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Microbiological Profile of Corneal Ulcers at a Tertiary Referral Center

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ABSTRACT

The aim of this study was to describe patient demographics, microbiological profile, and antibiotic susceptibility of corneal ulcer at a tertiary referral center to improve and optimize diagnosis and treatment of this potentially blinding entity and to reduce antibiotic misuse. Detailed external and slit-lamp bio-microscopic examination of 123 consecutive patients with suspected corneal ulcer was performed at an ophthalmology clinic. Corneal scraping was carried out under slit-lamp bio-microscopy. The obtained material was inoculated on culture media and smeared on a slide for Gram's staining for morphological identification of bacteria and fungus. For samples that developed colony in culture media, antibiotic susceptibility testing was performed. In a significant percentage of patients (72%) neither bacterial agents nor fungi were the cause of corneal ulcer. Of the 34 culture-proven corneal ulcers, in 79% of the cases, bacteria were detected while in 21% of cases, fungi were found. Of the 27 bacterial corneal ulcers, the majority were (67%) caused by Gram-positive bacteria, of which 50% were Streptococcus pneumoniae, and in the Gram-negative bacterial corneal ulcers, most of the cases (44%) were caused by Pseudomonas aeruginosa. In the antibiotic susceptibility report, Streptococcus pneumoniae, Staphylococcus aureus, Pseudomonas aeruginosa, and Escherichia coli were resistant to Cotrimoxazole (TS), Streptococcus pneumoniae to Erythromycin (E), Staphylococcus aureus to Peniciline (PG), Pseudomonas aeruginosa to Ceftriaxone (CRO) and Nitrofurantoin (NI), and finally, Escherichia coli to Gentamicin (GM). In conclusion, in a significant number of the patients neither bacterial agents nor fungi were offending microorganisms and bacteria were the most common agent of microbiological corneal ulcer, found in 79% of culture-proven corneal ulcers, followed by fungus, found in 21% of culture-proven corneal ulcers.

KEYWORDS

Microbiological Profile; Antibiotic Susceptibility; Corneal Ulcer; Tertiary referral center

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INTRODUCTION

A "silent epidemic" is a term that refers to corneal ulceration in developing countries and its incidence is approximately ten times that of the United States [1], which annually imposes a heavy financial burden on the health-care system, even in developed countries [2]. Even though the most common offending agent is Herpes Simplex Virus type 1, other common infectious agents in this potentially blinding condition are bacteria, including Streptococcus pneumonia, Staphylococcus aureus, coagulase-negative staphylococci, and Pseudomonas aeruginosa; fungus including Candida albicans, Aspergillus flavus, Fusarium solani, Penicillium species and Aspergillus fumigates as well as parasites [3, 4]. Environmental and geographical factors can influence the pattern of infection and there are numerous differences in the profile of corneal ulcers in various regions [5].

Although effective treatment with fortified antibiotics based on the results of corneal smear and culture is necessary, in the management of this sight-threatening condition, intensive treatment with several topical broad-spectrum fortified antibiotics is used while culture results are pending [6]. However, the emergence of a wide variety of antibiotic resistance has been reported in the recent decades, such as fluoroquinolones resistance in case of culture-proven corneal ulcer with prior use of this antibiotic or moxifloxacin resistance in corneal ulcer caused by *Pseudomonas aeruginosa* [6-10]. These findings highlight the necessity of conducting epidemiological studies to examine antibiotic susceptibility in different geographical areas so that a cost-effective approach can be designed for the management of this infective condition. Meanwhile, such studies will help the selection of effective antibiotics for optimal treatment of corneal ulcer in that specific geographical area based on previous studies conducted in this area.

The aim of this study was to describe patient demographics, microbiological profile, and antibiotic susceptibility of corneal ulcer at a tertiary referral center to improve and optimize diagnosis and treatment of this potentially blinding entity and also to reduce antibiotic misuse as well as antibiotic resistance of the microbes.

METHODS

This retrospective cross-sectional study was conducted between 23rd of September 2017 to 23rd of September 2018 at the pathology laboratory of Shaheed Labbafinejad Hospital, Shaheed Beheshti University of Medical Sciences, Tehran, Iran. The study obtained ethical approval from the Ethical Committee of the

Shahid Beheshti University of Medical Sciences (the research ethic's code is: IR.SBMU.RETECH.REC.1397.863). Following the Declaration of Helsinki, the samples were collected as a routine practice of ophthalmology clinic for suspected infectious corneal ulcers.

Detailed external and slit-lamp bio-microscopic examination of 123 consecutive patients with suspected corneal ulcer was performed by a resident of ophthalmology. Corneal scraping was carried out under slit-lamp magnification, following instillation of one drop of 0.5% tetracaine hydrochloride (Anestocaine, Sinadaru, Tehran, Iran) in an aseptic condition, from the leading edge and base of the ulcer, using a sterile No. 15 Bard-Parker.

The obtained material was inoculated on culture media, including MacConkey agar (Pronadisa Lab. Conda, Madrid, Spain), Blood Agar Pronadisa medium (Pronadisa Lab. Conda, Madrid, Spain), chocolate agar (Pronadisa Lab. Conda, Madrid, Spain), and Sabouraud dextrose agar (Pronadisa Lab. Conda, Madrid, Spain) for culture; the methods were based on instructions of the manual of American Society for Microbiology [11] and smear on the slide for Gram's stain for morphological identification of bacteria and fungus.

For samples that developed colonies in culture media, antibiotic susceptibility testing was performed using the following Mast Discs (Mast Group, Merseyside, UK): cefazolin (30 μg) - CZ30; streptomycin (10 μg) - S10; penicillin G (10 μg) - PG10; novobiocin (5 μg) - NO5; Cefepime (30 μg) - CPM; erythromycin (15 μg) - E15; nitrofurantoin (200 μg) - NI200; cefoxitin (30 μg) -FOX30; cotrimoxazole (Trimethoprim/sulfamethoxazole 1.25/23.75 μg) - TS 25C; Amikacin (30 μg) - AK30; Ceftazidime (5 μg) - CAZ5; ceftriaxone (30 μg) - CRO30; Meropenem (10 μg) - MEM10; Ampicillin (25 μg) - AP25; Imipenem (10 μ g) - IMI10; vancomycin (30 μ g) - VA30; gentamicin (10 μg), - GM10; and ciprofloxacin (1 μg) -CIP1C. An antibiotic disc was selected based on Gram staining of a developed colony in culture media and specific features of colonies showing the potential microorganism. In the microbiological report of this test, the sensitivity or resistance to the antibiotic was reported for each sample.

The obtained data was entered in Microsoft Office Excel, 2016 and Statistical Package for Social Science software (ver. 22.0, SPSS, Inc., Chicago, IL) was used for analysis of data. Statistical analysis included both descriptive and analytical statistics. Frequency, percentage, mean, and standard deviation (SD) for data were reported. In analytical statistics, P <0.05 was considered significant.



RESULTS

Overall, the mean \pm SD age of 123 enrolled patients in this study was 53.88 \pm 16.76 years old, of which 68 were men and 55 were women, with a mean \pm SD age of 58.21 \pm 15.03 years and 48.53 \pm 17.37 years, respectively, where women were significantly younger than men (P < 0.001).

In total, among 89 (72%) cases of corneal ulcers, no microorganisms in Gram stain and culture media were detected, and in the remaining 34 patients, who had culture-proven infectious corneal ulcer (Table 1), the following microorganisms were identified in order of frequency: Streptococcus pneumoniae in nine patients (six men and three women), Staphylococcus aureus in seven patients (seven women), Pseudomonas aeruginosa and Candida albicans; each in four patients (Pseudomonas aeruginosa; three men and one woman and Candida albicans; one man and three women), Escherichia coli in three patients (two men and one

woman), Aspergillus flavus in two patients (two men), and Acinetobacter spp, Alternaria spp, Klebsiella pneumoniae, alpha-hemolytic Streptococcus and Streptococcus agalactiae, each detected in one case (one woman, one man, one woman, one woman, and one man, respectively). According to the Gram staining results, out of 123 smears, eighteen Gram-positive and nine Gram-negative bacteria were detected. In order of frequency, out of 34 infectious corneal ulcers, 18 (53%) were caused by Gram-positive bacteria, nine (26%) by Gram-negative bacteria, and seven (21%) by fungus.

The results of antibiotic susceptibility testing for 27 bacterial keratitis out of 34 infectious corneal ulcers is shown in Table 2. These data were collected from the microbiology laboratory notebook, and no extra information other than the name of resistant or sensitive antibiotic was found.

Table 1: Microbiological Results for Culture-proven Infectious Corneal Ulcers (n = 34)

Organism	Number
Gram –positive bacteria	
Streptococcus pneumoniae	9
Staphylococcus aureus	7
alpha-hemolytic streptococcus	1
Streptococcus agalactiae	1
Total	18
Gram –negative bacteria	
Pseudomonas aeruginosa	4
Escherichia coli	3
Klebsiella pneumoniae	1
Acinetobacter spp	1
Total	9
Fungus	
Candida albicans	4
Aspergillus flavus	2
Alternaria spp	1
Total	7

Table 2: Result of Antibiotic Susceptibility testing for 27 Culture-Proven Bacterial Corneal Ulcers

Organism	Resistant	Sensitive
Streptococcus pneumoniae	TS, E	VA, CRO, MEM, S
Staphylococcus aureus	TS, PG	VA, CZ, NO, FOX
Pseudomonas aeruginosa	TS, CRO, NI	AK, GM, CIP, IMI,CPM, CAZ
Escherichia coli	TS, GM	CRO, CIP, CAZ, MEM, IMI, AK
Acinetobacter spp	-	AK, CAZ, CRO, GM, IMI, MEM
Klebsiella pneumoniae	-	AK, CAZ, CRO, CIP, GM, IMI
Alpha-hemolytic streptococcus	-	VA, E, CRO, MEM
Streptococcus agalactiae	-	AP, IMI, VA, GM, CIP

PG: Peniciline; S: Streptomycin; CZ: Cefazolin; NO: Novobiocin; CPM: Cefepime; E: Erythromycin; NI: Nitrofurantoin; FOX: Cefoxitin; TS: Cotrimoxazole; AK: Amikacin; CAZ: Ceftazidime; CRO: Ceftriaxone; MEM: Meropenem; AP: Ampicillin; IMI: Imipenem; VA: Vancomycin; GM: Gentamicin; CIP: Ciprofloxacin.



DISCUSSION

In the present study, the majority of patients were male, who were significantly older than females, and in a significant percentage of patients (72%), neither bacterial agents nor fungi were the cause of corneal ulcer. Of the culture-proven corneal ulcers, microorganisms in 79% of the cases were bacteria and in 21% of the case, these were fungi. Of the 27 bacterial corneal ulcers, the majority (67%) were caused by Grampositive bacteria, of which 50% were Streptococcus pneumoniae, and in the Gram-negative bacterial corneal ulcers, most of the cases (44%) were caused by Pseudomonas aeruginosa. Among seven cases of fungal corneal ulcers, Candida albicans was detected in four patients (57%), Aspergillus flavus in two patients (29%) and Alternaria spp in one patient (14%) as the offending fungus. In antibiotic susceptibility report, Streptococcus pneumoniae, Staphylococcus aureus, Pseudomonas aeruginosa, and Escherichia coli were resistant to TS, Streptococcus pneumoniae to E, Staphylococcus aureus to PG, Pseudomonas aeruginosa to CRO and NI, and finally, Escherichia coli to GM.

In the current study, culture proven microbial corneal ulcers were detected in 28% of corneal scrapings. This was lower than reports from Nepal [12], Ghana [13], South India [14], and Bangladesh [15], which was 45.5%, 57.3%, 70.6%, and 81.7%, respectively.

In the present study, bacteria were the most common agents of microbiological corneal ulcer in 79% of the culture-proven corneal ulcers and 22% of all patients with clinical diagnosis of corneal ulcer, followed by fungus in 21% of culture-proven corneal ulcers and 6% of all clinical diagnoses of corneal ulcer. In studies by Tewari et. al [16] and Suwal et. al. [4] similar results were found. In Tewari's study, bacterial agents caused 65% of the culture-proven corneal ulcers, and the fungus was the offending microorganism in 35% of the cases, and similar to the present study, Gram-positive cocci were the most common bacteria, and among the Gram-negative bacteria, Pseudomonas aeruginosa was the most common. Unlike the present study, however, among the gram-positive bacteria, Staphylococcus aureus and among fungal causes, Aspergillus spp. followed by Fusarium spp. were the most common, which could be due to differences in risk factors and geographical distribution among the two studies. However, because of the retrospective nature of the current study design, the risk factors were not recorded, thus, the causes of these differences are not certain. In Suwal's study, bacteria were detected in the majority of the culture-proven corneal ulcers, amongst which Streptococcus pneumoniae was the most common bacteria, as in the present study. Although the fungus was the second common agent of microbiological corneal ulcers, similar to the present study, the most common fungi were Fusarium species followed by Aspergillus flavus, which differs from the findings of the current study; the most likely justification for this difference is that mentioned previously.

The emergence of antibiotic resistance is a major public health issue and the distribution of microorganisms causing corneal ulcers, especially those which are resistant to antibiotics, varies according to time, geographical location, and hospital [17]. In the study of Mohammadpour et al. [18] on antibiotic susceptibility patterns in pseudomonas corneal ulcers, this microbe was susceptible to CAZ, CIP and AK, as was in the current study. Although the current study was conducted at a different tertiary referral hospital, the study city and the population studied were the same in both studies, which could explain the similarity of the results. In another study conducted in Iran [19] at a tertiary referral center, all Pseudomonas aeruginosa, such as the current study, were susceptible to CAZ and AK. In that study, only about 7% were resistant to CIP, whereas, in the present study, all Pseudomonas aeruginosa were resistant to TS, CRO, and NI. Also, their study results showed that Streptococcus pneumoniae were resistant to AK in 2.6% and in all cases sensitive to CZ, CAZ, cefixime, and cephalothin. However, in the current study, all Streptococcus pneumoniae were resistant to TS and E and sensitive to VA, CRO, MEM, and S.

The study design, recruitment of subjects, and its short duration can be cited as the limitations of this study. Due to the retrospective nature of this study, important information, which could be considered as possible risk factors of corneal ulcer, such as socioeconomic status of patients, history of contact lens wear, and occupational exposure, besides consequent complications of corneal ulcer was not available in the patients' records. Likewise, the choice of antibiotic discs to detect antibiotic sensitivity was out of the preference of the authors of this paper, and the authors only recorded the information in the microbiology laboratory notebook. Therefore, in order to make thoughtful decisions to reduce the risk of corneal ulcer and manage this potentially blinding condition, the need for stronger study design and a longer follow-up period is considered necessary besides examining predisposing factors, such as occupational exposure, history of contact lens wear, and socioeconomic level of patients and vigilance



selection of antibiotic discs for antibiotic sensitivity testing.

CONCLUSIONS

Overall, in a significant percentage of patients neither agents nor fungi were offending microorganisms and bacteria were the most common agent of microbiological corneal ulcer in 79% of the culture-proven corneal ulcers, followed by fungus in 21% of the culture-proven corneal ulcers. In antibiotic susceptibility Streptococcus report, pneumoniae, Staphylococcus aureus, Pseudomonas aeruginosa, and Escherichia coli were resistant to TS, Streptococcus pneumoniae to E, Staphylococcus aureus to PG, Pseudomonas aeruginosa to CRO and NI, and finally, Escherichia coli to GM.

DISCLOSURE

Ethical issues have been completely observed by the authors. All named authors meet the International Committee of Medical Journal Editors (ICMJE) criteria for authorship of this manuscript, take responsibility for the integrity of the work as a whole, and have given final approval for the version to be published. No conflict of interest has been presented.

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