

Not surprisingly, the main bulk of scientific contributions currently present in the literature come from research performed in developed nations. The lack of government funding, unclear health research priorities, and problems retaining scientific researchers, are among the reasons why this disparity has historically held true. These arguments lead underdeveloped countries to blindly accept the erroneous fact that patient care and scientific research do not belong in the same equation. Medical research has not been a health reform priority in most of these nations, and thus has been traditionally underfunded. The allocation of government funds to this field and the priority it is given reflects its unequal development among many regions of the world. Consequently, resources destined to train scientists and improve the research infrastructure and equipment are nowhere near the top of the priority list.

In the past few years, regional and global changes have been developed with the purpose of enhancing health research opportunities in non-traditional countries. There seems to be a growing consensus that many answers to common health problems can be provided by performing research. In light of this, the idea of creating knowledge-based economies in many emerging nations (i.e. Brazil, India, Russia, and China) has made research development one of its critical components. Similarly, the globalization phenomenon has motivated these nations to understand the role of research as a potential driving force for their economic growth, as has happened with many other industries in the past half century.¹²

Having experienced these challenges first hand, we have previously proposed how they can be translated into opportunities to increase the scientific contributions of other nations.³ For many years, our commitment to promote this change in paradigm has focused on education through the development of initiatives aimed to promote medical research in developing countries. A few years after introducing a novel program on global clinical research training using innovative web tools,⁴ we decided to enhance our offering and go beyond the books and lectures by creating the Principles and Practice of Clinical Research Fellow Practice (formerly known as the Latin American initiative). Through this program, we coupled our theoretical training with hands-on research experience by giving medical scientists from developing nations the chance of working as research fellows in world-renowned laboratories. Our goal: to train the future investigators that will become regional and global leaders for international collaboration in clinical research and medical education. As our training program has grown steadily throughout the years, so has our class of fellows that have had the opportunity to join us in Boston. In only three years, more than forty young doctors from many parts of the world have become part of this ever-growing family of professionals committed to adding their contribution to the advancement of their field. After going through a laborious application process, which commonly starts every June, each February we welcome and assign fellows to either basic science, clinical, or translational research laboratories in Harvard Medical School-affiliated hospitals.

Thanks to the outstanding performance of past fellows, we have been able to increase, year after year, the number and variety of specialties offered. We are proud to be part of those changes that are taking place worldwide that will bring forward the scientific advancement of developing countries. By planting the seed through means of education and training, we feel certain that the drive force to scientific development will reverberate and expand globally.

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Editorial

We commend the editorial board of the International Journal of Medical Students on their noble initiative of providing the much needed resources to promote the scientific growth and contribution of doctors-in-training from around the world. We proudly share the same motivating force that looks for the development of well-rounded professionals committed to the progress of the medical sciences.


References
Knowledge, Attitudes and Practices towards Medication Use among Health Care Students in King Saud University

Abdullah T. Eissa.1,2

Abstract
Background: Health sciences students are expected to have appropriate knowledge and attitudes toward medication use. However, literary evidence of such expertise among health sciences students of King Saud University is unknown. This study was completed to assess the knowledge about medicines and behavior of health science students towards safe use of medications. It also aims to assess the health knowledge, attitude and practices of the students. Methods: This cross-sectional study used a questionnaire consisting of 24 questions. This was administered by the researcher between October and December 2009 in the colleges of medicine, dentistry, pharmacy, applied medical science and nursing of the King Saud University. The survey consisted of three parts: Ten questions assessed the students’ knowledge on drug safety (Part 1). Four questions assessed student attitude toward medication consultations by the pharmacist (Part 2) and ten questions involved medication use practices and consultation with pharmacists (Part 3). A stratified sampling method was used to select participants. Results: Pharmacy students had better medication knowledge compared to other health sciences students especially regarding antihypertensive drugs, antibiotics, paracetamol and antacids (p<0.05). Pharmacy students showed a positive attitude regarding the trustworthiness of a pharmacist to give a consultation. Nearly all other health science students showed a negative attitude about dispensing and consultation concerning nutritional supplements by a pharmacist. All health sciences students had a similar perception toward medication use and practice. Conclusion: Pharmacy students had better knowledge about medication practice compared to other health sciences students. All other health sciences students lacked the appropriate attitude and practice related to the safe use of medications.

Key Words: Health Knowledge, Attitudes, Practice, Students, Health Occupations (Source: MeSH-NLM)

Introduction
King Saud is one of the largest universities in Middle East, from which a large number of health care providers graduate. It is ranked first in the Arab and Islamic world, the Middle East and Africa, according to the international Spanish webometrics ranking system and Academic Ranking of World Universities. Health sciences students’ knowledge about over-the-counter prescription medications and herbal products use is very important.1

An American national survey showed that medical students have huge gaps in knowledge towards the health care system, and the students feel that these deficiencies are not appropriately treated in the medical school curriculum.4

A questionnaire-based survey for first-year medical students of the Arabian Gulf University, Bahrain (including some Saudi students) suggested that these students had a poor knowledge about adequate self-medication whereas the knowledge of medication usefulness and harms was adequate. The attitude towards self-medication was positive and although the practice of self-medication was common, it was in most cases inadequate.1

About 83% of American medical schools include teaching on complementary and alternative medicine in their curricula, mostly in the form of electives.7 In addition, the Centers for Disease Control and Prevention have identified that colleges and universities are essential settings to deliver health education and services.7

The growing direct-to-consumer advertising of medicinal products targets the young population, a generation exposed to large amounts of media directing them to self-medicate. Lack of correct knowledge about medicines may directly lead to dangerous outcomes, such as overuse or non-compliance to treatment programs.8 For example, early self-discontinuation of antibiotics can lead to drug resis-
tance and challenges to effective therapeutics for future infections.7

Health science students, including medical, dental, pharmacy and nursing students, are expected to have appropriate knowledge and attitudes toward medication use to prevent inappropriate and harmful use of the medications. For this reason, the study objective was to assess the knowledge, practice and attitude of health care students of King Saud University (KSU).

Materials and Methods
This cross-sectional study utilized a structured, self-administered questionnaire that intended to evaluate the knowledge and behavior of medical, dental, pharmacy and nursing students at King Saud University toward common medication in Saudi Arabia. The survey was validated to ensure clear questions and simple and comprehensible language.

The questionnaire was distributed in paper and electronic format and was divided in three parts. (1) Questions to evaluate health science students’ knowledge of drug safety, (2) questions to assess students’ attitude toward medication consultations by the pharmacist, and (3) questions to identify medication use practices and consultation with pharmacists.

Part one utilized true and false questions to assess drug safety knowledge in five critical areas in our country: (1) anti-hypertensive drugs and regimens, (2) antibiotic use, (3) antacid use, (6) medication storage, and (7) vitamins, non-prescription drugs and herbal product use.

Part two included the items of the attitude assessment that were four questions answered to indicate how students trust pharmacists regarding the consulting and dispensing of medications, herbs and food. Items were scored on a five-point Likert scale, where higher scores indicated more positive attitudes toward medication consultation provided by pharmacists.

Part three included ten items of the practice questionnaire that were divided into three subscales: two items on self-care management, four items on appropriate use, and four items on prescription filling and medication consultation with a pharmacist. Items were scored on a five-point Likert scale. The coding for medication consultation with a pharmacist was the reverse, where (1) was never, (2) seldom, (3) sometimes, (4) usually, and (5) always. A higher total score represented better practices associated with safer medication use.

Table 1. Distribution of pharmacy, nursing, applied medical science, medical, and dentistry students in self-administered questionnaire by percent and frequency.

<table>
<thead>
<tr>
<th>Students</th>
<th>Percent</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pharmacy students</td>
<td>37.3%</td>
<td>76</td>
</tr>
<tr>
<td>Nursing students</td>
<td>15.2%</td>
<td>31</td>
</tr>
<tr>
<td>Applied medical science students</td>
<td>18.1%</td>
<td>37</td>
</tr>
<tr>
<td>Medical students</td>
<td>27.5%</td>
<td>56</td>
</tr>
<tr>
<td>Dentistry students</td>
<td>2.0%</td>
<td>4</td>
</tr>
<tr>
<td>Total</td>
<td>100.0%</td>
<td>204</td>
</tr>
<tr>
<td>Pharmacy students</td>
<td>37.3%</td>
<td>76</td>
</tr>
<tr>
<td>Non-pharmacy health students</td>
<td>62.7%</td>
<td>128</td>
</tr>
</tbody>
</table>

An overall internal consistency reliability coefficient (Spearman’s rho) was calculated for the questionnaire. Respondents were classified into two groups, which included the pharmacy students and other health students.

Results
Of 450 surveys distributed to five different health college students of King Saud University, including the colleges of medicine, pharmacy, nursing, applied medical science, and dentistry, 204 surveys were completed and returned to the researcher. Table 1 shows the distribution of respondents among different health students to compare the responses of 76 (37.3%) pharmacy students with 128 (62.7%) other health sciences students which includes 56 (27.5%) medical students, 31 (15.2%) nursing students, 37 (18.1%) applied medical science students and 4 (2.0%) dentistry students. Table 2 shows health sciences students’ responses to questions regarding knowledge of medication use. Although pharmacy students had better knowledge about medication use in the majority of the questions, the main difference between both groups was knowledge about anti-hypertensive drugs, antibiotics, paracetamol, antacids and drug interactions (p<0.05). In the question relating to the antihypertensive medication, pharmacy students performed significantly better than other colleges (p=0.000). In the question relating to the antibiotics, pharmacy students demonstrated a significantly higher level of knowledge compared to other health college students (p=0.013). In addition, the question regarding the toxicity of paracetamol and antacids and the interference of drugs with other substances showed significant differences between the two groups (p=0.013, p=0.000, p=0.000, respectively). In the remaining questions, which were related to antacid administration, vitamins, storing of ointment, storing of syrup and dosage of cough syrup, there were no significant differences between the two groups.
Table 2: Responses of pharmacy, nursing, applied medical science, medical, and dentistry students regarding knowledge of medication use by percent and frequency (N=204).

<table>
<thead>
<tr>
<th>Question</th>
<th>Pharmacy students</th>
<th>Other health students</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 - Antihypertensive drugs could be discontinued when blood pressure returns to a normal range.</td>
<td>10/39(25.6%)</td>
<td>12/39(30.8%)</td>
<td>0.404</td>
</tr>
<tr>
<td>2 - You can discontinue the use of antibiotics by yourself when the symptoms of fever or sore throat are relieved.</td>
<td>5/39(12.8%)</td>
<td>24/39(61.6%)</td>
<td>0.013</td>
</tr>
<tr>
<td>3 - Overdose of Panadol® (acetaminophen) will cause liver toxicity.</td>
<td>59/60(98.3%)</td>
<td>3/60(5%)</td>
<td>0.013</td>
</tr>
<tr>
<td>4 - Antacids should be chewed before swallowing to achieve a better effect.</td>
<td>29/39(74.4%)</td>
<td>10/39(25.6%)</td>
<td>0.000</td>
</tr>
<tr>
<td>5 - Antacids should be added into all prescriptions to avoid GI upset.</td>
<td>7/39(17.9%)</td>
<td>32/39(82.1%)</td>
<td>0.000</td>
</tr>
<tr>
<td>6 - Vitamins are a health food, so overusing them will not cause negative effects to the human body.</td>
<td>13/39(33.3%)</td>
<td>26/39(66.7%)</td>
<td>0.292</td>
</tr>
<tr>
<td>7 - Storing ointment or gel in the refrigerator could extend the expiration date.</td>
<td>28/39(71.8%)</td>
<td>12/39(30.8%)</td>
<td>0.879</td>
</tr>
<tr>
<td>8 - Storing syrup in the refrigerator could extend the expiration date.</td>
<td>30/39(76.9%)</td>
<td>9/39(23.1%)</td>
<td>0.284</td>
</tr>
<tr>
<td>9 - Dosage of cough syrup is one bottle per week.</td>
<td>20/26(77.0%)</td>
<td>6/26(23.1%)</td>
<td>0.872</td>
</tr>
<tr>
<td>10 - Taking some medicines with food, drink or tea will interfere with the effect of medicine.</td>
<td>70/92(76.1%)</td>
<td>22/92(23.9%)</td>
<td>0.000</td>
</tr>
</tbody>
</table>

Table 3 shows health college students’ attitudes toward medication consultation with a pharmacist. Pharmacy students showed a positive attitude and agreed with the perception that a pharmacist is a trusted person to provide consultation about drugs while other health sciences students tended to neither agree nor disagree. Regarding dispensing medication and consultation by a pharmacist, other health sciences students showed negative attitudes, while pharmacy students did not. Other health sciences students also showed negative attitudes regarding herbal medicine consultation by pharmacists. Both pharmacist and other health sciences students showed negative attitudes about dispensing and consultation of nutritional supplements by the pharmacist.

Table 4 shows medication use and practice among pharmacy and other health college students. Both pharmacy students and other health sciences students had the same practice level toward seeking non-prescription medications from community pharmacies, giving prescription medicine to others and combining herbal medication with prescription medicine. Other related questions provided similar responses between both groups.

Discussion
Pharmacy students demonstrated higher levels of knowledge about medication practices compared to other health sciences students according to the significance of the results of many questions. Results shown in Table 2 suggest that patients should seek the appropriate management from licensed physicians or pharmacists. Despite the fact that nurses, dentists, and applied science specialists are...
considered health care providers, they cannot prescribe and consult patients regarding medication-related concerns. In addition, Table 2 suggests that the curricula of these health colleges do not provide appropriate teaching of pharmacology. Undergraduate pharmacy students seem to still be unable to deal with all medications appropriately because there were some questions that were not solved correctly by the majority of them. Dental students’ responses suggested little knowledge about the items of the questionnaire, leading them to be classified similarly to the general public.

Table 3 shows that other health college students express neutral responses when asked about the perceived need to consult a pharmacist to adequately use and dispense medications, herbals, and food products. This is considered an inappropriate attitude because the use of medications, herbals, and food products to some extent is considered to fall under the responsibilities of pharmacists. Pharmacy college students agreed that the pharmacists are knowledgeable and can dispense drugs but the students lacked an appropriate attitude towards consulting pharmacists for the use of herbal and food products.

Table 4 shows that both pharmacy students and other health students lacked appropriate practices related to the safe use of medications. That may be because the curricula emphasize gaining knowledge more than clinical practice. Further training and interventions are warranted to improve the attitude and practice related to the safe use of medications in both student groups.

References

Acknowledgements
Thanks to Mansour Adam for his advice.

Conflict of Interest Statement & Funding
The Authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as:
Survival and Prognostic Factors in Adults Acute Myeloid Leukemia: A Retrospective Analysis of 119 Cases

Alina Maria Gridjac,1 Cristian Daniel Pirlog,2 Anca Simona Bojan3

Abstract

Background: Acute myeloid leukemia (AML) is a malignant disease with significant identified prognostic factors. Therefore our aim was to develop an Assessment Scheme of Prognosis in AML based on prognostic factors. In some countries, such as Romania or other less-highly developed countries, this scheme would be beneficial particularly when cytogenetic testing is unavailable or time-intensive. Methods: We analyzed 119 adult patients with AML during a five year-period from a single-center in Romania. We retrospectively collected and analyzed data with Epi info and Excel using patient medical records. Results: According to age, the group A1 (<60 years) had a 40 months survival, in contrast with the group B1 (≥60 years) with a survival of 19 months (p=0.0063). The group A2 (secondary AML) survived 15 months, whereas the group B2 (AML de novo) survived 40 months (p=0.0021). Additionally, the group A3 (mild comorbidities) achieved a 40 months survival, the group B3 (moderate comorbidities) survived 19 months, whereas the group C3 (severe comorbidities) survived 7 months (p=0.0059). According to WBC and blast number, the group A4 (high levels) had a 25 months survival, whereas the group B4 (low levels) survived 40 months (p=0.0057). Conclusion: The prognostic factors studied are useful to identify the risk level of AML disease for each patient at diagnosis. We developed an assessment scheme of prognosis with three risk groups according to age, secondary AML, comorbidity, WBC and blasts and cytogenetic examination.

Keywords: Acute Myeloid Leukemia, Survival Analysis, Prognosis, Risk Assessment, Age Factor (Source: MeSH-NLM)

Introduction

Acute myeloid leukemia (AML) is a clonal disease characterized by the proliferation and accumulation of myeloid progenitor cells in bone marrow, which ultimately leads to hematopoietic failure.1,2 The incidence of AML increases with age, and older patients typically have worse treatment outcomes than younger patients.3 Diagnosis of AML is based on cellular morphology, immunology, cytogenetics, and molecular features.4

A number of prognostic factors have been identified in AML, including age, performance status, organ dysfunction, secondary AML, white blood cell and blast count at presentation, karyotype, and molecular abnormalities.5,6 Most relevant studies, based on large multicenter trials have definitively demonstrated that age and cytogenetics at diagnosis are the most significant prognostic determinants for patients with AML.7

This study focuses on survival rate, prognostic factors and the evolution of AML in patients. The purpose of this study was to formulate an Assessment Scheme for Prognosis of AML. Even if these prognostic factors were previously studied our goal was to develop a score, which is easy to calculate, for early prognostic assessment at diagnosis. Especially in Romania or other less-highly developed countries, this scheme would be beneficial particularly when cytogenetic testing is unavailable or time-intensive.

Methods

We performed a retrospective longitudinal study on 119 patients, aged 21-85 years with a median age of 60. Using statistical tests such as Excel and Epi Info we managed to associate, analyze and compare the variables. Additionally, this study was analytical and observational.

The length of the study was five years, from January 2007 to November 2011. The main feature of all patients was the presence of AML. The demographic characteristics included the patients admitted at the Department of Hematology of the Oncology Institute “Prof. Dr. Ion Chişcitu”, Cluj-Napoca, Romania. Although it was a single-center study, it comprised a whole region of Romania, because Cluj-Napoca is the most important medical centre in Transilvania.
We aimed to identify the most important prognostic factors, therefore to perform a risk assessment for patient with AML. According to Kaplan-Meier’s method, we analysed the patient’s survival rate to illustrate the differences between specific variables and the characteristics of the patients. We formulated five research hypotheses, which confirmed a shorter survival for 60 year olds, secondary AML patients, patients with severe comorbidities and patients with elevated levels of WBC and blasts.

The data used in the study comprised: gender, age, place of origin, date of admission, FAB type, WHO classification, cytogenetic examination, immunophenotyping, laboratory data (hemoglobin, erythrocytes, leukocytes, platelets, LDH, blasts), symptoms, comorbidity, other malignancies, dysplasia, treatment, complications, date of death and status censorship. Status of censorship coding included two groups of patients. The first group was “censored” if at the end of study, some patients were lost, whereas the second group was “complete” if during the study, some patients died.

Almost 14% presented with other malignancies requiring radio-chemotherapy. As a result, patients with these malignancies developed secondary AML. Moreover, half of all cancer patients presented with genitourinary malignancy whilst a third were gastrointestinal in origin. Analysis of the survival of primary and secondary AML patients revealed significant differences. As shown in Figure 2, there was a 15 month survival for the first group (A2 =Secondary AML) and 40 month survival for the last group (B2 =AML de novo, p=0.0021).
A proportion of 64% of cases had associated comorbidities as they were present especially within the elderly patients. The most frequent were the heart diseases, which comprised 60 cases. Digestive and respiratory illnesses ranked next with 28 and 21 cases respectively. The next frequent disease was diabetes with 19 cases and the less frequent diseases were renal and neurological disorders with 10 and 9 cases respectively. We separated patients, according to comorbidities into three groups seen in Figure 3. Group A3 comprised patients with mild comorbidity with a survival of 40 months. Group B3 covered patients with moderate comorbidities who survived 19 months, whereas group C3 included patients with serious comorbidities who survived only 7 months (p=0,0059).

As shown in the Figure 4, in Epi Info, we analyzed two categories: group A4 = WBC >15,000/mm³ + blasts >40% and group B4 = WBC >15,000/mm³ + blasts >40%. Survival in group A4 was up to 40 months, whereas the group B4 had a survival of 25 months (p=0,0057).

Only 26% of patients received cytogenetic examination. The prognosis distribution of results followed the WHO classification, therefore 71% of patients correspond to intermediate risk, whereas 29% of patients correspond to unfavorable risk with chromosomal abnormalities (Figure 5).

Discussion

Various clinical and biological features identified previously as useful factors in the prediction of clinical outcomes, could help guide therapy choices. A subgroup of patients with hyperleucocytosis was identified with a shorter rate of survival in these cases. In our study, many laboratory analyses were performed, but we chose to combine WBC and blast count for a significant difference on our patients survival. Patients were divided into groups according to low or high levels of WBC and blasts. Patients with values of WBC and blasts >40% had much shorter survival than those with low values.

Karyotype abnormalities are probably the most important prognostic determinants in AML. It is reported that half of adult AML patients present with a normal karyotype. In our study, a proportion of 71 per cent presented a normal karyotype and 29 per cent had unfavorable cytogenetic abnormalities. In our study, we were unable to demonstrate the significance of cytogenetic testing due to the small number of patients who benefited from cytogenetic examination. Finally, we propose that all AML patients should receive cytogenetic examination for a better management of AML.

Our single-center study evaluated and confirmed the significant predictors of outcome of AML. The age of patients, the presence of secondary AML, comorbidities and WBC count and blasts were the most important prognostic parameters. Furthermore, we demonstrated this in our study illustrated a surprisingly low life expectancy. As a result, patients with AML de novo lived twice as long as the patients with secondary AML.
Finally, this classification would also help direct the precise therapy for this patient. Furthermore, we can assign patients with AML according to their prognosis, to categorize patients into risk groups. Moreover, we are going to check this scheme on a larger patient population to demonstrate its significance. On some patients we have already evaluated this scheme and we obtained a precise prediction of survival rates among the different groups of patients.

Conclusion

The understanding of dominant prognostic factors in patients with AML is rapidly evolving, but until we find more accurate factors, we should clarify the known ones. This single-center study evaluated the significance of pretreatment factors, and found that the age of patients, the presence of secondary AML, comorbidities, WBC, blasts and cytogenetics were factors which influenced the outcomes of survival in patients with AML. Therefore, we propose an Assessment Scheme of Prognosis for patients with AML, which would be useful to apply a therapeutic regimen and to categorize patients into risk groups. Furthermore, we can assign patients with AML according to their prognosis, in three categories: Low risk, intermediate risk and high risk. We included cytogenetic examination with data from medical literature, as we were not able to come up with our own results, due to our small number of patients who received cytogenetic examination. Moreover, we are going to check this scheme on a larger patient population to demonstrate its significance. On some patients we have already evaluated this scheme and we obtained a precise prognosis. The scheme works by adding the points if the factor is present. For instance, a patient who is aged ≥60 years (3 points), has AML de novo (1 point), moderate comorbidities (2 points), WBC=15000-100.000/mm3 and blasts=40-80% (2 points) and an unfavorable chromosomal abnormality (3 points) would be classified as high risk. Finally, this classification would also help direct the precise therapy for this patient.

Acknowledgements

None.

Conflict of Interest Statement & Funding

The Authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as:


References

Health Care Expenditure of Rural Households in Pondicherry, India

Poornima Varadarajan, Lopamudra Moharana, Murugan Venkatesan

Abstract
Background: Shortcomings in healthcare delivery has led people to spend a substantial proportion of their incomes on medical treatment. World Health Organization (2005) estimates reveal that every year 25 million households are forced into poverty by illness and the struggle to pay for healthcare. Thus we planned to calculate the health care expenditure of rural households and to assess the households incurring catastrophic health expenditure. Methods: A cross-sectional study was conducted in the service area of Sri Manakula Vinayagar Medical College and Hospital from May to August 2011. A total of 100 households from the 4 adjoining villages of our Institute were selected for operational and logistic feasibility. The household’s capacity to pay, out of pocket expenditure and catastrophic health expenditure were calculated. Data collection was done using a pretested questionnaire by the principal investigator and the analysis was done using SPSS (version 16). Results: The average income in the highest income quintile was Rs 51,885 but the quintile ratio was 14.98. The median subsistence expenditure was Rs 4,520. About 18% of households got impoverished paying for health care. About 81% of households were incurring out of pocket expenditure and 66% were facing catastrophic health expenses of 40%. Conclusion: There was very high out of pocket spending and a high prevalence of catastrophic expenditure noted. Providing quality care at affordable cost and appropriate risk pooling mechanism are warranted to protect households from such economic threats.

Key Words: Health expenditures, health services needs and demand, India (Source: MeSH-NLM).

Introduction
The promotion of health is of fundamental value in itself. It is a vital public good and a basic human right. In this regard, delivery of healthcare is very important for providing preventive, promotive and curative services to the community.

There have been substantial achievements in healthcare in past few decades. However, technological innovation in the health sector has improved the quality of life but has also increased costs especially in middle and low income countries. Shortcomings in healthcare delivery have been largely designated as fragmented care, misdirected care and impoverishing care. In countries that have no social insurance and where the role of the state is limited, people spend a substantial proportion of their incomes on seeking medical treatment, and in the process get impoverished, thus widening disparities in the health status. The unpredictability of illness, the lumpiness of health consumption, and the irregular and seasonal nature of incomes make it virtually impossible for the poor to finance their health needs, resulting in a denial of care and poverty.

According to the World Health Organization (2005) estimates, every year 25 million households (more than 100 million people) are forced into poverty by illness and struggle to pay for healthcare. The decline in public investment in health and the absence of any form of social insurance have heightened insecurities. Considering the Indian scenario, a report by the National Health Accounts reveals that 71% of the health budget is contributed by the private sector; of which households alone spend about 69%. It is well known that health expenditure in India is dominated by private spending and this is a reflection of inadequate public spending. The relationship between poverty and ill-health is indisputable. Even relatively small expenditure on health can be financially disastrous for poor households. High out of pocket payment, an absence of risk pooling mechanism in health financing systems, and high level of poverty can result in catastrophic health expenditure. Thus, the present study was conducted to quantify the health care expenditure of households in rural Pondicherry with these objectives: i) to note the health care expenditure of rural households and ii) to assess if any family is undergoing catastrophic health expenditure.

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Materials and Methods

Study Setting

Pondicherry (Puducherry), district of India has a total population of 946,600 (census 2011) of which the rural population is about 292,208 (30.87%). Per-capita income of Pondicherry at 2009-10 current prices is in Indian rupee (Rs) 72,917 (draft annual plan 2011-12, Pondicherry) and about 22% are below poverty line (2004-05 Pondicherry estimate). Pondicherry ranks quite high compared to India in terms of fulfillment of several health infrastructure indicators. The total health care expenditure is Rs 80 but the per-capita expenditure is Rs 783 (budget estimate, 2003-04).

Study Design and Sampling

We planned to study the total health spending so that the burden on households can be commented on. Moreover, to get a better picture of rural areas we studied the rural households. A cross-sectional study was designed and conducted from May to August 2011. We covered four villages, situated within 2-4 kilometers of our Institute (Sri Manakula Vinayagar Medical College and Hospital), for operational and logistic feasibility. The villages were Madagadipet (1724), Kalitheerthalkuppam (1120), Kuchipalayam (158) and PS palayam (441) which constituted our sampling frame and the list of houses of the respective villages were collected from the PHC registers (2011). We planned to survey a total of one hundred households. From the total number of households of the four villages (2892), all sampled households were included in the study. A proportionate sampling method was adopted to draw the sample from the household numbers of each village. The streets in each village were selected by simple random technique. Systematic random sampling method was adopted to select every third household in each street. Initially, a pilot study was conducted to assess and modify the logistic problems expected during the main study.

Parameters

The total household income and expenses were calculated. For expenditure on health, both direct and indirect expenses were assessed. Direct expenses were costs incurred for the defined medical problem (consultations, investigations, medicines etc.) and the indirect cost included collateral expenses due to the illness (travel, food, loss of wages etc.). We considered the expenses incurred by the households both for their outpatient and inpatient consultations. To avoid recall bias in expenditure, a one month recall period for any OPD consultations and 3 months for any in-patient admissions was considered. For our calculations, we used equivalent household size (household size 0.56) instead of average household size. The various heads of expenses of the households were determined and for accuracy of food expenditure, equivalized food expenditure was calculated. The subsistence expenditure per (equivalent) capita or the poverty line was determined and the subsistence expenditure (poverty line*equivalent household size) was calculated. The household’s capacity to pay (non subsistence effective income of households), out of pocket expenditure (OOP), burden of health payment and catastrophic health expenditure (CHE) was calculated adopting the methodology described by Xu K et al. OOP expenditure is defined as the payment made by families for health care and include out of pocket spending on deductibles and other forms of cost sharing such as co-payments and co-insurance and direct expenditure of health care services equipments and supplies not covered by insurance. OOP in our study was the net of insurance reimbursements and did not include indirect expenses (health-related travel and food). CHE is defined as the level of OOP expenditure that exceeds some fixed proportion of household income or household capacity to pay. For the purpose of our study, if a household’s total OOP equaled or exceeded 40% of the household’s capacity to pay (non subsistence effective income of the household or income available after basic needs have been met), it was considered to be facing CHE. We also calculated the households that are poor (total household expenditure less than its subsistence spending) and the non-poor households that were impoverished by health payments.

Results

The socio-demographic profile of the total households surveyed is described in Table 1. We noted that the majority belonged to the Hindu religion (94%). The various castes were OBC (30%), MBC (44%) and the forward castes (6%). About 81% houses were pucca houses and the majority (93%) were nuclear families. The median “total” and “per-capita” income of the households were Rs 10,000 and Rs 2,333 respectively. The average income in the highest (5th) quintile was Rs 51,885 but the “quintile ratio” (richest to poorest) was 14.98. A majority (72%) possessed pink ration cards and 8% did not have any ration card. However, those who participated in different income-generating activities and in self-help groups were 7% and 2% respectively. None of the households had health insurance. The “equivalent household size” was 2.26 and the median “equivalized per capita household expenditure” was Rs 2323 where as the median “equivalized food expenditure” was Rs 1379. From the food expenditure, the subsistence expenditure per (equivalent) capita or the poverty line was calculated as Rs 2,080 and thus the median “subsistence expenditure” was calculated to be Rs 4,520.

The health facilities preferred by the majority of the households were governmental (24%) and private (74%). About 1% preferred both private and governmental facilities and the remaining 1% preferred chemist shops. About 69% had reported outpatient illnesses and 35% had inpatient illnesses within our specified period of recall. The common outpatient illnesses were fever, cough and cold, diarrhea, body and joint pains, gastritis, poor vision, pregnancy, allergy, TB,
Table 1. Socio-demographic characteristics of the study participants

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>%</th>
<th>n = 100</th>
</tr>
</thead>
<tbody>
<tr>
<td>Religion</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hindu</td>
<td>94</td>
<td></td>
</tr>
<tr>
<td>Muslim</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Christian</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Housing type</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pucca</td>
<td>81</td>
<td></td>
</tr>
<tr>
<td>Kutcha</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Semipucca</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Family type</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nuclear</td>
<td>93</td>
<td></td>
</tr>
<tr>
<td>Joint</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>Possession of ration cards</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pink</td>
<td>72</td>
<td></td>
</tr>
<tr>
<td>Yellow</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>No card</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Income generating activities</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>93</td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>Participation in SHG</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>98</td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Health Insurance</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Income Quintile</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1st Quintile (Rs 3462)*</td>
<td>26</td>
<td></td>
</tr>
<tr>
<td>2nd Quintile (Rs 6875)*</td>
<td>18</td>
<td></td>
</tr>
<tr>
<td>3rd Quintile (Rs 10,188)*</td>
<td>16</td>
<td></td>
</tr>
<tr>
<td>4th Quintile (Rs 16,781)*</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>5th Quintile (Rs 51,885)*</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>Quintile Ratio (Richest to poorest)</td>
<td>14.98</td>
<td></td>
</tr>
<tr>
<td>Total income (Median)</td>
<td>10,000</td>
<td></td>
</tr>
<tr>
<td>Per capita monthly income (Rs, median)</td>
<td>2333</td>
<td></td>
</tr>
<tr>
<td>Equivalent household size’ (Mean ± SD)</td>
<td>2.26 ± 0.4</td>
<td></td>
</tr>
<tr>
<td>Equivalized household expenditure (Rs per capita, median)</td>
<td>2323</td>
<td></td>
</tr>
<tr>
<td>Equivalized food expenditure’ (Rs, median)</td>
<td>1379</td>
<td></td>
</tr>
<tr>
<td>Subsistence expenditure (Rs, median)</td>
<td>4520</td>
<td></td>
</tr>
</tbody>
</table>

* Figures in parentheses are average income in the corresponding income quintile.

* Equivalent household size = 1/6 (Adopted from Xu K et al.).

* Equivalized food exp = Food expenditure / equivalent household size.

Abbreviations: OOP: Out of Pocket expenditure, Rs: Indian rupee, SHG: Self-help group, SD: Standard deviation.

Table 2. Reported illness, capacity to pay and consequence of health expenses incurred per households.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>%</th>
<th>n = 100</th>
</tr>
</thead>
<tbody>
<tr>
<td>Health facility preferred</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Private</td>
<td>74</td>
<td></td>
</tr>
<tr>
<td>Governmental</td>
<td>24</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Chemist shop</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Outpatient illness (Reported within 2 months)</td>
<td>69</td>
<td></td>
</tr>
<tr>
<td>Inpatient illness (Reported within 3 months)</td>
<td>35</td>
<td></td>
</tr>
<tr>
<td>Household spending on health (n=84, who reported any illness, as % of their total income)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;20%</td>
<td>35</td>
<td></td>
</tr>
<tr>
<td>20-50%</td>
<td>14</td>
<td></td>
</tr>
<tr>
<td>&gt; 50%</td>
<td>35</td>
<td></td>
</tr>
<tr>
<td>Household’s capacity to pay</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rs 10,000</td>
<td>71</td>
<td></td>
</tr>
<tr>
<td>Rs 10,000-30,000</td>
<td>18</td>
<td></td>
</tr>
<tr>
<td>Rs &gt;30,000</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>OOP incurred</td>
<td>81</td>
<td></td>
</tr>
<tr>
<td>OOP share of total income</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤50%</td>
<td>48</td>
<td></td>
</tr>
<tr>
<td>&gt;50%</td>
<td>33</td>
<td></td>
</tr>
<tr>
<td>OOP share of total expenses</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤50%</td>
<td>33</td>
<td></td>
</tr>
<tr>
<td>&gt;50%</td>
<td>48</td>
<td></td>
</tr>
<tr>
<td>OOP share of capacity to pay</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤20%</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>20-40%</td>
<td>15</td>
<td></td>
</tr>
<tr>
<td>&gt;40%</td>
<td>60</td>
<td></td>
</tr>
<tr>
<td>Poor (Below poverty line)</td>
<td>29</td>
<td></td>
</tr>
<tr>
<td>Impoverishment (Due to health expenses)</td>
<td>18</td>
<td></td>
</tr>
</tbody>
</table>

* Note: Median OOP spending per household was Rs 3000. Median subsistence spending (expenditure) of the households was Rs. 4520. The households’ capacity to pay is calculated as the non-subsistence effective income of households. CHE is calculated as OOPCTP ≥ 40%. About 18% are impoverished paying for their healthcare expenses. Abbreviations: OOP: Out of Pocket expenditure, Rs: Indian rupee.

Table 3. Prevalence of Catastrophic Expenditure by Cut off Levels (of OOPCTP)

<table>
<thead>
<tr>
<th>Households' OOPCTP</th>
<th>No of Households (n = 81)</th>
<th>% of Households with illness</th>
<th>Household Expenditure/m (Median)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 20%</td>
<td>6</td>
<td>7</td>
<td>520</td>
</tr>
<tr>
<td>20-40%</td>
<td>15</td>
<td>19</td>
<td>600</td>
</tr>
<tr>
<td>&gt;40%</td>
<td>60</td>
<td>74</td>
<td>8025</td>
</tr>
</tbody>
</table>

* CHE, i.e. OOPCTP ≥ 40% was incurred by 66%.

* Illnesses include both outpatient and inpatient categories.

Abbreviations: OOP: Out of Pocket expenditure, OOPCTP: OOP share of capacity to pay.
diabetes and hypertension. The common inpatient illnesses were accidents, abdominal pathologies and major abdominal surgeries, obstetric and gynecological complaints, hernias, fractures, tumors and non healing ulcers. Out of the households who reported some illness (84%), about 49% were spending about 50% of their total income and 35% were spending even >50% of their incomes (direct and indirect expenses) on health (Table 2).

Analysis of the households’ capacity to pay showed that the majority had the capacity to pay up to Rs 10,000 (71%) while the remaining had the capacity to pay more. About 81% of the households were incurring OOP expenses (only direct health expenses were considered) and the median OOP spending per household was Rs 3,000 (Table 2). For 33% of households, the OOP incurred was 50% of their total expenses but for 48% of households, OOP incurred was even >50% of their total expenses. About 29% of the households whose total household expenditure was less than their subsistence spending were already poor, but about 18% were impoverished by paying for health expenses (Table 2).

The OOPCTP of ≥40%, i.e., CHE was incurred by 66% of households (Table 2). The percentage of households at catastrophic threshold of <20%, 20-40% and >40% cut off levels were 6%, 15% and 60% respectively (Table 3). We noted that households with higher proportion of reported illnesses and those belonging to higher median household expenditure categories were incurring high CHE. Across the different expenditure quintiles (Table 4, Figure 1), the median health expenditures and median OOP were noted to increase gradually. The OOPCTP and the OOPEXP were highest in the higher expenditure quintiles (4th and 5th quintiles). The OOPEXP was even more than 100% in the highest (5th) expenditure quintiles. We also noted that there was impoverishment in the middle expenditure quintiles (3rd and 4th); whereas lowest and highest ones did not show impoverishment. Table 5 represents a picture of the households under poverty and those who got impoverished by paying for health care expenses across income quartiles. We noted that the highest number of poor households (65%) were in the first quartile and the tendency gradually decreased towards the highest quartile. On the contrary, households that faced impoverishment (paying for health care expenses) were 27%, 21% and 24% in the 2nd, 3rd and 4th quartile respectively. There was no impoverishment noted in the first quartile.
We present the burden of health payments suffered by the rural households of Pondicherry, India. Information of household expenditure was gathered, particularly on healthcare expenses with respect to their total incomes and expenditures. Even though, most of the households belonged to Prasad’s class I (48%) of socio economic status, the high quintile ratio (14.98) indicates a gross inequality among the richest and poorest quintiles. About a third of households had a greater subsistence spending than the total expenditure (29%, poor), which unfortunately was also accompanied by scarce participation in income-generating activities and no health insurance benefits.

It was observed that within the specified recall period of our study, households that reported some illness were spending a major portion of their income on health. Considering the unpredictability and increased frequency of illnesses, health expenditure amounts to a major burden on the households. However, due to the cross-sectional nature of the study we could not determine the frequency of illness. Additionally, preference for private health care facility was supposed to pose a great burden on the households because it usually incurred higher healthcare costs. According to the emerging market report in India from 2007, the private sector accounts for more than 80% of total healthcare spending. Unless there is a decline in the combined federal and state government deficit, the opportunity for significantly higher public health spending would be limited. The majority of households in our study had low capacity to pay (71%) and most preferred a private health care facility (74%). On the other hand, households were spending even more than half of their total expenses on healthcare from out of pocket and the burden of payment faced was high. We also noted that most households in the first income quartile were already poor (65%) whereas those that got impoverished were more likely to be in the higher income quartiles. It could be due to the fact that higher income groups spent more for their healthcare because of higher affordability. This was further evidenced as shown in Table 4 and Figure 1; median health expenditure increased across the expenditure quintiles and OOP and OOPCTP also increased. The high burden of OOP was remarkable in our study; even well above households’ total expenses in the highest expenditure quintile (127%).

Table 5. Poverty and impoverishment across income quartiles (n = 100)

<table>
<thead>
<tr>
<th>Income Quartiles</th>
<th>Households (No)</th>
<th>Poverty, n (%)</th>
<th>Impoverishment, n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st</td>
<td>26</td>
<td>17 (65)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>2nd</td>
<td>30</td>
<td>7 (23)</td>
<td>8 (27)</td>
</tr>
<tr>
<td>3rd</td>
<td>19</td>
<td>4 (21)</td>
<td>4 (21)</td>
</tr>
<tr>
<td>4th</td>
<td>25</td>
<td>1 (4)</td>
<td>6 (24)</td>
</tr>
</tbody>
</table>

Discussion

According to a report of the National Commission on Macroeconomics and Health 2005, households undertook 75% of all health spending in the country. Analysis of health care spending in union territories of India in 2004-05 revealed the per-capita health expenditure to be Rs 598 with households spending about 85% and the government spending at just 9%. A study of catastrophic household expenditure on childhood illness in an urban slum of Karnataka, India, showed that all households were undergoing catastrophic expenses at a 5-20% threshold.

Conclusion

There was very high OOP spending and high prevalence of catastrophic expenditure noted in our study. Irrespective of the income and expenditure categories, households were incurring CHE and there was substantial amount of income spent on healthcare. We recommend improving the quality of primary care services to make it more accountable to community necessities which would minimize private healthcare expenses. Increased community awareness to participate in income-generating activities to strengthen their household economy is needed. There should be appropriate risk pooling mechanisms to protect households from incurring catastrophic expenses.
The Authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as
The Risk of Contact Lens Wear and the Avoidance of Complications

Farihah Tariq,1 Peter Koay2

Abstract
Contact lenses are lenses placed on the surface of the cornea to correct refractive errors such as myopia (short-sightedness), hypermetropia (far-sightedness) and astigmatism. Lens-related complications are becoming a greater health concern as increasing number of individuals are using them as an alternative to spectacles. Contact lenses alter the natural ocular environment and reduce the efficacy of the innate defences. Although many complications are minor, microbial keratitis is potentially blinding and suspected cases should be rapidly diagnosed and referred to an ophthalmologist for treatment. Several risk factors have been identified with extended wear, poor hand hygiene, inadequate lens and lens-case care being the most significant. Promotion of good contact lens hygiene and practices are essential to reduce the adverse effects of contact lens wear.

Key Words: Contact Lenses, Complications, Keratitis, Patient Compliance (Source: MeSH-NLM)

Introduction
Ametropic disorders of vision affect between 800 million to 2.3 billion individuals globally.1 Around 140 million users worldwide, including 3.3 million in the United Kingdom, wear contact lenses for the correction of refractive errors.2,3 The British contact lens market value has risen from £33 million in 1992 to £198 million in 2009.4 They are becoming increasing popular because of the clearer vision achieved, for cosmetic reasons, for sports and convenience. Contact lenses are, however a medical device and wearing contact lenses incurs risks with an estimated 6% of users developing complications.4 We will discuss the pathophysiology of contact lens-associated complications and their avoidance.

Search strategy and selection criteria
Soft contact lenses are the focus of this paper. In depth discussion on other types of lenses such as rigid gas permeable, PMMA lenses were out of the scope of this paper. We identified the papers in this review by a computerised search of the PubMed database using the queries “contact lens complications” and “contact lens keratitis”. We gathered other information from contact lens manufacturers’ data sheets and used evidence from published abstracts, major international scientific meetings and textbooks as well as reference collections.

How do contact lenses affect the ocular surface?
Contact lens wearers are sixty times more likely to develop ocular disorders than the general population, with the users of extended wear at greatest risk.5-8 An estimated 1 per 2500 persons per year using daily wear and 1 per 500 persons per year using extended wear will develop presumed microbial keratitis.9 Incidences of complications compiled by Morgan and colleagues is presented in Table 1.7

Contact lenses influence the allergic and inflammatory responses, alter the ocular microbiota, cause metabolic and mechanical trauma, reduce ocular surface wetting and can exacerbate pre-existing ophthalmic disorders.10,11

Contact lenses alter the natural ocular environment
Contact lenses are foreign objects in the eye, altering the natural environment by introducing a bio-burden of microorganisms to the ocular surface from contaminated hands, lens and lens-care solution.12-14 Insertion of the lens initiates the formation of a biofilm which not only attracts pathogenic flora but increases antibiotic resistance by almost one thousand fold.13 Bacteria adhere to the contact lens; this propensity is stimulated by deposits on the lens surface.14 Within 30 minutes of insertion, approximately 50% of the lens accumulates materials on or into the lens matrix.15 Such spoilage by the constituents of the tear film is not
The contact lens directly impedes oxygen transmission

Contact lenses cause micro-trauma attributed to hypoxia. The cornea receives oxygen fundamental to cellular function primarily through the atmosphere and a small quantity from the limbal and aqueous vasculature. 

Hypoxia causes oedema, altering the epithelial and endothelial morphology predisposing the cornea to cellular breakdown. Reduced oxygen permeability correlates with diminished corneal sensation and increased risk of keratitis. The greater oxygen permeable silicone hydrogel lenses have a five-fold reduced risk of severe keratitis compared with hydrogels.

Scarce distribution of oxygenated tear film due to reduced blinking whilst users are performing visual tasks like wor-
Non-associated blepharitis, or sterile infiltrates

With the natural barriers threatened, damage to the cornea arises. However, with the availability of contact lenses, particularly soft non-silicone hydrogel lenses, the corneal surface is under a constant threat of infection. Contact lenses introduce pathogens to the corneal surface, leading to decreased visual acuity, particularly if central visual axis is involved.24 Acute hypoxia can lead to overwear syndrome whilst chronic hypoxia can instigate corneal neovascularisation contributing to decreased visual acuity, particularly if the central visual axis is involved.14,25 However, with the availability of more permeable lenses such problems have been reduced.6

**Contact lenses introduce pathogens**

The corneal surface is under a constant threat of infection from a barrage of pathogens and at any instance up to 63% of contact lenses yield a positive culture consisting of normal commensals.26 Reduced efficacy of the defence mechanisms coupled with change in the concentration and variety of bacteria can contribute towards pathogenic processes.6 With the natural barriers threatened, damage to the intact cornea allows bacteria to adhere to the cell membrane; a vital step in the infectious process as it aids colonisation.25 Recent research has shown there is upregulation of surface-binding receptors further augmenting bacterial adherence.6 Contact lenses, particularly soft non-silicone hydrogel lenses, potentiate their infiltration by inducing changes in corneal epithelium (e.g. reduced desquamation and mitotic activity) making it thinner and increasing the risk of infection.24,28

Although a variety of organisms have been isolated from corneal infections, gram negative infections are most common and sight threatening.24,28 Infectious keratitis arising due to the ubiquitous Pseudomonas aeruginosa has the greatest associated morbidity.29 This is attributed to a large number of genes dedicated to virulence regulation, environmental adaption and resistance to antimicrobial drugs (e.g. aminoglycosides).29,30 Although rare, 5% of contact lens-related microbial keratitis is attributed to Acanthamoeba.11 This opportunistic pathogen is found in soil and air; but the main perpetrator is contaminated water (e.g. swimming pools, hot tubs, water tanks, lakes and contaminated cleaning solutions).29,31 Acanthamoeba exists in two forms; a feeding and replicating trophozite which can form amoebic-resistant dormant cysts.32,33 Acanthamoeba keratitis was associated with a poor prognosis before the introduction of topical polyhexamethylenebiguanide (PHMB), proamidine isethionate, and chlorhexidine; 30% of patients who had reduced visual acuity (6/18 or less), 50% underwent surgery whilst enucleation was performed in resistant cases.34,35 More recently, early diagnostic techniques and timely treatment with anti-amoebics have improved prognosis; 90% of patient retain visual acuity of 6/12 or better and less than 2% become blind.36

Correctly differentiating microbial keratitis from the less serious sterile corneal infiltrates is crucial.6 Sterile infiltrates tend to be present on the periphery and may be symptomatic or asymptomatic.4 They may be the consequence of lens wear itself, from bacterial endotoxins present in conditions related microbial keratitis is attributed to Acanthamoeba.11 This opportunistic pathogen is found in soil and air; but the main perpetrator is contaminated water (e.g. swimming pools, hot tubs, water tanks, lakes and contaminated cleaning solutions).29,31 Acanthamoeba exists in two forms; a feeding and replicating trophozite which can form amoebic-resistant dormant cysts.32,33 Acanthamoeba keratitis was associated with a poor prognosis before the introduction of topical polyhexamethylenebiguanide (PHMB), proamidine isethionate, and chlorhexidine; 30% of patients who had reduced visual acuity (6/18 or less), 50% underwent surgery whilst enucleation was performed in resistant cases.34,35 More recently, early diagnostic techniques and timely treatment with anti-amoebics have improved prognosis; 90% of patient retain visual acuity of 6/12 or better and less than 2% become blind.36

**What are the risk factors for developing contact lens wear complications?**

There are a range of modifiable and non-modifiable risk factors involved in the development of complications.7 Non-modifiable risk factors are younger age (<25 years), older age (> 50 years), male gender, diabetes mellitus, low socioeconomic class and late winter months.5,4,3,6,8,13,19 Modifiable risk factors are those which can be influenced or altered and includes improper lens and case care, poor hand hygiene, smoking, swimming and showering wearing lenses, as well as extended and overnight wear.5,8,34

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**Figure 2. Relative risks and non-compliance for a range of compliance and usage factors**

Adapted with permission from from Baush & Lomb, 2010.36

**Table 1. Incidence of contact lens-associated complications.**

<table>
<thead>
<tr>
<th>Contact lens type</th>
<th>Complications (per 10,000)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Non-severe</td>
</tr>
<tr>
<td>Daily wear hydrogel</td>
<td>5</td>
</tr>
<tr>
<td>Extended wear hydrogel</td>
<td>96</td>
</tr>
<tr>
<td>Extended wear silicon hydrogel</td>
<td>20</td>
</tr>
</tbody>
</table>

**Table 2. Avoiding contact lens-related complications**

1. Regular review by contact lens provider
2. Take hygiene instructions seriously
3. Follow and understand the care protocol and regime
4. Avoid overnight wear unless extended wear lenses
5. Never shower or swim wearing contact lenses
Non-Compliance

Dissenting behaviour amongst contact lens wearers is paramount when considering the main reason for complications. A large well conducted study undertaken on behalf of Bausch & Lomb across Europe highlighted that 98% of all lens wearers were non-compliant in at least one aspect of their lens-care regime (Figure 2).

(1) Hand Hygiene: Although inadequate handwashing before lens handling has been associated with a significant increase in risk of infection, the effect is not instantaneous as it takes weeks to remove micro-organisms embedded on the hands. Perhaps as Morgan suggested, formal training should be provided as this has proven to improve infection control in hospital settings.

(2) Care Regime & Solutions: One in three lens-related complications arise directly from inadequate lens care. Cleaning regimes are either hydrogen peroxide or multipurpose solution based. Multipurpose solution, dubbed as the ‘no rub’ solution is the most widely used. However, rubbing and rinsing is an imperative step as it removes up to 99.9% of bacteria, thereby adding a safety margin of up to 100,000 times. Interestingly, recent studies have demonstrated, hydrogen peroxide based cleaning regimes have superior disinfecting capabilities than using multipurpose solution alone. They reduce the risk of corneal inflammation by ten-fold and disinfects against amoebic cysts. However, for maximal benefit lenses must be exposed to the peroxide solution for a longer time and must be neutralised before wear to avoid ocular toxicity.

(3) Personal Habits: Other unsafe practises include using lenses beyond their recommended replacement schedule, inadequate lens-case care and topping up contaminated solution. The risk increases four-fold compared with appropriately discarded lenses.

Unsupervised wear

Another recent social trend was the use of zero-powered or plano tinted cosmetic lenses designed to change the colour of the eye. They were being bought from unlicensed vendors over the internet without prescription, proper fitting, inadequate information on use, hygiene and complications and no ongoing supervision. Complications associated with the use of such lenses were first reported in 2003. In 2005, further cases reported users sharing lenses between multiple wearers without adequate cleaning. Subsequently, in 2006, Food and Drug Administration (FDA) introduced guidance in the USA, whereby plano lenses could only be purchased under the supervision of a registered practitioner.

Orthokeratology

Orthokeratology is the practice of temporary reduction in myopia by the programmed application of rigid gas-permeable contact lenses, usually at night whilst sleeping.

How to reduce the risk of complications

Education

Patient education, particularly regarding the handling and maintenance of contact lenses, is vital in improving overall

Of late, there has been a resurgence of this phenomenon particularly in East Asia and there are growing concerns about the risk of microbial keratitis and loss of vision. Findings of fifty case studies showed 30% had Acanthamoeba keratitis from nocturnal orthokeratology compared with 5% from regular lens wear.

What are the implications?

Each year 0.02% to 0.04% of lens wearers can lose up to two lines of best correct visual acuity measured using the snellen chart. As well as the risk of losing sight, other significant morbidity associated includes hospital admission and/or intensive treatment, cost of therapy, visiting a health care provider, taking time off from work and inability to wear lenses. An Australian study estimated the median direct costs at Aus $760 [interquartile range $1859] and indirect median costs at Aus $468 [interquartile range $1810]. Not to mention, patients may claim compensation for negligence.

Adapted with permission from from Bausch & Lomb, 2010.
compliance. There is no statistically significant difference between patients receiving both verbal and written instructions and those receiving oral only. However, intense initial education has shown improvements in handwashing.

Morgan and colleagues reported that although, 88% were given lens care information, 23% were unable to recollect seeing any information regarding the risks and complications associated with lens wear. Thus, the practitioner must ensure the patient understands the associated risks, how these are best avoided, as well as early recognition of the signs and symptoms and how to proceed in an emergency.

A degree of non-compliance will always be present despite education. A small study amongst medical students in Malaysia showed that although 88% were aware of complications, only 84% were fully compliant with hygiene and lens-care, and 14% continued use despite experiencing eye symptoms.

To help the practitioners identify individuals with poor compliance Morgan has developed the “Traffic Light Model” (Figure 3). Green behaviour is equated to a fully compliant user whilst the red behaviour user is considered non-compliant. To maximize compliance both verbal and written information should be given and key aspects reinforced during follow-ups. Any literature disseminated should be clearly illustrated with sequential steps. Table 2 highlights some key aspects that should be reinforced.

Public awareness

Bausch & Lomb launched a novel and invigorating online campaign “Eyegiene” to promote the importance of maintaining good eye health and aid compliance. Their website (http://www.thinkleyegiene.com) features a multilingual virtual optician. Patients can further enhance their eye care whilst travelling, “On-The-Go-Flight-Pack” was also introduced. Such programmes help publicise good lens care to a wider audience.

Conclusion

Contact lens-associated complications can range from self-limiting to potentially sight-threatening, yet they are avoidable. The eye has various defense mechanisms to protect itself; however, the presence of contact lenses alters the natural environment increasing the risk of infection. The incidence of adverse effects of contact lens wear can be reduced by promoting good contact lens hygiene and practices.

References

The authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as:
Brugada ECG Sign & Chest Pain Mimicking ST Elevation Myocardial Infarction

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Abstract

Background: Management of patients with the brugada ECG sign who have no previous history of syncope is still negotiable. We present a case of a 57 year-old Caucasian lady who presented to the emergency department with substernal chest pain.

Results: Her past medical history showed that she had two previous episodes of lightheadedness, but no syncope. She had a family history of sudden death secondary to unknown cause in her aunt at the age of 61. Physical exam was unremarkable except for diaphoresis. Electrocardiography (ECG) showed ST elevation in the right precordial leads (V1-V2) with T inversion, mimicking a STEMI. Emergent cardiac catheterization revealed normal coronary arteries. Echocardiogram was normal. Again, interpretation of ECG revealed a Brugada type 1 pattern, characterized by coved-type, gradually descending ST-T segment, elevated J point of more than 2 mm and T wave inversion. Electrophysiological (EPS) testing with a Sodium channel blocker challenge showed a persistent Brugada type 1 pattern with non-inducible ventricular tachycardia. This patient had Brugada type 1 ECG pattern with no previous history of syncope (asymptomatic). Thus she was considered at low risk of developing a serious arrhythmogenic event in the future. Conclusion: A history of syncope remains the best available predictor for arrhythmogenic events. EPS testing in such patients, to stratify the risk and predict for any future events, is still controversial. It is still unjustified to place an implantable cardioverter-defibrillator in asymptomatic non-inducible individuals with the Brugada pattern. These patients should follow up closely with a cardiologist and be aware of the risk of possible triggers of ventricular arrhythmias.

Key Words: Brugada syndrome, sudden cardiac death, myocardial infarction, sodium channels, cardiac electrophysiologic technique (Source: MeSH-NLM)

Introduction

The Brugada pattern is characterized by ECG changes alone, without any of the clinical features of Brugada Syndrome. The ECG changes can be one of three types and can be dynamic. However, the patient should have the type 1 ECG pattern “coved-type” to make the diagnosis of Brugada. The following case report and discussion focuses on the presentation, diagnosis, management and prognosis of the asymptomatic patient with a Brugada ECG pattern.

The Case

A 57-year-old Caucasian lady presented to the emergency department (ED) with substernal chest pain of one day duration. It was associated with headache, backache and mild dyspnea. She reported 2 episodes of lightheadedness in the past month when she felt she was about to pass out. The episodes lasted for a few seconds and were relieved spontaneously. She denied syncope, dizziness, cough, palpitations, nausea or vomiting.

Her past medical history included hypertension, migraine, depression, attention deficit hyperactivity disorder, chronic sinusitis and asthma. Her home medications included lisinopril, gabapentin, trazodone, adderall, celebrex and flonnet diskus. She has history of allergy to codeine sulfate. Her family history included hypertension and coronary artery disease, however it was significant for sudden death of unknown cause in her aunt at the age of 61. She also had a 48-pack-year history of smoking, but quit 7 years ago. She used to work as a hairstylist and consumed alcohol occasionally.

On physical examination, the patient was diaphoretic and in moderate distress in the ED. The ECG was interpreted by the on-call ER physician as a STEMI (Figure 1) and the cardiac catheterization team was activated. Coronary angiography revealed normal coronary arteries. Laboratory results revealed hypokalemia K = 2.9, and otherwise normal CBC, cardiac enzymes and other electrolytes. Transthoracic echocardiography showed a normal left ventricular ejection fraction and no structural heart disease.
Case Report

Figure 1. Electrocardiography of 57 years-old lady showing a Brugada type 1 pattern, characterized by the coved-type, gradually descending ST-T segment elevation, J wave and T inversion in leads V1-V2 and incomplete RBBB in the emergency department.

Figure 2. Electrocardiography of the same lady showed a persistent Brugada type 1 ECG pattern on the 5th day of admission. The patient was afebrile and recovering from a community acquired pneumonia on antibiotics.

The electrophysiology service was consulted and reevaluation of the ECG revealed coved-type J point and ST elevation in the right precordial leads (V1-V2) with incomplete right bundle branch block (RBBB), illustrating Brugada type 1 pattern. Cardiac MRI study showed no evidence of arrhythmogenic right ventricular dysplasia (ARVD). Genetic testing for SCN5A mutation wasn’t performed.

The patient was transferred to telemetry for observation. Electrophysiological study (EPS) was planned for the next day due to a Brugada type 1 pattern on ECG with non-sustained ventricular tachycardia, when the patient was found to have fever (101.2 F), tachycardia, shortness of breath and cough.
Physical examination and chest X-ray showed left lower lobe infiltrate with left pleural effusion. Labs showed leukopenia and a positive urine test for the Streptococcus pneumonia antigen. She was diagnosed with community acquired pneumonia and antibiotics were started.

On the 6th day of admission, the patient was doing well and afebrile. A persistent Brugada type 1 ECG pattern was shown in (Figure 2). EPS was performed with provocative testing using procainamide. It revealed Brugada type 1 ECG pattern but no VT was induced with up to triple ventricular extra stimuli from right ventricular apex and base, on and off isuprel. The patient was asymptomatic and no implantable cardioverter defibrillator (ICD) was implanted. She was discharged home and scheduled to follow up closely with her cardiologist.

Discussion
Sudden cardiac death (SCD) in patients with normal hearts is an uncommon occurrence.1 Brugada syndrome (BrS) was described in 1992 as one of the causes of SCD in patients with normal heart structure.2,3 This syndrome of high incidence of life-threatening ventricular arrhythmias is associated with ECG changes consistent with ST segment elevation in leads V1 to V3 and incomplete right bundle branch block.4-6 Due to the availability of a preventable mode of therapy, an implantable cardioverter-defibrillator (ICD) should be considered when definitive diagnosis of BrS is made.7

The Brugada pattern is characterized solely by the typical Brugada ECG changes, excluding the following clinical features: documented ventricular fibrillation (VF), self-terminating polymorphic ventricular tachycardia (VT), family history of SCD at the age <45 years, type 1 ST segment elevation in family members, electrophysiologic inducibility of VT, unexplained syncope suggestive of tachyarrhythmia or nocturnal agonal respiration. The Brugada ECG pattern has a coved-type, gradually descending ST segment, J wave amplitude of more than 2mm and T wave inversion.5,9 Characteristics of patients with the Brugada ECG pattern included higher prevalence in the Asian population.5

However, the exact prevalence varied among studies from different countries.11,14-18,21 It was mainly observed in adult males11,20 and a mutation in SCN5A gene on chromosome 3p21-24, that codes for the alpha-subunit of cardiac sodium channels, was found in about 18-30% of families with BrS.10,11,15,21

The relation between the Brugada ECG sign in asymptomatic patients (no previous history of syncope) and the future risk of developing arrhythmogenic events has been investigated, especially over the past decade. One study revealed that the cardiac event rate per year in asymptomatic patients was 0.5%, compared to 1.9% in patients with a history of syncope and 7.7% in patients with aborted SCD.19 Further reports of extended follow up of the asymptomatic population over a period of 3 to 7 years showed none to a very low arrhythmogenic event rate.20-22 Variations in the baseline characteristics of asymptomatic patients over a mean follow up of 2 to 3 years, revealed different rates of arrhythmias, with events arising in 8% of the asymptomatic population studied by Brugada et al.,20-22 compared to 0.8% in a later study by Eckardt et al.23

Multiple predictors of future arrhythmic events in patients with the Brugada pattern were studied, with male gender, mutation of the SCN5A gene and a positive family history of SCD found to be non-predictive.24-26 The prognostic value of the clinical, ECG and EPS variables was analyzed in a population of spontaneous type 1 Brugada ECG patterns and no previous cardiac arrest. The cohort with negative EPS (non-inducibility of VT/VF) had a 1.8% risk of developing arrhythmic events (SCD or documented VF), compared to 14% in those with a positive EPS.27 Further studies revealed a 0.9% rate of significant cardiac events in patients with non-inducible arrhythmias.28-30 A recent prospective multicenter study showed that a positive EPS is not predictive for arrhythmic events, and data reported that the spontaneous ECG pattern and the history of syncope are the best available predictors of such events.31-33 It also showed that QRS fragmentation and a ventricular effective refractory period of less than 200ms are risk indicators. These findings are imperative indicators for the need of an implantable cardioverter defibrillator in such populations.34-36 Additional literature review revealed the low significance of EPS in stratifying the risk of future arrhythmic events, and that its role remains an area of investigation and debate.37,38,39,40

Currently, pharmacological therapy for the prevention of SCD in Brugada patients is not well established, however some reports showed that quinidine and hydroquinidine may be beneficial in such patients.41-43 The ICD has multiple complications, including the relatively high rate of inappropriate shocks (6-9% per year), thus it seems to be unjustified to give an ICD to the asymptomatic non-inducible individual with an abnormal ECG pattern.44-46 There is no definite evidence to suggest that individuals who have no personal history of syncope or any family history of SCD are at a higher risk of SCD than the general population.47,48,49,50

In our case, genetic testing was deferred as only a small percentage of patients with the BrS do test positive for the SCN5A gene mutation,51-53 and recent reports showed that it will not guide the management of such cases.54 We excluded other important causes of SCD, including ARVD, where the Brugada ECG pattern can be an early subclinical manifestation of this genetic disorder.55 Moreover, our patient was afebrile on presentation and didn’t have any of the known provoking factors that induce a Brugada-like
ECG pattern. There was no previous history of syncope along with non-inducible VT on provocative EPS testing. Lack of such possible predictive factors reflects a low risk of developing arrhythmic events and SCD. Thus ICD therapy wasn’t recommended in this lady as the weight of evidence didn’t support it. To conclude, our patient was advised to follow up carefully with the cardiologist and to be aware of the risks of possible triggers of ventricular arrhythmias.

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Acknowledgements
None.

Conflict of Interest Statement & Funding
The Authors have no funding, financial relationships or conflicts of interest to disclose.

A Case of Hyperbaric Oxygen as Adjunct to Fasciotomies in Compartment Syndrome, Ischemia-Reperfusion Injury, and Delayed Secondary Infection

Dawnielle C. Endly, 1 Joan V. Eggert 2

Abstract

Background: Fasciotomies are a therapeutic treatment for compartment syndrome, but they also allow reperfusion to tissues that have been hypoxic. We report a case study of a 52-year-old male with an ischemic leg. Despite prophylactic fasciotomies, ischemia-reperfusion injury resulted in delayed myonecrosis and progressive necrotizing fasciitis. Results: After two hyperbaric oxygen treatments, edema was markedly reduced and all visible wound tissues were well perfused without evidence of ascending infection, allowing the patient more time for further evaluation. The patient did opt for an above the knee amputation in a non-emergent setting and now successfully utilizes a prosthesis for ambulation. Conclusion: Ischemia-reperfusion injury may result in delayed tissue loss in spite of appropriate fasciotomies. Hyperbaric oxygen may be a useful adjunct therapy even when initiated days after the initial injury.

Key Words: Reperfusion Injury, Hyperbaric Oxygenation, Surgical Decompression. (Source: MeSH-NLM).

Introduction

Acute compartment syndrome of an extremity often occurs after a traumatic injury. As pressure builds within the compartment, the arterioles collapse rendering the tissue hypoxic. 1 A fasciotomy can be limb saving as it allows decompression of the involved compartments. 2 Paradoxically, however, the decompression allows return of blood flow which can result in further tissue damage known as ischemia-reperfusion injury. Hyperbaric oxygen therapy (HBOT) is useful as an adjunct to fasciotomies where it is likely that ischemia-reperfusion injury will result. 3 We present a case where HBOT was utilized eight days after a traumatic injury when progressive myonecrosis and necrotizing fasciitis occurred secondary to ischemia-reperfusion injury.

The Case

Eight days after a high speed motorcycle accident, a healthy 52-year-old male had deteriorating signs of myonecrosis and evidence of a rapidly ascending necrotizing fasciitis of the left lower extremity. He had been revascularized after transection of the proximal popliteal artery approximately seven hours after injury with prophylactic fasciotomies performed at time of initial revascularization. Extensive irrigation and debridement was performed on three occasions due to the wound being highly contaminated and the presence of leukocytosis of 13.5 K/ul on day one. He was placed on Penicillin and Gentamicin prophylactically the day of the injury.

Five days after injury, negative pressure wound therapy (V.A.C® Therapy, KCI, San Antonio, TX) was utilized to control the drainage and Piperacillin-tazobactam was added for continued fevers and a white blood cell (WBC) count that began increasing again. He was extubated on day six, but it became evident that he could not move the foot on command. On day seven, an infectious disease (ID) consult was ordered as the patient had continued fever, WBC count of 14.0 K/ul, a foul odor coming from the leg, and dirty dishwater like secretions exceeding the wound V.A.C. canister capacity. ID recommended adding Linezolid and Ciprofloxacin given the clinical course and wound cultures grew Escherichia coli, Pseudomonas aeruginosa, and Stenotrophomonas maltophilia. By day eight, the WBCs increased to 18.8 K/ul and tissue continued to devitalize and dissolve causing a clinical suspicion of a life threatening necrotizing fasciitis and the surgeon adamantly advised an above knee amputation.

The patient had an altered mental status and refused to consent for amputation. The surgeon requested a psychiatric consult as family members weren’t comfortable with consenting for amputation and requested that he be transported to his home state for a second opinion before deciding that the leg was non-viable.
HBOT was initiated as an alternative to immediate amputation. The patient received two hyperbaric oxygen treatments at 2.8 atmospheres absolute (ATA) and 2.4 ATA respectively within 12 hours in an attempt to slow the myonecrosis and ascending infection. After HBOT, all visible wound tissues were pink, well perfused, and viable (Figures 1a and 1b). Edema was reduced and there was no evidence of a progressive ascending infection so he was immediately transferred by air to the care of a trauma surgeon at a specialty hospital closer to his home. After further assessment, it was determined the sciatic nerve was completely severed above the knee, and the leg would be incapable of regaining motor activity. The patient did opt for an above the knee amputation in a non-emergent setting with family support and full informed consent. The limb healed well after amputation and he now successfully utilizes a prosthesis for ambulation.

Discussion

Fasciotomies have long been the standard of care in patients suffering from acute compartment syndrome. However, this case illustrates that resulting ischemia-reperfusion injury may cause delayed tissue necrosis in spite of appropriate fasciotomies and HBOT can be utilized as a viable therapeutic adjunct. The mechanisms behind ischemia-reperfusion injury begin with cellular hypoxia which creates a shift to anaerobic metabolism and accumulation of anaerobic byproducts. Fasciotomies re-introduce oxygen into this distorted environment. As a result, free radical formation increases, inflammatory mediators aggregate, the mitochondria sustain structural damage, the sodium-calcium pumps are disrupted, and the complement system is activated.  

In addition, endothelial cells increase the expression of adhesion molecules, such that re-perfusion increases neutrophil attachment. The resulting release of reactive oxygen species from bound neutrophils activates complement and ultimately induces tissue necrosis.  

While HBOT does not replace a fasciotomy, it has several mechanisms of action that are useful to compromised and contaminated tissue. First, the hyperoxia improves oxygen delivery to tissues in need and prevents infections due to anaerobic microorganisms. Leukocytes also utilize hyperoxia for the neutrophil’s oxygen dependent bacterial killing. Additional oxygen in reperfused tissues results in the generation of free radical scavengers such as superoxide dismutase to neutralize the waste products of anaerobic metabolism. Moreover, hyperbaric oxygen inhibits the diapedesis of neutrophils on post-capillary venules by antagonizing the Beta 2 integrin system. Overall, HBOT maintains oxygenation of the tissues that have become compromised, reduces risk of infection, and down-regulates the inflammatory response.

Ideally HBOT is started within hours of injury, but this requires the patient to be stable enough to tolerate the treatment. Most hyperbaric centers in the United States, including ours, primarily treat stable outpatients, not intubated and critically ill patients. Thus, we could not have treated this patient before he was extubated. In this case HBOT was utilized to treat a secondary progressive myonecrosis and necrotizing fasciitis to allow the patient and family time to make an educated decision on moving forward with amputation of the limb. Further discussion about the use of HBOT as an adjunct to fasciotomies is needed before conclusions regarding the expected clinical course and outcome can be made.
Case Report

References

Acknowledgements
None.

Conflict of Interest Statement & Funding
The Authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as:
“How Can I Help You?”: Perspectives from a Patient with a Hearing Loss

Rachel Kolb

The Experience

“So, how are you feeling today?” I retrieve the question off the doctor’s lips and compose myself before launching into an explanation. She is a stranger to me, and though she can easily see on my medical records that I am deaf, I wonder if she knows what that entails for our conversation. I wonder if she knows that, for me, deafness is not just a medical condition, a malady to be treated with prescriptions and therapeutic remedies. It is a dominant force that shapes the way I communicate, the way I understand the world. It has influenced my life experiences as much as any physical characteristic can, even while in another sense it is no different than, say, wearing contact lenses. I see through the lens of my deafness, and I use a particular set of strategies to surmount any barriers it sets before me, even while it is a deep and inescapable part of who I am.

I also wonder if my doctor knows that, despite the “profound hearing loss” designation that must dominate my medical file, I come in today just as anyone else would, wanting to have a direct conversation about my body and my condition. I am sure she does, since she is treating me as she would any other patient. She listens, speaks normally, and looks down at her clipboard from time to time. I appreciate this; I am simply myself, and I am here for a routine physical. Yet difficulties seep in when she treats me like she would treat any other person. This is the paradoxical reality of my life: in many ways I feel and function just like anyone else, but I must often behave differently if I am to communicate on an equal plane.

My doctor continues looking down to write notes as I speak, but sometimes she throws me off balance by saying something when she is not looking at me. Sometimes, later in the examination, she moves around the room to where I cannot see her as clearly, often to retrieve a new implement, but does not think to stop talking. The part of the physical when she stands behind me with a stethoscope and asks me to breathe deeply never works too well. My strategy, in the lack of any visual cues, is to take deep breaths at random and hope they give her some useful information. In dentist’s offices or other medical situations, she places a surgical mask over her face, cutting me off from communication entirely.

There is an easy solution for this: I can politely interrupt what she is doing and say, “I’m reading your lips. Could you please slow down, look at me, and speak more clearly?” In most cases, this works well. She adjusts her mannerisms and I breathe a sigh of relief as the words flow off her clear, articulate lips. Still, lipreading is less of a fix-all solution than it initially appears, for it is not quite like reading the words in a book. Years of speech therapy and immersion in the hearing world have taught me to work with the facial features of most people I meet, to contrive the appearance of functioning like a hearing person myself, but work is the key word in this statement: lipreading is incredibly hard work.

When I lipread, in the medical setting or elsewhere, I settle down and fixate on my companion’s face, observing the minute ways its muscles move. Some people’s features are more naturally expressive than others, and their words reach my eyes more clearly. What to do with the others? The men who have facial hair that obstructs what they are saying, the people who mumble or speak quickly or who have thin lips that hardly move at all? What about the people whose accents are foreign to me, whether they hail from Boston or somewhere farther away like Scotland or Singapore, and whose way of articulating their words clashes with my mental model of what speech looks like? When I meet people whose physiques challenge my ability to adapt, I resort to guesswork. While still attempting to pay attention to what they are saying in real time, I process what they have just said, what it could have meant according to the context and according to what I’m seeing right now. But guesswork doesn’t only figure into conversations with people whose faces make them hard to lipread. It is an inherent part of lipreading itself.
Experience

When I lipread, I enter the difficult territory of trying to use one sense to retrieve the information that was meant for another. My sight informs me of what my hearing cannot grasp, but spoken words were not meant for sight alone. I have read that only thirty percent of spoken English is clearly discernable from lipreading alone.4 Beyond that threshold, whatever accuracy I achieve is due to guesswork, context, determining whether that word I saw contained a “b” or a “p,” a “t” or a “d.” (Their only difference: one is voiced and one is not.) I use information I know, or preexisting clues I’ve been given, to try to figure out what I did not catch. But how am I supposed to guess what word I’m seeing when my doctor is using medical terminology I do not know, or when she prescribes a medication I have never heard of before and cannot even pronounce?

Unfamiliar terminology surfaces in all situations, making lipreading more difficult, but in the medical setting I feel heightened pressure to get it right. The conversations I have with my doctor are not like the small talk I might have at a café. When I visit her office, I enter an intimate conversation about my body and its condition; I yearn to grasp her insights. My failure to communicate well because I am deaf is not only potentially hazardous for my health. It also fosters a mindset in which I feel far less empowered to look after my own physical well-being. Or, to ask a broader question: if I cannot understand a doctor’s insights about my body, what happens to my sense of autonomy and control over my life?

One might ask, at this point, why I do not use a sign language interpreter to help me function better in the medical setting. At one point, I did. When I was a child, my mother always came along to my doctor’s appointments to be my interpreter. But even with her support, learning to communicate for myself was paramount: I longed after the independence that comes with direct communication. To take my communication into my own hands, although I know the option is there. I prefer to communicate well on my own. My speech and lipreading skills enable me to communicate well one-on-one without a professional interpreter, although I know the option is there. I prefer to keep my communication into my own hands, although I realize that this is a matter of personal choice.

Some other deaf and hard-of-hearing people do not (or cannot) make this choice – and, in any case, when I or anyone else speaks of “hearing loss,” we speak of a dizzying spectrum of experiences and abilities, from the deaf to the hard-of-hearing, from those born with their hearing loss to those whose hearing fades later in life. Nearly one in eight Americans has a bilateral hearing loss, and among people aged 70 and older the number rises to three in five.4 Some of those individuals sign; others might prefer to lipread. Some might strongly identify as culturally Deaf; others might view themselves as hearing people who have difficulty communicating in certain situations. Many people with hearing loss lipread to some extent, though when one’s deafness is profound enough, using sign language is often a far clearer mode of communication. The diversity among people with hearing loss is staggering, making flexibility in the medical setting all the more crucial.

When I was young, I relied heavily on sign language support, and the medical system could sometimes seem ill-equipped to accommodate this need. When I was in first grade, my family left a reputable local pediatric dentist because her office would not allow parents to be present during medical procedures – even for a child with a disability. My mother explained my situation: I would be sitting in the dentist’s chair unable to understand who was poking around in my mouth and what she was about to do to me. Couldn’t the office make an exception and allow her to interpret? When they refused, we went elsewhere. I have heard about other deaf people who have also struggled with medical professionals’ seeming unwillingness to provide full communication access. Refusing to write things down or to provide an interpreter is only the worst of it. Other behaviors, like failing to maintain eye contact or treating the patient as invisible while conversing with the interpreter instead, can feel nearly as demeaning. With or without interpreters, and medical procedure notwithstanding, visiting a doctor’s office needs to be something that is done for me, not to me.

Sometimes, it is not even my conversation with the doctor herself that poses the biggest challenge in my medical visit. I arrive in the office and check in at the receptionist’s desk, with someone who is usually sitting in front of a computer who is usually paying more attention to the screen than she is to me. Her words can be fleeting and difficult to understand, but fortunately experience has taught me what will happen: I will say my name and the name of the doctor I am here to see and when, and she will thank me and tell me. She needs to be something that is done for me, not to me.

While in the waiting room, I take care to position myself where I can see the nurse when she comes out and calls my name. I do the same in airports and other public places: even when I ask the people staffing the front desk to alert me when something important happens, more often than not they forget. Sometimes I pick up a magazine to pass the time, but always my eye lingers on the doorway. The process of calling patients back to the exam room is an auditory rather than a visual one, and more often than not I do not catch my name the first time the nurse says it, or erroneously respond when she has called someone else.
When I reach the exam room, the nurse, even though I have told her I read lips, sometimes turns to her notepad or computer and asks me a rapid succession of questions – Your height? Your weight? Have your medications changed? – without bothering to look at me. Then she catches me off-guard by asking for my arm to test my blood pressure, or asking me to sit up so she can listen to my heart. I have had this happen even in otolaryngology and audiology clinics, where hearing loss is prevalent. Often the situation improves when I stop the nurse and explain my situation, but already before I meet my doctor I feel embarrassed, worried about my ability to understand, and generally ill at ease. Even for hearing patients, for whom communication is not such a barrier, must it not feel disorienting to be treated in this brisk, laboratory-object manner? Slow down, I want to say. Look into my face and let me tell you what I need. The doctors who do this – the ones who walk into the room and pay instant attention to my needs, who take care to maintain the connection, who ask how they can make communication easier – are the best ones. I have been fortunate enough to have many doctors like this. Even if I have never met a doctor who knew sign language, their good communication skills can help surpass the language barrier that tends to isolate people with hearing loss. And, even since I received a cochlear implant several years ago, which has made lipreading easier, I continue to believe that the doctors who exhibit such keen awareness of their patients’ overarching needs tend to have excellent medical insights as well. After all, a good doctor ought not to be simply an ivory tower of knowledge, to whom patients must make a pilgrimage. He or she ought to be someone willing, humble, and flexible enough to ask those patients, quite clearly and directly, How can I help you?

References

Acknowledgements
Some parts adapted from “Seeing at the Speed of Sound,” Stanford magazine, March/April 2013, p.50-56.

Conflict of Interest Statement & Funding
The authors have no funding, financial relationships or conflicts of interest to disclose.

Cite as: