

Original Article

Drug resistance of isolated strains of *Pseudomonas Aeruginosa* from burn wound infections to selected antibiotics and disinfectants

Parviz Owlia¹, Horieh Saderi¹, Sadegh Mansouri¹, Sirus Salemi¹, Hossein Ameli¹

1. Department of Microbiology, School of Medicine, Shahed University, Tehran

ABSTRACT

Background and Objectives: Infection is the most common problem following burn injury. Selection and dissemination of intrinsic and acquired resistance mechanisms increase the probability of burn wound colonization by resistant species including *Pseudomonas aeruginosa*. Multi-drug resistant *Pseudomonas aeruginosa* has frequently been reported as the cause of nosocomial outbreaks of infection in burn wards or as colonizers of the wound of burned patients. Therefore, this research study was conducted to compare the activity of various antibiotics and disinfectants against clinically important strains of *P. aeruginosa*.

Materials and Methods: One hundred strains of *P. aeruginosa* were obtained as clinical isolates from burn wound infections. The antimicrobial activity of antibiotics was tested by disk diffusion method of Kirby-Baur. For disinfectants, 30 µl of each of them was placed on sterile blank disk and studied by disk diffusion method.

Results: The frequency of resistant strains to kanamycin, tobramycin, amikacin, cefotaxime, carbenicillin, ceftazidime, ceftizoxime, cefixim, ciprofloxacin, cefazolin, cephalaxine, and ceftriaxone was 100, 93, 95, 81, 84, 95, 94, 100, 99, 100, 100, and 92 respectively. The averaged diameter of inhibition zone for chlorhexidine (0.2%), povidone iodine (10%), cetrimide-C (3.5%), dekonsept, hypochlorite (10%), micro 10⁺ (2%), deconex 53⁺ (2%), and ethanol (70%) was 14.4 ± 1.9 mm, 10.6 ± 1.3 mm, 9.1 ± 2.6 mm, 8.6 ± 2.2 mm, 26.9 ± 5.2 mm, 6.58 ± 1.5 mm, 8.3 ± 2.2 mm, and 6 ± 0.0 mm respectively.

Conclusion: The high frequency of resistance to antibiotics and sensitivity to a few disinfectants suggests to restrict the spread of *P. aeruginosa* and to limit administration of these antibiotics and to use of hypochlorite and chlorhexidin as disinfectant as a preventive treatment.

Key words: *Pseudomonas aeruginosa*, Burn infection, Antimicrobial resistance

Received: 20 February 2006

Accepted: 1 May 2006

*Address communications to: Dr. Parviz Owlia (PhD), Department of Microbiology, School of Medicine, No. 29, Abdollahzadeh St., Keshavarz Blvd, Shahed University, Tehran-Iran

E-mail: owlia@shahed.ac.ir

Introduction

Burns remain a huge public health issue, at least in terms of morbidity and long term disability throughout the world, especially in the developing countries. Burn injuries still produce a significant morbidity and mortality in Iran (1). Burn-wound infection is one of the most common causes of death and serious problem after burn injury (2, 3). Burn predisposes the body to infection by damaging the protective barrier function of the skin, thus facilitating the entry of pathogenic microorganisms and by inducing systemic immunosuppression. In this respect, *Pseudomonas aeruginosa* (*P. aeruginosa*) is an opportunistic pathogen that produces a number of unique virulence factors. Extracellular toxin, proteases, hemolysins, and exopolysaccharides are a few types of virulence factors that have been implicated in the pathogenicity of *P. aeruginosa* (4 – 6). In addition, *P. aeruginosa* is a major cause of burn injury colonization and serious wound infection. Furthermore, *P. aeruginosa* remains the leading pathogen causing wound infection in the Tohid burn center (Tehran, Iran), and the frequency of its infection is higher than reported from other countries (7, 8). Multi-drug resistance of *P. aeruginosa* is another complication in affected patients (9).

Therefore, the purpose of this study was to determine the susceptibility of clinically isolated *P. aeruginosa* from burn infection to selected antibiotics and disinfectants.

Materials and Methods

One hundred strains of *P. aeruginosa* were obtained as clinical isolates from burn wound infections during a period of 10 months from Tohid burn center (Tehran, Iran). *P. aeruginosa* ATCC 27853 and *Staphylococcus aureus* ATCC 25923 were included as control strains to verify the accuracy of the antibiotic susceptibility test procedure. The antimicrobial activity of kanamycin (30 µg), tobramycin (10 µg), amikacin (30 µg), gentamicin (10 µg), cephotaxim (30

µg), carbenicillin (100 µg), ceftazidime (30 µg), ceftizoxime (30 µg), cefixime (5 µg), ciprofloxacin (5 µg), cefazoline (30 µg), cephalaxine (30 µg), and ceftriaxon (30 µg) (Padtan Teb Co, Iran) against *P. aeruginosa* were tested by disk diffusion method of Kirby-Baur. Bacterial suspensions with 10⁶ CFU/ml were spread on the surface of Muller-Hinton agar plates. Within 15 minutes after the surface of the agar has been inoculated, antimicrobial disks were applied with sterile forceps. The plates were inverted and incubated at 35 °C for 18 h and then diameter of each zone of inhibition was measured (10).

The antimicrobial activity of chlorhexidine (0.2%), povidone iodine (10%), cetrimide-C (3.5%), dekosept, hypochlorite (10%), micro 10⁺ (2%), deconex 53⁺ (2%), and ethanol (70%) were tested by disk diffusion method. For disk diffusion method, 30 µl of each of above disinfectants was placed on sterile blank 6 mm-diameter disk. Then, the disks were manipulated with sterile forceps. Thereafter, plates were incubated at 35 °C for 18 h and then diameter of each zone of inhibition was measured (11).

Results

From 100 isolated strains of *P. aeruginosa*, the frequency of resistant strains to kanamycin, tobramycin, amikacin, cefotaxime, carbenicillin, ceftazidime, ceftizoxime, cefixim, ciprofloxacin, cefazolin, cephalaxine, and ceftriaxon were between 84 and 100 (Table 1).

The results of antimicrobial effects of disinfectants had shown that mean of diameter of inhibition zone for chlorhexidine (0.2%), povidone iodine (10%), cetrimide-C (3.5%), dekosept, hypochlorite (10%), micro 10⁺ (2%), deconex 53⁺ (2%), and ethanol (70%) were 14.4 ± 1.9 mm, 10.6 ± 1.3 mm, 9.1 ± 2.6 mm, 8.6 ± 2.2 mm, 26.9 ± 5.2 mm, 6.6 ± 1.5 mm, 8.3 ± 2.2 mm, and 6 ± 0.0 mm respectively by disk diffusion method. Inhibition zone for ethanol was not seen and all strains were completely resistant to this disinfectant (Table 2).

Table 1. The results of antibiogram for isolated strains of *P. aeruginosa*

Antibiotics	Number of strains of:		
	R	I	S
Kananycin	100	0	0
Toramycin	93	5	2
Amikacin	95	2	3
Gentamicin	96	2	2
Cefotaxime	81	17	2
Carbenicillin	84	9	7
Ceftazidime	95	2	3
Ceftizoxime	94	2	4
Cefixime	100	0	0
Ciprofloxacin	99	5	6
Cefazoline	100	0	0
Cephalexine	100	0	0
Ceftriaxon	92	6	2

R=Resistant I= Intermediate S= Sensitive

Table 2. Means of inhibition zone of disinfectants in disk diffusion method

Disinfectants	Means of inhibition zone (mm)
Chlorhexidine	14.4 ± 1.9
Povidone iodine	10.6 ± 1.3
Cetrimide- C	9.1 ± 2.6
Dekosept	8.6 ± 2.2
Hypochlorite	26.9 ± 5.2
Micro 10 ⁺	6.6 ± 1.5
Deconex 53 ⁺	8.3 ± 2.2
Ethanol	6 ± 0.0*

Discussion

It has been reported that *P. aeruginosa* remains the leading pathogen causing wound infection in Tohid burn center (Tehran, Iran) (8). The incidence of *P. aeruginosa* infection in Tohid burn center has been higher than reported from other countries (7). In addition, an increasing trend in its resistance is a major problem in *Pseudomonas* infections. In this study, 100% of isolated strains

were resistant to the kanamycin, cefixime, cefazoline, and cephalexine. In a similar study by Rastegar Lari et al in 1995, it had been shown that the rates of resistance for amikacin, ciprofloxacin, and gentamicin were 48.9%, 45.2%, and 88.5% respectively (7). In our study (in 2002), we found that resistance to the mentioned antibiotics increased to 95%, 99%, and 96% respectively.

We also used the zone of inhibition method to test the efficacy of various disinfectants against isolated strains. It was found out that the most effective disinfectants were hypochlorite, povidone iodine, chlorhexidine, cetrimide- C and deconex 53⁺. These results have also been confirmed by other studies. In this regard, Sterla et al and Leonardo et al had shown that chlorhexidin is effective against *P. aeruginosa* (12, 13). Meanwhile, Chogawala et al (1990) had shown that povidone iodine (1%) is effective on *P. aeruginosa* (14). A few studies have also proved the efficacy of commercial sodium hypochlorite against *P. aeruginosa* (13, 15-16). The important point to consider is that ethanol 70% has not been effective in this respect, although it has been commonly used as disinfectant. The high incidence and wide spread of *P. aeruginosa* and its high resistance to antibiotics as mentioned above indicates the necessity for urgent measures to be taken to restrict the spread of this pathogen and to limit administration of these antimicrobial agents. Also, we suggest using disinfectants hypochlorite and chlorhexidin as a preventive treatment.

Conclusion

considering the high frequency of resistance of *P. aeruginosa* to the current antibiotics, it is possible that the infections of this pathogen may be common, and so, molecular epidemiology for determining its clone in Tohid burn center is clinically very essential.

References

1. Panjeshahin MR, Rastegar Lari A, Talei AR, et al. Epidemiology and mortality of burn in the South West of Iran. *Burns* 2001 27: 219-26

2. Mason AD, McManus AT, Pruitt BA, et al. Association of burn mortality and bacteremia. Arch Surg 1986 121: 1027-31
3. McManus AT, Mason AD, McManus WF, et al. Twenty five-years review of *Pseudomonas aeruginosa* bacteremia in a burn center. Eur J Clin Microbiol 1985 4: 219-33
4. Demko CA, Thomassen MG. Effect of mucoid property of antibiotic susceptibility of *Pseudomonas aeruginosa*, Curr Microbiol 1980 4: 69-73
5. Masuda N, Goton N, Ishii C, et al. Interplay between chromosomal β -lactamas and Mex AB – OprM Efflux system in intrinsic resistant to β -lactamas in *Pseudomonas aeruginosa*. Antimicrob Agent Chemother 1999 43: 400-2.
6. Slack MPE, Nichols WW. The penetration of antibiotics through sodium alginate and through the exopolysaccharide of a mucoid strain of *Pseudomonas aeruginosa*. Lancet 1981 2: 502-3
7. Rastegar Lari A, Bahrami Honar H, Alaghebandan R. *Pseudomonas* infections in Tohid Burn Center, Iran. Burns 1998 24: 637-41
8. Rastegar Lari A, Alaghebandan R. Nosocomial infection in an Iranian burn care center. Burns. 2000 26: 737- 40
9. Eykyn SJ. Treatment of *Pseudomonas aeruginosa* infection. Br J Hospital Med. 1984 0: 147-40
10. Acar FJ, Goldstein . Disk susceptibility test. In: Lorian V, eds: Antibiotics in laboratory medicine, 4th ed. Baltmor: Williams & Wilkins, 1996 1- 34
11. Kovacs BJ, Aprecio RM, Ketring JD, et al. Efficacy of various disinfectants in killing a resistant strain of *Pseudomonas aeruginosa* by comparing zones of inhibition: Implication for endoscopic equipment reprocessing. Am J Gastroentero 1998 93 (11): 2057-9
12. Esterla C, Ribeiro RG, Esterla CR, et al . Antimicrobial effect of 2% sodium hypochlorite and 2% chlorhexidin tested by different methods. Braz Dent J 2003 14(1): 58-62
13. Leonardo MR, da Silva LA, Filho MT, et al. Invitro evaluation of the antimicrobial activity of a castor oil-based irrigant. J Endod 2001 27(12): 717-9
14. Chogawala Z, Furtado D. Invitro and invivo bactericidal activities of 10%, 2.5% and 1% povidon iodine solution. Am J Hos Pharm 1990 47:1562-66
15. Merritt K, Hitchins VM, Brown SA. Safety and cleaning of medical materials and devices. J Biomed Mater Res 2000 53(2): 131-6
16. Cotter JL, Fader RC, Lilley C, et al. Chemical parameters, antimicrobial activities, and tissue toxicity of 0.1 and 0.5% sodium hypochlorite solutios. Antimicrob Agents Chemother 1985 25(1); 118-22
17. Coates D. Comparison of sodium hypochlorite and sodium dicloro-isocyanarate disinfectants: neutralization by serum. J Hosp Infect 1988 11(1): 60-7