

Characterization of Carbapenemases in Extensively Drug Resistance *Acinetobacter baumannii* in a Burn Care Center in Iran

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The emergence of multidrug resistance (MDR) and extensively drug resistance (XDR) in *Acinetobacter baumannii* has made an important challenge in the treatment of infections caused by this organism. The ability of carbapenemase production is one of the main mechanisms for the emergence of MDR and/or XDR in *A. baumannii*. The aim of this study was to detect carbapenemase producer *A. baumannii*. In this study, 65 imipenem resistant *A. baumannii* were collected from burned patients. Biochemical identification, antibiotic susceptibility test and multiplex polymerase chain reactions for the detection of carbapenemases genes were performed. The results showed that all strains carried *bla*_{OXA-51}. 83%, 12.5% and 9.23% strains harbored *bla*_{OXA-23}, *bla*_{VIM} and *bla*_{KPC} genes, respectively. None of the isolates carried *bla*_{IMP}, *bla*_{OXA-48}, *bla*_{NDM-1} and *bla*_{SPM-1} genes. The results of this study indicate the emergence of *Klebsiella pneumoniae* Carbapenemase (KPC) in *A. baumannii* causing nosocomial infections in burned patients which can be important for hospital infection prevention systems in Iran.

Key words: *Acinetobacter baumannii*, carbapenemases, KPC

A *Acinetobacter baumannii* has been recognized as one of the important causes of nosocomial infections in hospitalized patients, particularly in burned ones in recent years (1- 3). During the past decade, in some countries, this opportunistic pathogen had high rate of infection in burned patients and was reported to be the second most common cause of nosocomial infections in burned patients (1, 2, 4-7). *A. baumannii* has been shown to

acquire fast antibiotic resistance elements (8, 9). Recently, this Gram negative bacilli has shown resistance to the most available antibiotics followed by the emergence of multiple (MDR) and extensive drug resistance (XDR) strains (2, 8-10). This has partly been due to the extensive use of broad spectrum antibiotics especially in burned patients (9). Carbapenems are generally used for antibiotic therapy in infections due to MDR *A. baumannii*.

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Characterization of Carbapenemases in Drug Resistance *A. baumannii*

However, the emergence of carbapenem resistant isolates has made it difficult to treat such infections (2, 3, 8, 11-14). One of the most common and important mechanisms in carbapenem resistance in *A. baumannii* is their ability to produce carbapenemase enzymes (2, 3, 6-8). Among the carbapenem hydrolyzing β -lactamases, class D oxacillinases (OXA type) is the most prevalent in *A. baumannii* strains (3, 9, 14). On the other hand, some studies have reported class B **beta lactamases** (metallo- beta lactamases) in *A.*

baumannii as the most prevalent carbapenemase after OXA- type (8). The significant importance is the worldwide prevalence of *Klebsiella pneumoniae* Carbapenemase (KPC) producers (15, 16). Gram negative KPC producers can be resistant to all beta lactam antibiotics except aztreonam (17). Most of these carbapenemases have been located in transferable genetic elements and can spread among *A. baumannii* and even into other Gram- negative bacilli (2, 8). Therefore, the determination of carbapenemase- producing strains in healthcare

Table 1. Sequence of primers

Primer	Sequence (5'→3')	PCR pduct size (bp)	References
VIM F	TTGACACTCCATTTACDG	390	
VIM R	GATYGAGAATTAAGCCACYCT		
Imp F	GATGGTGTTTGGTCGCATA	139	8
Imp R	CGAATGCGCAGCACCAG		
OXA-23 F	GATGTGTCATAGTATTCGTCGT	1050	
OXA-23 R	TCACAACAACATAAAAGCACTGT		9
OXA-48-ike F	CCAAGCATTTTTACCCGCATCKACC	389	
OXA-48-like R	GYT TGA CCA TAC GCT GRC TGC G		
NDM-1 F	CCCGGCCACACCAGTGACA	129	
NDM-1 R	GTAGTGCTCAGTGTCGGCAT		9
SPM-1 F	GGGTGGCTAAGACTATGAAGCC	447	
SPM-1 R	GCCGCCGAGCTGAATCGG		
OXA-51-like F	TAATGCTTTGATCGGCCTTG	353	7
OXA-51-like R	TGGATTGCACTTCATCTTGG		
KPC F	GTATCGCCGTCTAGTTCTGC	636	18
KPC R	GGTCGTGTTCCCTTTAGCC		

Table 2. PCR conditions for *bla* genes amplification

Gene	Initial denaturation Temperature (time)	Denaturation Temperature (time)	Annealing Temperature (time)	Extension Temperature (time)	Final extension Temperature (time)	No. Cycles
<i>bla</i> _{VIM} & <i>bla</i> _{IM:}	94 °C (10 min)	94 °C (40 sec)	60 °C (40 sec)	72 °C (1 min)	72 °C (7 min)	30
<i>bla</i> _{OXA-23} & <i>bla</i> _{OXA-48}	94 °C (1 min)	94 °C (30 sec)	55 °C (1 min)	72 °C (1 min)	72 °C (7 min)	30
<i>bla</i> _{NDM-1} & <i>bla</i> _{SPM-1}	94 °C (1 min)	94 °C (30 sec)	60 °C (40 sec)	72 °C (1 min)	72 °C (1 min)	30
<i>bla</i> _{KPC}	94 °C (5 min)	94 °C (1 min)	56 °C (1 min)	72 °C (1 min)	72 °C (5 min)	30
<i>bla</i> _{OXA-51}	94 °C (5 min)	94 °C (45 sec)	58 °C (1 min)	72 °C (1 min)	72 °C (5 min)	30

systems could help in the eradication of *A. baumannii* antibiotic resistant strains.

Materials and methods

Bacterial isolates

This cross-sectional study was conducted in Motahari Hospital which is the only referral burn center in Tehran, Iran. Sixty-five imipenem resistant *A. baumannii* isolated from burn wound were collected from admitted patients from April to July 2013. Primary identification of these strains was performed by conventional biochemical tests such as oxidase, TSI, SIM and growing at 44 °C as *A. baumannii*. The identification of isolates was confirmed by API 20 NE (Bio-Merieux, Lyon, France) strip test.

Antimicrobial susceptibility testing

The antibiotic susceptibility testing was performed by disc diffusion method on Mueller–Hinton agar using cefotaxime (30 µg), ceftazidime (30 µg), imipenem (10 µg), ticarcillin (75 µg), ticarcillin- clavulanic acid (75/10 µg), piperacillin (100 µg), piperacillin- tazobactam (100/10 µg), ciprofloxacin (5 µg), gentamicin (10 µg), tobramycin (10 µg), amikacin (30 µg), tetracycline (30 µg), trimethoprim (5 µg) and trimethoprim-

sulfamethoxazole (1.25/23.75 µg) according to clinical and laboratory standards institute (CLSI) 2011. The standard antibiotic disks used in this study were from MAST Company (Mast Diagnostics, UK). *P. aeruginosa* ATCC 27853 was used as control strain in the antibiotic susceptibility testing. Detection of extended spectrum beta lactamase (ESBL) has been conducted by combination disc method by using ceftazidime and ceftazidime plus clavulanic acid. Strains with increasing at least 5 mm in the diameter of inhibition zone around ceftazidime plus clavulanic acid in contrast of ceftazidime alone were considered as an ESBL producer.

PCR amplification for carbapenemases production genes

Extraction of bacterial DNA was performed with a plasmid Minikit (Qiagen GmbH, Hilden, Germany) according to the manufacturer's instructions.

Three different sets of multiplex PCR for *bla_{VIM}*, *bla_{IMP}* and *bla_{OXA-23}*, *bla_{OXA-48}* and *bla_{NDM-1}*, *bla_{SPM-1}* genes and common PCR for *bla_{KPC}* and *bla_{OXA-51}* genes were designed. The lists of primers used are shown in Table 1 and amplification conditions are represented in Table 2.

Table 3. Antibiotic resistance patterns

patterns	Antibiotic Resistance patterns	Number	Percentage
1	CTX CAZ CEF IMI PTZ PYR TC TC-C GM AK TO TM SXT CI	36	55
2	CTX CAZ CEF IMI PTZ PYR TC TC-C GM AK TO TM SXT CI T	16	25
3	CTX CAZ CEF IMI PTZ PYR TC TC-C AK TM SXT CI T	7	11
4	CTX CAZ CEF IMI PTZ PYR TC TC-C GM TO TM SXT CI	3	5
5	CTX CAZ CEF IMI PTZ PYR TC TC-C AK TO TM SXT CI T	2	3
7	CTX CAZ CEF IMI PTZ PYR TC TC-C AK TO TM SXT CI	1	1

CTX: cefotaxime, CAZ: ceftazidime, CEF: CEFEPIME, IMI: imipenem, PTZ: piperacillin-Tazobactam, PRL: piperacillin, TC: ticarcillin, TC-c: ticarcillin- clavulanic acid, GM: gentamicin, AK: amikacin, TO: tobramycin, TM: trimethoprim, SXT: trimethoprim-sulfamethoxazole, CI: ciprofloxacin, T: tetracycline

Table 4. Number and percentages of detected *bla* genes

genes	<i>bla_{OXA-51}</i>	<i>bla_{OXA-23}</i>	<i>bla_{VIM}</i>	<i>bla_{KPC}</i> and <i>bla_{OXA-23}</i>	<i>bla_{VIM}</i> and <i>bla_{OXA-23}</i>
NO.(%)	63(100%)	42(66.6%)	2(3.1%)	6(9.52%)	6(9.52%)

Characterization of Carbapenemases in Drug Resistance *A. baumannii*

PCR products were analyzed by electrophoresis on 1.5% agarose gel with SYBR safe staining. Direct sequencing of amplicons was carried out using ABI 3730X capillary sequencer (Genfanavarun, Macrogen, Seoul, Korea).

A. baumannii ATCC 19606 was used as a negative control in all PCR run and an internal positive control was used for each gene

Results

A total of 65 *A. baumannii* were isolated from 65 hospitalized burned patients, 15 (23%) females and 50 (77%) males. The age of patients were between 9 months to 72 years and total body burned surface area (TBSA) was more than 6%. Causes of burn were classified as follows: fire 6 (11%), hot water 6 (11%), electricity 10 (17%) gas 10 (17%), 6 (11%) chemical materials and others 19 (33%). Based on antibiotic susceptibility test, 97% of strains were resistant to all tested beta lactam antibiotics and 80% of them showed resistance to all tested aminoglycoside antibiotics. The results of antibiotic susceptibility tests indicated seven antibiotic resistance patterns (Table 3). Pattern one was observed in 55% of isolates. Those strains were resistant to all tested antibiotics except colistin and tetracycline. In pattern two, all strains were resistant to the tested antibiotics which were observed in 25% of strains. In 20% strains,

resistance to at least one of aminoglycoside antibiotics were observed according to antibiotic resistance patterns.

The Verona integron-encoded metallo- β -lactamase (VIM) and KPC producing strains were observed just in two antibiotic resistance patterns. OXA-23 and OXA-51 producing strains showed all seven antibiotic resistance patterns.

The ability of ESBL production was indicated just in one strain in combination disc method.

Sequence analysis showed that *bla*_{OXA-51} gene was detected in all isolates, 54 (83%) isolates included *bla*_{OXA-23} gene. Eight (12.5%) and 6 (9.23%) strains carried *bla*_{VIM} and *bla*_{KPC} genes respectively (Figures 1-4). Six out of eight VIM-producing strains contained both *bla*_{OXA-23} and *bla*_{vim} genes (Table 4). All of KPC producer isolates were combined by OXA-23. On the other hand, none of them contained *bla*_{IMP}, *bla*_{OXA-48}, *bla*_{NDM-1} and *bla*_{SPM-1} genes.

Discussion

According to many studies, *A. baumannii* is known as one of the most common Gram negative bacteria that can cause nosocomial infection in health care centers especially in burned hospitalized patients (2, 3, 8). Treatment of infection due to XDR *A. baumannii* is extremely difficult and causes more morbidity and motility in hospitalized burn



Fig. 1. Multiplex PCR amplification fragments for the detection of *vim* and *imp* gene among *Acinetobacter baumannii* isolates. Lane 1: negative control, lanes 2, 3: positive *vim* strains, lane 4: positive control of *vim* and *imp* genes and M: 1kb DNA size marker

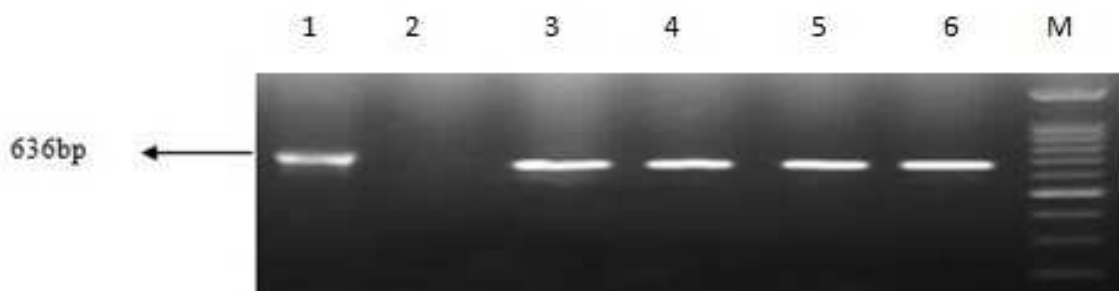


Fig. 2. PCR amplification fragments for the detection of *kpc* gene among *Acinetobacter baumannii* isolates. Lane 1: positive control of *kpc* gene, lane 2: negative control, lanes 3- 6: positive *kpc* strains and M: 1kb DNA size marker



Fig. 3. PCR amplification fragments for the detection of *oxa-23* and *oxa-48* genes among *Acinetobacter baumannii* isolates. Lane 1- 4 and 7, 8: positive *oxa-23* strains, lanes 5, 6: negative control of *oxa-23* and *oxa-48* genes, lane 9: positive control of *oxa-23* and *oxa-48* genes and M: 1kb DNA size marker

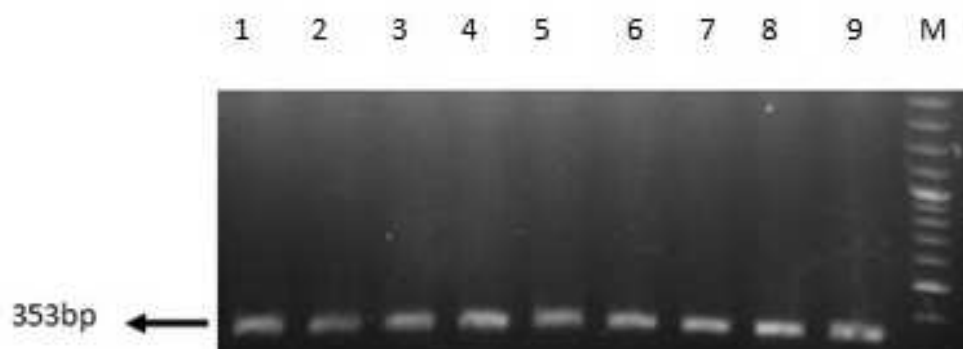


Fig 4. PCR amplification fragments for the detection of *oxa-51* gene among *Acinetobacter baumannii* isolates. Lane 1: positive control of *oxa-51*, lanes 2- 9: positive strains of *oxa-51*.and M: 1kb DNA size marker

patients (13, 19). The results of this study indicated that 98.4% of *A. baumannii* were resistant to all tested beta lactam antibiotics and 55% of them remained susceptible just to colistin and tetracycline. Generally, carbapenemases producer strains of *A. baumannii* convert to XDR strains and can be resistant to most antibiotics used for the treatment of *A. baumannii* infections. Wounds of burned patients generally infected by multiple bacteria harboring antibiotic resistance genes located in transferable genetic elements, can be a

disaster in burn care units. Resistance to all tested antibiotics in 50% of isolated KPC- producer *A. baumannii* and to all tested antibiotics except tetracycline in the remaining isolates is a significant point and creates limitation for treatment choices of this kind of infection. OXA type is the common carbapenemase in *A. baumannii* and OXA-23 is the common one in Iran (3, 8, 14, 20). Outbreaks of OXA-23 producing *A. baumannii* have been reported in several Asian countries such as China, Thailand, Taiwan (8, 6, 11). On the other hand,

metallo- beta lactamase (MBL) can be considered as a second common carbapenemase in *A. baumannii*. In the research which was carried out by Szejbach in 2013 in Poland on MBL- producer *A. baumannii*, it was shown that 10.3% of isolates carried *bla_{IMP-like}* gene. No *bla_{VIM-4}* was detected in the isolates (7). On the contrary, in the same study in India in 2012, 47% of isolated *A. baumannii* carried the *bla_{VIM}* and 0.9% of them harbored *bla_{IMP}*. The variants result may be related to different ecology situation, antibiotic therapy program and variants antibiotype patterns in two different countries (21). In 2013 in China, OXA-51 gene was detected in all isolated *A. baumannii* and 94% carried *oxa-23* gene (22). Also, the results of a study in 2012 on 68 imipenem-resistant *A. baumannii* in Iran indicated that OXA-51 and OXA-23 types were detected in all strains (20). But in the current study, 83% of isolates were identified as an OXA-23-producer which can be associated to the source of isolated bacteria. In this study, all isolates were collected from wounds of burned patients but in the study carried out in 2012 in Iran (20), the isolates were obtained from different medical specimens. On the other hand, in 2012 in another study in Iran, 88.7% of imipenem resistant *A. baumannii* carried *bla_{OXA-23-like}* gene (14) and these results are similar to ours.

In Korea in 2013, 97% of *Acinetobacter* isolates harbored the *bla_{OXA-23-like}* gene (23).

Mutation in OXA type enzymes because of overuse and/or abuses of carbapenem antibiotics in clinic can lead to high carbapenem activity by OXA type carbapenemase enzymes. Because they generally have low activity of carbapenem hydrolyzing activity (9, 24). In addition, we had a report from the prevalence of OXA-48- producer *K. pneumonia* at the same time and same burn unit and this gene was located in mobile genetic elements. Thus, we have detected specifically *oxa-48* gene in isolated *A. baumannii* (25). Consequently, it can be an alarming threat in health in Iran. Moreover, as

the gene of this carbapenemase is located in a mobile genetic element and can be transferred to *A. baumannii*, it can impose more complications in burn patient's therapy. In the current study, 97% of carbapenemase producer strains subsequently were resistant to all beta lactam antibiotics and 80% of them were resistant to aminoglycoside. These results can be a concern for physicians because 78% of the strains were sensitive to colistin and tetracycline only. The results of this study indicated that the ability of carbapenemase production can be one of the most important reasons for the emergence of XDR *A. baumannii* in nosocomial infections. The infection is important in burned patients because their skin lose a first protective barrier, putting the patients at high risk for multiple infections.

Regarding to the mentioned above study and our results, it seems that *A. baumannii* is becoming one of the important and challengeable Gram-negative microorganism and *P. aeruginosa* is replaced by *A. baumannii* in nosocomial infection. The emergence of their XDR type makes significant therapeutic problem especially in burned patients. According to our results and another study (26), colistin remains a more effective antibiotic for the treatment of infection caused by these XDR and/or MDR microorganisms. The spread of carbapenemase producer *A. baumannii*, as a second important pathogen that can make nosocomial infection in burned patients can be alarming for public health organization and health care systems.

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Conflict of interest

The authors declared no conflicts of interest.

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Characterization of Carbapenemases in Drug Resistance *A. baumannii*

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