

## Bacterial Susceptibility Pattern of the Bacteria That Cause Skin and Soft Tissue Infections to Cephalexin and Co-Trimoxazole

<sup>1</sup>Alireza Sharif, <sup>2</sup>Mohammad Reza Sharif and <sup>3</sup>Javad Alizargar

<sup>1</sup>Department of Infectious Disease, Kashan University of Medical Sciences, Kashan, Iran

<sup>2</sup>Department of Pediatrics, Kashan University of Medical Sciences, Kashan, Iran

<sup>3</sup>Student Research Committee, Kashan University of Medical Sciences, Kashan, I.R. Iran

**Abstract:** Skin and soft tissue infections (SSTIs) are a common reason for presentation to outpatient practices. In our hospital settings, there are reports of high antibiotic resistance and lots of antibiotics have become useless because of this reason. We planned to study the pattern of infection and bacterial resistances of the bacteria that cause SSTI in the patients that are referred to infectious disease clinic. The overall bacterial susceptibility of the 98 growth shows that only 49(50%) of them were sensitive to cephalexin, 14(14.2%) were intermediate and 35(35.8%) were resistant. The results regarding co-trimoxazole were 34(34.6%), 11(11.3%) and 53(54.1%) for sensitive, intermediate and resistant species respectively. Strategies like treatment after getting antibiogram results seem to be necessary in treatment of SSTIs.

**Key words:** Antimicrobial • Bacteria Resistance

### INTRODUCTION

Skin and soft tissue infections (SSTIs) are a common reason for presentation to outpatient practices, emergency rooms and hospitals [1]. The incidence of skin and soft tissue infections in the ambulatory setting has more than doubled over a ten-year period to 3.4 million emergency department visits in 2005 [2]. The main reason for this increase is emergence of resistance species especially methicillin-resistant *Staphylococcus aureus* [3, 4].

SSTIs are usually treated with topical or oral antibiotics, but comorbid conditions like diabetes mellitus or invasion of the infection to deeper tissues can necessitate intravenous antibiotic therapy [5].

The need to control antibiotic resistance is felt now is more than ever [6]. There are reports of high antibiotic resistance in our hospital settings and lots of antibiotics have become useless because of that [6-12].

Treatment of SSTIs should include drainage of the abscess if possible and starting a proper antibiotic or a

combination of antibiotics [13]. If these antibiotics be chosen correctly there would be lesser antibiotic resistance due to fully eradication of the bacterial cause, thus microbiological studies concerning these infections are important [14]. In this study we are to evaluate bacterial resistance pattern of the bacteria that cause SSTI in the patients that are referred to infectious disease clinic of Shahidbeheshti clinic of infectious diseases.

### MATERIALS AND METHODS

In this retrospective study, by referring to the Shahidbeheshti Hospital laboratory, data of SSTI samples including sex, age and antibiotic susceptibility to cephalexin and co-trimoxazole of every case [15] has been added to checklists and all data were analyzed with SPSS software version 11.5. Names of the patients remained unrevealed. Antimicrobial susceptibility was evaluated by the Kirby-Bauer disk diffusion method in guidelines of Clinical and Laboratory Standards Institute [16].

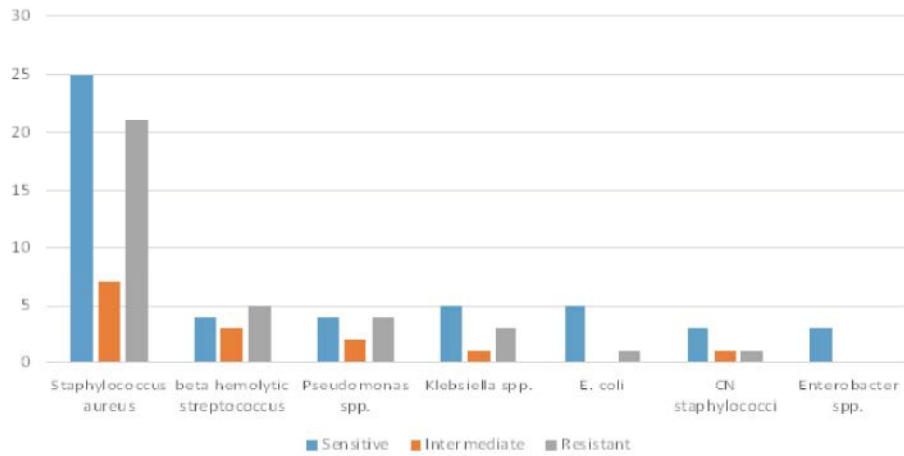


Fig 1: Distribution of different bacterial type susceptibility pattern to cephalixin in patients with SSTIs CN *Staphylococci*= coagulase-negative *Staphylococci*

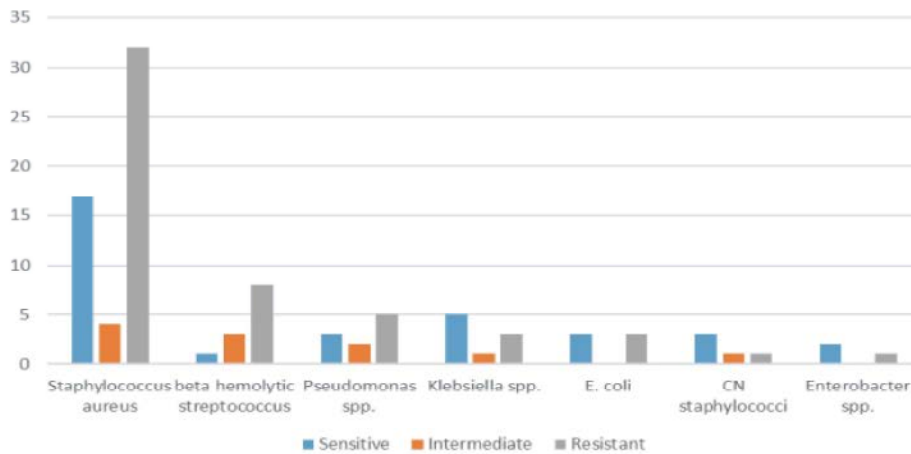


Fig 2: Distribution of different bacterial type susceptibility pattern to co-trimoxazole in patients with SSTIs CN *Staphylococci*= coagulase-negative *Staphylococci*

**RESULTS**

Data of 98 patients entered our checklists. There were 47 (48%) male patients and 51 (52%) female ones. Mean age of the patients was 59.3 years. The overall bacterial susceptibility of the 98 growth shows that only 49(50%) of them were sensitive to cephalixin, 14(14.2%) were intermediate and 35(35.8%) were resistant. There results regarding co-trimoxazole were 34(34.6%), 11(11.3%) and 53(54.1%) for sensitive, intermediate and resistant species respectively. Distribution of different bacterial type susceptibility pattern to the tested antibiotics can be seen in figure 1 and 2 and resistance rate of different bacterial types can be seen in Table 1.

Table 1: Resistance rate (%) of different bacterial types to cephalixin and co-trimoxazol in patients with SSTIs

Bacterial type	Cephalixin	Co-trimoxazole
<i>Staphylococcus aureus</i>	39.6	60.3
Beta hemolytic <i>Streptococcus</i>	41.6	66.6
<i>Pseudomonas</i> spp.	40	50
<i>Klebsiella</i> spp.	33.3	33.3
<i>E. coli</i>	16.6	50
CN <i>Staphylococci</i>	20	20
<i>Enterobacter</i> spp.	0	33.3
Overall	35.8	54.1

Table 2: Comparison of the results of different species sensitivity pattern of bacteria causing SSTI in the present study and the study of Sepehri *et al.* [21]

Bacterial type	Present study		Sepehri <i>et al.</i> study	
	Ce.	Co.	Ce.	Co.
<i>Staphylococcus aureus</i>	60.4	39.7	68.3	34.4
Beta hemolytic <i>Streptococcus</i>	58.4	33.4	40	15.4
<i>Pseudomonas</i> spp.	60	50	0	8.3
<i>Klebsiella</i> spp.	66.7	66.7	11.1	55.6
<i>E. coli</i>	83.4	50	33.3	33.3
CN <i>Staphylococci</i>	80	20	-	-
<i>Enterobacter</i> spp.	100	66.7	-	-

Ce=Cephalexin, Co=Co-trimoxazole

### DISCUSSION

This study evaluated 98 bacterial isolates recovered from SSTI patients referred to Shahidbeheshti clinic of infectious diseases and analyzed their antimicrobial resistance pattern to cephalexin and co-trimoxazole.

In a study that was conducted by Goh *et al.* [17] there were 100% sensitivity to cephalexin and co-trimoxazole in *Staphylococcus aureus*. But our results showed high resistance of *Staphylococcus aureus* against it. Regarding beta hemolytic *Streptococcus* Jain *et al.* [18] reported 12.2% resistance against co-trimoxazole that is much less than ours 66.6% resistance. Resistance rate of *Pseudomonas* spp. to co-trimoxazole was announced 86% in the study of Nikokar *et al.* [19] that is higher than that reported in our study. *Enterobacter* spp. have been reported to have high resistance rates in previous studies [20], but all of our 3 isolates of *Enterobacter* was sensitive to cephalexin. This may be because of our limited study number.

Another study that conducted in Iran on antibiotic susceptibility of isolates that cause SSTI by Sepehri *et al.* [21] showed that the overall antibiotic susceptibility pattern of different bacterial types that cause SSTI to cephalexin was 41.8% sensitive, 17.7% intermediate and 40.5% resistant and 26.3% sensitive, 19.5% intermediate and 54.2% resistant for co-trimoxazole. Our results are in concordance with these results and show the same antibiotic usage pattern of these two antibiotics in the two studies. The results of comparing species sensitivities of the two studies are illustrated in table 2.

Our results of antibiotic resistance of bacteria that cause SSTI to cephalexin and co-trimoxazole are high

especially regarding cephalexin resistance of the most prevalent bacterial causes of SSTI and these antibiotics are somehow useless because of more than 50% or even near 100% resistance rates in some species. Strategies like treatment after getting antibiogram results seem to be necessary in treatment of SSTIs.

### REFERENCES

1. Rajan, S., 2012. Skin and soft-tissue infections: Classifying and treating a spectrum. *Cleveland Clinic Journal of Medicine*, 79: 57-66.
2. Pallin, D.J., D.J. Egan, A.J. Pelletier, J.A. Espinola, D.C. Hooper and C.A.Jr. Camargo, 2008. Increased US emergency department visits for skin and soft tissue infections and changes in antibiotic choices, during the emergence of community-associated methicillin-resistant *Staphylococcus aureus*. *Ann. Emerg. Med.*, 51: 291-298.
3. Fridkin, S.K., J.C. Hageman, M. Morrison, L.T. Sanza, K. Como-Sabetti, J.A. Jernigan, K. Harriman, L.H. Harrison, R. Lynfield, M.M. Farley and Active Bacterial Core Surveillance Program of the Emerging Infections Program Network, 2005. Methicillin-resistant *Staphylococcus aureus* disease in three communities. *N. Engl. J. Med.*, 352: 1436-1444.
4. Klevens, R.M., M.A. Morrison, J. Nadle, S. Petit, K. Gershman, S. Ray, L.H. Harrison, R. Lynfield, G. Dumyati, J.M. Townes, A.S. Craig, E.R. Zell, G.E. Fosheim, L.K. McDougal, R.B. Carey, S.K. Fridkin and Active Bacterial Core Surveillance (ABCs) MRSA Investigators, 2007. Invasive methicillin-resistant *Staphylococcus aureus* infections in the United States. *JAMA*, 298: 1763-1771.
5. Wilson, S.E., 2001. Clinical trial results with linezolid, an oxazolidinone, in the treatment of soft tissue and postoperative gram-positive infections. *Surg. Infect (Larchmt)*, 2: 25-35.
6. Sharif, MR., J. Alizargar and A. Sharif, 2013. Antimicrobial resistance among Gram-negative bacteria isolated from different samples of patients admitted to a University hospital in Kashan, Iran. *Advances in Biological Research*, 7: 199-202.
7. Alizargar, J., M.R. Sharif and A. Sharif, 2013. Risk factors of methicillin-resistant *staphylococcus aureus* colonization in diabetic outpatients, a prospective cohort study. *International Journal of Microbiological Research*, 4: 147-151.

8. Sharif, MR., J. Alizargar and A. Sharif, 2013. Prevalence of methicillin-resistant *Staphylococcus aureus* nasal carriage in children admitted to Shahidbeheshti hospital. World Journal of Medical Sciences, 9: 109-112
9. Sharif, M.R., J. Alizargar and A. Sharif, 2013. Prevalence and antimicrobial susceptibility pattern of *Staphylococcus aureus* isolates at Shahidbeheshti hospital. World Journal of Medical Sciences, 9: 84-87.
10. Sharif, MR., J. Alizargar and A. Sharif, 2013. Prevalence and antibiotic susceptibility pattern of microbial agents that cause urinary tract infection. Middle-East J. of Scientific Research, 17: 1512-1515.
11. Sharif, M.R., J. Alizargar and A. Sharif, 2013. Antibiotic susceptibility of *Staphylococcus aureus* in isolates of the patients with osteomyelitis. World Journal of Medical Sciences, 9: 180-183.
12. Sharif, A., M.R. Sharif and J. Alizargar, 2014. Susceptibility of gram negative bacteria to three frequently used antibiotics from specimens sent to the laboratory of Shahidbeheshti hospital. Middle-East Journal of Scientific Research, 20: 1041-1045.
13. Bosu, W.K., S. Acquah, 1996. Susceptibility of urinary tract bacteria to antibiotics in Cape Coast, Ghana. East African Medical Journal, 73: 468-470.
14. Agulur, G.M., 1992. The emergence of highly fluoroquinolone resistant *Escherichia coli* in community-acquired urinary tract infection. J. Antimicrob Chemother, 29: 369-370.
15. Dan, M., I. Somer, B. Knobel and Gutman, 1988. Cutaneous manifestation of infection with *Corynebacterium* group. Rev Infect Dis., 10: 1204-1207.
16. CLSI, 2013. Performance Standards for Antimicrobial Susceptibility Testing; Twenty-Third Informational Supplement. CLSI document M100-S23. Wayne, PA: Clinical and Laboratory Standards Institute.
17. Goh, C.L., J.S. Wong and Y.C. Giam, 1997. Skin colonization of *Staphylococcus aureus* in atopic dermatitis patients seen at the National Skin Centre, Singapore. Int. J. Dermatol, 36: 653-657.
18. Jain, A., V.K. Shukla, V. Tiwari and R. Kumar, 2008. Antibiotic resistance pattern of group-a beta-hemolytic *Streptococci* isolated from north Indian children. Indian J. Med. Sci., 62: 392-396.
19. Nikokar, I., A. Tishayar, Z. Flakiyan, K. Alijani, S. Rehana-Banisaeed, M. Hossinpour, S. Amir-Alvaei and A. Araghian, 2013. Antibiotic resistance and frequency of class 1 integrons among *Pseudomonas aeruginosa*, isolated from burn patients in Guilan, Iran. Iran J. Microbiol, 5: 36-41.
20. Wenzel, R.P., D.F Sahn, C. Thornsberry, D.C. Draghi, M.E. Jones and J.A. Karlowky, 2003. *In Vitro* Susceptibilities of Gram-Negative Bacteria Isolated from Hospitalized Patients in Four European Countries, Canada and the United States in 2000-2001 to Expanded-Spectrum Cephalosporins and Comparator Antimicrobials: Implications for Therapy. Antimicrob Agents Chemother, 47: 3089-3098.
21. Sepehri GH.R., S. Dabiri and Z. Zeinadini Meimandi, 2004. Antimicrobial resistance of isolates of cutaneous, soft tissue and visceral abscesses. Hakim, 6: 65-71.