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A STUDY OF DIABETIC FOOT ULCER WITH MULTIDRUG-RESISTANT ORGANISM INFECTION:

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Abstract:

Due to the increased resistance rate of *Escherichia coli* and *Pseudomonas aeruginosa* among gram-negative bacteria, the risk factors of drug-resistant negative bacilli in patients should be evaluated in the development of initial anti-infective treatment regimen in clinical practice. In this study, multidrug-resistant bacteria accounted from the strains obtained by wound cultures, which may be related to the patients with serious illness admitted to our hospital as a tertiary care hospital and the more complex history of antibacterial drug use. This study puts in a sincere effort to find the risk of diabetic foot ulcer with multidrug-resistant organism infection.

Keywords: Drug, Resistance, Foot Ulcer.

Introduction:

Earlier studies reported that Gram-positive bacteria such as Staphylococcus aureus and Enterococcus faecalis were considered as the most common flora of diabetic foot infections, followed by Gram-negative bacteria such as Escherichia coli and Pseudomonas aeruginosa [1, 2]. In recent years, the bacterial spectrum of diabetic foot infections has shifted considerably with the evolution of diabetes epidemiology and changes in the application of antimicrobial drugs [3, 4]. Staphylococcus aureus according to a report is the most common bacterium among all aerobic bacteria, which was consistent with the bacteriological characteristics of skin and soft tissue infections [5]. In a study the analysis of bacterial resistance, it was found that Staphylococcus aureus was the main multi-drug resistant bacteria in diabetic foot ulcer infection, and the resistance rate of Escherichia coli was the highest among gram-negative bacteria, followed by Pseudomonas aeruginosa, [6]. Infection with multidrug-resistant bacteria in diabetic foot ulcers increases the difficulty of treatment by reducing the clinical effect of antibiotics therapy, leading to amputations or deaths [5, 7, 8, increased resistance rate of Escherichia coli and Pseudomonas aeruginosa among gram-negative bacteria, the risk factors of drug-resistant negative bacilli in patients should be evaluated in the development of initial anti-infective treatment regimen in clinical practice. This study puts in a sincere effort to find the risk of diabetic foot ulcer with multidrug-resistant organism infection.

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Aims and Objectives:

To study the diabetic foot ulcer with multidrug-resistant organism infection.

Materials and Methods:

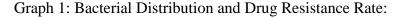
This study included 130 patients with diabetic foot ulcer in Srinivas Institute of Medical Sciences, Mangalore from Feb 2012 to Jan 2014.

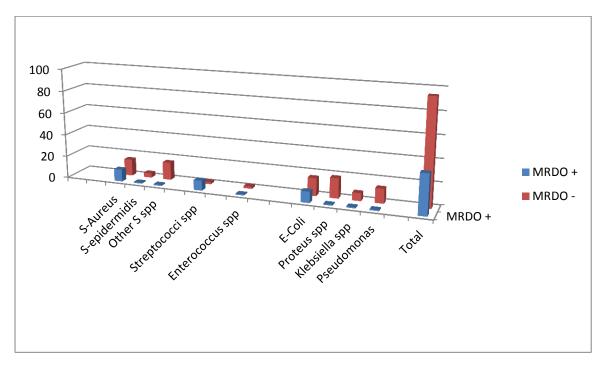
On admission, the skin around the wound was cleaned with povidone iodine solution. After careful irrigation of the wound with normal saline, remove necrotic tissue as needed. Irrigate the wound again, press a sterile cotton swab firmly into the wound and rotate to collect deep secretions. After 30 minutes, the swabs were transported to the microbiology laboratory in a sterile container. The specimens were inoculated and cultured, and the strains were identified.

Results:

Table 1: Bacterial Distribution and Drug Resistance Rate:

Bacteria	MRDO+	MRDO-
Staphylococcus spp		
S-Aureus	11	15
S-epidermidis	01	04
Other S spp	01	16
Streptococci spp	09	02
Enterococcus spp	01	02
Gram Negative		
E-Coli	10	16
Proteus spp	01	18
Klebsiella spp	01	07
Pseudomonas	01	14
Total	36	94





Discussion:

Diabetic patients often have a high prevalence of diabetic foot infection (DFI) due to associated neuropathy, peripheral vascular disease, immune abnormalities, etc. Some studies have shown that about 15% of diabetic patients can develop foot ulcers during their lifetime [10]. Diabetic foot ulcer wounds contain a large number of protein and carbohydrate nutrients, which can be a nidus for infection. About 40% to 80% of patients with diabetic foot suffer from a concurrent infection. 25% of patients with mild infection develops with severe deep infection. Clinically, the disease develops rapidly for a certain proportion of patients with mild infection and they need attention [11]. The types of diabetic foot infections are complex and varied. In addition to the common cellulitis, complications can involve osteomyelitis and gangrene [12]. Since the increasingly severe form of prevention and treatment of diabetic foot ulcer infections is associated with a high rate of detection of multidrug resistant bacteria, it is important to focus on assessing the risk factors of multi-drug resistant bacterial infections in order to find more effective treatment [13,14,15]. Multi-drug resistance (MDR) organisms refer to bacteria that exhibit drug resistance to three or more commonly used antibiotics that are usually sensitive. Multiple drug resistance also includes extensive drug resistance (XDR) and pan-drug resistance (PDR). Clinically, Common MDR methicillin-resistant staphylococcus aureus (MRSA), vancomycin-resistant enterococcus (VRE), bacteria that produce ultra-broad spectrum β-lactamase (ESBLs) in enterobacteriaceae (such as escherichia coli and klebsiella pneumoniae), carbapenemresistant enterobacteriaceae, multidrug-resistant Pseudomonas aeruginosa (MDR-PA), multidrug resistant acinetobacter baumannii (MDR-AB), etc. [16]. The aim of this study is to

analyze the bacteriological characteristics of diabetic foot ulcers, drug resistance of isolates, and to reveal the potential risk factors for multi-drug resistant bacteria. This allows for more effective assessment and treatment plans for patients with MDR diabetic foot infections.

Conclusion:

Early empiric antibiotic treatment is necessary for patients at high risk of MDRO infection. This study puts in a base for other studies so that a regional data grows so that it would be useful for the practising physicians and surgeons.

References:

- 1. Mantey I, Hill RL, Foster AV, Wilson S, Wade JJ, Edmonds ME. Infection of foot ulcers with Staphylococcus aureus associated with increased mortality in diabetic patients. Commun Dis Public Health. 2001;3(4):288–90.
- 2. Fejfarová V, Jirkovská A, Skibová J, Petkov V. Vliv rezistentních patogenů a ostatních rizikových faktorů na cetnost amputací dolních koncetin u pacientů se syndromem diabetické nohy [Pathogen resistance and other risk factors in the frequency of lower limb amputations in patients with the diabetic foot syndrome]. Vnitr Lek. 2002;48(4):302–6.
- **3.** Saltoglu N, Ergonul O, Tulek N, Yemisen M, Kadanali A, Karagoz G, et al. Influence of multidrug resistant organism on the outcome of diabetic foot infection. Int J Infect Dis. 2018;70:10–4.
- **4.** Chen Y, Ding H, Wu H, Chen HL. The relationship between osteomyelitis complication and drug-resistant infection risk in diabetic foot ulcer: a meta-analysis. Int J Low Extrem Wounds. 2017;16(3):183–90.
- **5.** Katz DE, Friedman ND, Ostrovski E, Ravid D, Amrami N, Avivi D, et al. Diabetic foot infection in hospitalized adults. J Infect Chemother. 2016;22(3):167–73.
- 6. Cassatella MA. Neutrophil-derived proteins: selling cytokines by the pound. Adv Immunol. 1999;73:369–509.
- 7. Chen Y, Ding H, Wu H, Chen HL. The relationship between osteomyelitis complication and drug-resistant infection risk in diabetic foot ulcer: a meta-analysis. Int J Low Extrem Wounds. 2017;16(3):183–90.
- **8.** Katz DE, Friedman ND, Ostrovski E, Ravid D, Amrami N, Avivi D, et al. Diabetic foot infection in hospitalized adults. J Infect Chemother. 2016;22(3):167–73.
- 9. Cassatella MA. Neutrophil-derived proteins: selling cytokines by the pound. Adv Immunol. 1999;73:369–509.
- 10. BenMoussa M, Khalfallah M, BoutibaBenBoubaker I, Nouira R, Slim A, Jerraya H, et al. Bacteriological and therapeutic profile of diabetic foot infection: a prospective study of 100 patients. La Tunisie Médicale. 2016;94(2):95–101.
- 11. Tang ZY. The comprehension to the part of diabetic foot infection in Chinese guideline on prevention and management of diabetic foot (2019 edition). World Clini Drugs. 2019;40(9):599–602.

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- 12. Price Bianca L, Lovering Andrew M, Bowling Frank L, Dobson CB. Development of a Novel Collagen Wound Model To Simulate the Activity and Distribution of Antimicrobials in Soft Tissue during Diabetic Foot Infection. Antimicrob Agents Chemother. 2016;60(11):6880–9.
- 13. Wayne Pa: Clinical and Laboratory Standards Institute: M2-A9[S]. Clinical and Laboratory Standards Institute. Performance standards for antimicrobial disk susceptibility tests 2006
- **14.** Agbi KelechiE, Carvalho Maria, Phan Ha, Tuma Cristiane. Case report: diabetic foot ulcer infection treated with topical compounded medications. Int J Pharm Compd. 2017;21(1):22–7.
- **15.** Belefquih B, Frikh M, Benlahlou Y, Maleh A, Jadid L, Bssaibis F, et al. Diabetic foot infection in Morocco: microbiological profile. Wounds A Compendium of Clin Res and Pract. 2016;28(3):89.
- 16. Huang X, Deng ZD, Ni YX, Deng M, Hu BJ, Li LY, et al. Chinese experts' consensus on prevention and control of multi-drug resistance organism healthcare-associated infection. Chin J Infect Control. 2015;14(1):741–7.