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Genetic Characterization of Carbapenem Resistant *Acinetobacter baumannii* in Tertiary care settings of Lahore, Pakistan

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Abstract

Background: *Acinetobacter baumannii* is major cause of ventilator associated pneumoniae (VAP) as it is an opportunistic nosocomial organism. The current study was to find out the antibiotic resistance pattern of *Acinetobacter baumannii*, its phenotype and the genetic characterization of Metallo- β -Lactamase (MBL) genes that are responsible for carbapenem resistance.

Methods: One hundred and fifty Carbapenem resistant *Acinetobacter baumannii* (CRAB) specimens were isolated and PCR amplification of organism specific *bla*-OXA-51 gene was performed and antibiotic susceptibility was checked. Phenotypic susceptibility analysis was performed by Modified Hodge Test (MHT) and Imipenem-EDTA Double Disc Synergy Test (IMP-EDTA DDST). The carbapenemases and MBL producing genes were amplified by PCR.

Results: CRAB showed high resistance against piperacillin/tazobactam (99.3%), cefepime and ceftazidime (99.3% each), amikacin (91.3%), ciprofloxacin (96.7%) and levofloxacin (96.7%). Only one isolate showed resistance to colistin. The isolates positive for both MHT and DDST (n=70) were further characterized to detect metallo- β -lactamase genes. Molecular characterization revealed the presence of *bla*-OXA-51 gene in all tested isolates (100%) followed by *bla*-VIM 89%, *bla*-OXA-23 64%, respectively and so on. Few genes coexisted with each other including *bla* VIM, *bla* OXA 23, *bla* OXA 51 and *bla* NDM-1. None of the isolate was found positive for *bla*-IMP gene.

Conclusion: It is concluded that CRAB isolates exhibited a high rate of resistance towards antimicrobials because of the presence of drug hydrolyzing enzymes, carbapenemases and MBLs. This is among the rare study reported recently indicating CRAB isolates co-harboring many resistant genes are very difficult to treat. There is a dire need to develop novel antibiotics against resistant *A. baumannii* to minimize its prevalence. Moreover, it is recommended that colistin treatment in the clinical settings should be continuously monitored in order to prevent the development of resistance.

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Keywords:

Acinetobacter baumannii; Carbapenem resistance; Metallo-beta-Lactamases; Resistant genes

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Introduction

Acinetobacter species are universal, free-living bacilli found in sewage, soil, water and contaminated food [1]. *Acinetobacter* are gram negative coccobacilli, non-motile, glucose non-fermenter, non-fastidious, catalase positive and oxidase negative. The genus contains pathogenic species causing significant nosocomial infections [2]. *Acinetobacter baumannii* (*A. baumannii*) has the ability to survive in the low-moist environment and also to develop resistance against antimicrobials [3]. Scientific literature highlights the *Acinetobacter* as principal pathogen of hospital acquired infections however, fewer reports also document infections in general populations [4,5]. Prolonged hospitalization, venous catheters, nursing home residence, and an impaired immune system are some risk factors for *A. baumannii* infection [6]. It causes infections like pneumonia, wound sepsis, septicemia, urinary tract infections, endocarditis meningitis and endocarditis [7].

ESKAPE bacterial pathogens are the notorious critters known for their difficult to treat properties. They are *S. aureus*, *E. faecium*, *K. pneumoniae*, *A. baumannii*, *P. aeruginosa*, and *Enterobacteriaceae* family. These bacteria have a higher risk of developing antimicrobial resistance. Among this ESKAPE group, *A. baumannii* is the worst to treat due to its unique antibiotic resistance pattern [8]. *A. baumannii* isolates are becoming more resistant to many antimicrobials, including beta-lactams, aminoglycosides, and fluoroquinolones posing serious public health concerns [9]. Carbapenem is an antibiotic of class of β -lactam and is the recent resort therapeutics for the control of bacterial infections particularly, associated with *A. baumannii* [10]. In recent years, the reports suggested the increase in resistance of *A. baumannii* strains to carbapenem. Carbapenems are the drugs to treat these infections [11]. High mortality rate (50%) associated to carbapenem resistant bacteria has been reported [12]. Carbapenemase is one of the most frequent enzymes reported in carbapenem resistant *A. baumannii* [13]. Carbapenem resistance in *A. baumannii* is because of the production of class B, C and D carbapenemases [14,15]. Due to the production of OXA-type carbapenemase and Metallo β -Lactamases (MBLs) most *Acinetobacter* spp. develop the carbapenem resistance [16,17]. Among OXA-type carbapenemases *bla* OXA-23 like, *bla* OXA-40 like, *bla* OXA-58 like and *bla* OXA-51 like are commonly reported, while *bla* OXA-51 like β -Lactamase is used for species identification and is intrinsically resistant to *A. baumannii* [18,19]. Various metallo- β -lactamase genes such as IMP, VIM, SPM, KPC, OXA, and NDM have been reported in *A. baumannii* conferring AMR to carbapenems [15]. Antimicrobial resistance is a major threat in developing

countries leading to greater number of treatment failure in nosocomial infections [20]. By 2050, antimicrobial resistance is expected to cause 10 million deaths if the current situation continues [21].

Keeping in consideration the antimicrobial resistance and a raised level of nosocomial infections, in current study antibiotic resistance pattern, phenotypic characterization and Molecular identification, and sequencing of carbapenemase producing genes *bla* IMP, *bla* VIM, *bla* OXA 23 and *bla* OXA 21, *bla* OXA 51, *bla* OXA 58 and NDM-1 from CRAB were studied.

Methods

The study was jointly conducted by Institute of Microbiology, University of Veterinary and Animal Sciences, (UVAS) Lahore, Pakistan and Institute of molecular biology and biotechnology department of University of Lahore (UOL). Strains of *A. baumannii* were collected from various pathological laboratories and tertiary care hospitals in Lahore including Chughtai Institute of Pathology from April 2021 to April 2022. The design of the study was approved by the ethical review committee of University of Lahore (UOL).

Isolation of *A. baumannii*

A total of 150 Carbapenem resistant *A. baumannii* strains were isolated from various clinical specimens including blood, pus, wound swabs, sputum, tracheostomy swabs, tracheal secretions, cerebrospinal fluid, CVP tip, bronchoalveolar lavage, drain fluid, and tissues.

Identification of *A. baumannii*

At initial, isolates were cultured on blood agar medium and MacConkey agar and were incubated at 37°C for 24-48 hours. Different biochemical tests like Gram staining, catalase test, oxidase test and Analytical Profile Index (API) were performed for identification purpose [22, 24].

Antibiotic resistance patterns of Carbapenem resistant *Acinetobacter baumannii* determination

Antibiotic susceptibility testing of *A. baumannii* was checked by Kirby Bauer Disc Diffusion method as mentioned by CLSI [25]. Antibiotic discs contained cefepime (30 μ g), ceftazidime (30 μ g), Imipenem (10 μ g), Meropenem (10 μ g), Amikacin (30 μ g), Gentamicin (10 μ g), Tobramycin (10 μ g), Doxycycline (30 μ g), Ciprofloxacin (5 μ g), Levofloxacin (5 μ g), Trimethoprim sulphate (25 μ g). They were placed onto the MHA plate and incubated at 37°C for 24 hours. After 24 hours zones of inhibition around antibiotic discs were evaluated and declared as sensitive, intermediate and resistant as per the guidelines of clinical and laboratory institute (CLSI). The antibiotic susceptibility testing of

colistin was carried out by Minimum Inhibitory Concentration (MIC) by broth microdilution method as per Clinical Laboratory Standard Institute CLSI using Muller Hinton Broth [26].

Phenotypic assays for the determining β -lactamases (Carbapenemases & Metallo- β -Lactamase)

The detection of Carbapenemase producing *A. baumannii* was performed by Modified Hodge test as mentioned earlier by Amjad et al [27]. According to them ATCC *Escherichia coli* 25922 equivalents to 0.5 McFarland was swabbed on Muller Hinton Agar. In the center of plate disc of meropenem 10 μ g was placed. From the edge of the antibiotic disc to the edge of the plate the test organism was streaked and the incubated at 37°C for 24 hours. MHT positive test showed a clover leaf like indentation of *Escherichia coli* 25922 growing along the test organism growth streak within the disk diffusion zone after 24 hours.

The identification of Metallo- β -Lactamase producing *A. baumannii* was carried out by IPM-EDTA double disc synergy test according to a study done by Young et al [28]. Inoculum equivalent to 0.5 McFarland of *A. baumannii* was swabbed onto Muller Hinton Agar. A 10 μ g of imipenem disc and a sterile blank disc were placed 10 mm apart from edge to edge. As a control another imipenem disc was placed far. A volume of 10 μ L of 0.5M EDTA solution was put in to the blank disc. After overnight incubation, the established zone diameter difference of ≥ 7 mm between imipenem disk and imipenem plus EDTA will be interpreted as EDTA synergy positive.

Molecular identification of Carbapenem resistance determinants

By PCR using gene specific primers for *bla* OXA-23, *bla* OXA-51, *bla* OXA-24, *bla* OXA-58, *bla* IMP, *bla* VIM and *bla* NDM-1 gene as shown in table 1. The characterization of Carbapenemase & MBL encoding genes in *A. baumannii* was performed. The primers were synthesized on order by Macrogen, Inc., Seoul Korea. Multiplex PCR was performed for *bla*-OXA genes. PCR conditions for *bla*-OXA genes and MBL encoding genes (*bla* IMP, *bla* VIM and *bla* NDM-1) were set as usual.

Sequencing analysis

Sequencing of amplified samples (*bla* OXA-23, *bla* OXA-24, *bla* OXA51, *bla* OXA-58, *bla*NDM-1, *bla* VIM) was done by services of Advance Biosciences International (ABI) Company. Sequences were analyzed by Chromas software and then got confirmed with the help of Basic Local Alignment Search Tool (BLAST). Nucleotide sequences were submitted to the Gen Bank for their accession numbers.

Gene	Primer Sequence	Reference
<i>bla</i> OXA-23	Forward GATCGGATTGGAGAACCAGA Reverse ATTTCTGACCGCATTTCAT	[29]
<i>bla</i> OXA-51	Forward TAATGCTTTGATCGGCCTTG Reverse TGGATTGCACCTCATCTTGG	[29]
<i>bla</i> OXA-24	Forward GGTTAGTTGGCCCCCTAAA Reverse AGTTGAGCGAAAAGGGGATT	[29]
<i>bla</i> OXA-58	Forward AAGTATTGGGCTTGTGCTG Reverse CCCCTCTGCGCTCTACATAC	[29]
<i>bla</i> IMP	Forward GGAATAGAGTGGCTTAAATCTC Reverse GGTTTAAAYAAAACAACCACC	[30]
<i>bla</i> VIM	Forward GATGGTGTGGTTCGCATA Reverse CGAATGCGCAGCACCAG	[30]
<i>NDM-1</i>	Forward GGG CAG TCG CTT CCA ACG GT Reverse GTA GTG CTC AGT GTC GGC AT	[31]

Table 1: Gene and respective primers.

Statistical analysis

The clinical analysis of data was done by using SPSS, version 26. To find any possible association between demographic variables, specimens, isolates, antimicrobial resistance and genome of CRAB Chi-square or fisher exact test were used.

Results

Identification and isolation of *A. baumannii*

One hundred and fifty Carbapenem resistant *A. baumannii* (CRAB) exhibiting the following morphological characteristics: Gram-negative short coccobacillus on gram staining, non-motile, non-pigmented, mucoid and domed shaped colonies on chocolate and blood agar, non-lactose fermenting colonies over MacConkey, were isolated. They appeared to be catalase positive and oxidase negative. *A. baumannii* were also identified by API 20 E. Further confirmation was done by presence of *bla* OXA 51 gene in the isolates.

Prevalence of CRAB from different wards

Out of 150 isolates, highest prevalence was observed from Medical ICU (28%) followed by Surgical ICU (18%), Surgical wards (16.6%) Pediatric ICU (16.6%), Medical ward (15.3%), and lowest was observed from Pediatric ward (5.3%).

Prevalence of CRAB from different specimens

The maximum percentage of CRAB were isolated from tracheal secretions (24.7%), followed by pus (20.00%), blood (16.00%), wound swab (14.7%), sputum (10.00%), tracheostomy swab (4.70%), CSF (4.00%), alveolar

lavage (2.00%), tissue samples (1.30%) and minimum percentage from bronchial washings (0.70%), pleural fluid (0.70%), drain fluid (0.70%) and CVP tip(0.70%) respectively Figure 1.

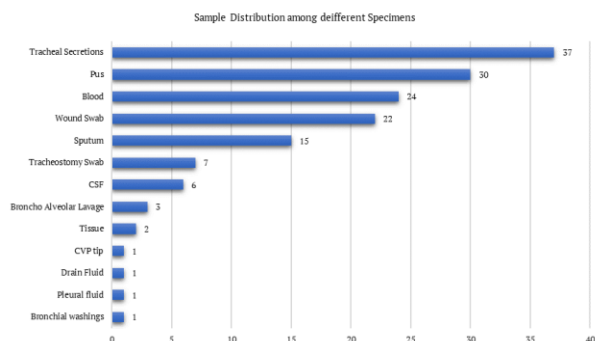


Figure 1: Frequency of *A. baumannii* from various types of clinical samples from tertiary care hospital.

Prevalence of CRAB from different age groups

The maximum percentage of CRAB isolates were obtained from the patients of 21 to 40 years of age group (34.00%) followed by 41 to 60 years of age group (28.00%) and the minimum percentage of isolates were isolated from less than 01 years (11.33%) and 01 to 20 years age group (11.33%) respectively.

Frequency of CRAB from gender

The maximum percentage of CRAB were isolated from males (60.66%) as compared to females (39.33%) respectively.

Antibiotic resistance pattern

Out of 150 CRAB isolates, highest level of resistance was found in piperacillin-tazobactam (99.30%), cefepime (99.30%), ceftazidime (99.30%), Ciprofloxacin (96.7%), levofloxacin (96.7%), amikacin (91.3%), gentamicin (88.7%), tobramycin (62%), trimethoprim-sulfamethoxazole (82.7%) and doxycycline (67.3%). Amongst all antibiotics only colistin (0.7%) showed better activity against CRAB respectively (Figure 2). All clinical isolates were found resistant to most antibiotic groups and were considered multi drug resistant.

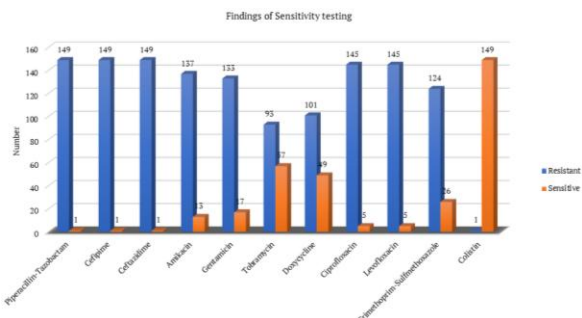


Figure 2: Frequency of antibiotic susceptibility testing of *A. baumannii* isolates

Phenotypic characterization of metallo- β-Lactamase production and carbapenemase

All the isolates (n=150) were characterized phenotypically for production of Carbapenemase by Modified Hodge Test (MHT). The maximum number of isolates 143 (95%) were positive for Carbapenemase production. By double disc synergy test (DDST) the detection of metallo-β-Lactamase producing *A. baumannii* was. Out of 150 isolates, 74 (49%) were positive for metallo-β-Lactamase production.

Molecular characterization of Metallo β-Lactamase gene

The isolates that were positive for Modified Hodge Test as well as IPM-EDTA double disc synergy test (n=70) were proceeded further for the genetic characterization of carbapenem resistant genes. The MBL genes *bla* NDM-1, *bla* OXA-23, *bla* OXA-24, *bla* OXA-51, *bla* OXA-58, *bla* VIM and *bla* IMP were amplified by using gene specific primers as seen in Figure 3.

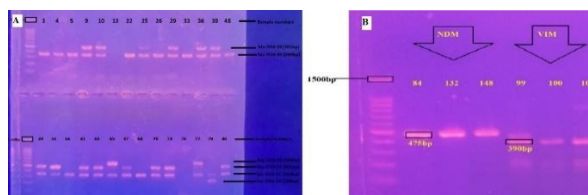


Figure 3: Molecular Characterization of Metallo-Beta-Lactamase genes in *A. baumannii* using gene specific primers. (A) Amplification of *bla*-OXA genes by multiplex PCR, (B) PCR Amplification of *bla*-NDM-1 and *bla*-VIM genes.

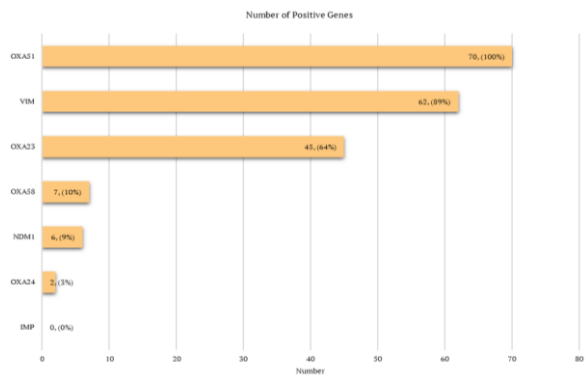


Figure 4: Frequency of Metallo-β-Lactamase genes, characterized from *A. baumannii* isolates in this study.

bla OXA 51 gene had the maximum rate of amplification and exhibited positive amplification for all 70 samples followed by *bla* VIM (62/70) 89%, *bla* OXA-23 (45/70) 64%, *bla* OXA-58 (7/70) 10%, *bla* NDM-1 (6/70) 8.5% and *bla* OXA-24 (2/70) 2.8%. None of the isolate was positive for *bla* IMP gene as shown in Figure 4. Furthermore, these genes coexisted with each other in combinations as well. The most common combination was a 3 gene combination in 38(54%) isolates harboring *bla* VIM, *bla* OXA 23 and *bla* OXA 51

genes. A combination of 4 genes occurred simultaneously in 3 (4%) isolates having *bla* VIM, *bla* OXA 23, *bla* OXA 51 and *bla* NDM-1 as shown in Figure 5.

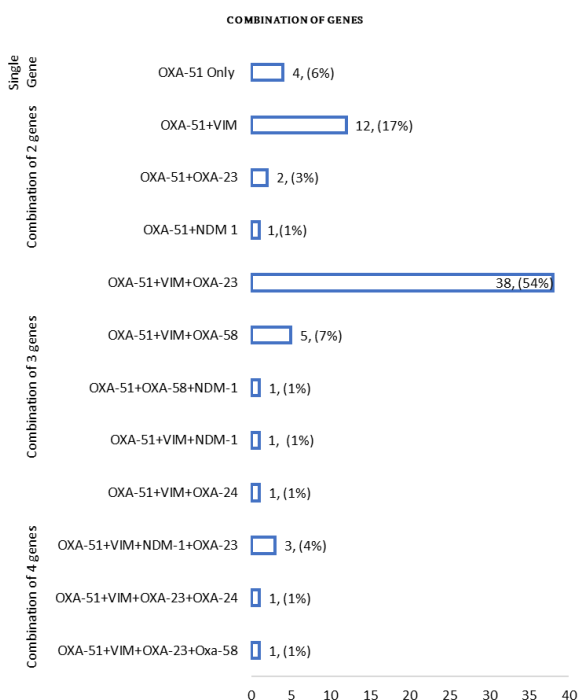


Figure 5: Distribution of multiple Metallo-β-Lactamase genes characterized from *A. baumannii* isolates in this study.

Discussion

β-Lactam antibiotics have been among the most effective medications used to treat bacterial infections in humans over past 60 years. *Acinetobacter* has developed as a significant disease class, posing ongoing hazards and problems to the global health care system [32-34]. In *A. baumannii* resistance to β-Lactams specially carbapenems and cephalosporins has been excessively seen globally and this is also the case in Pakistan.

The present study showed a high resistance pattern of different antibiotics towards carbapenem resistant *A. baumannii* isolated from various clinical specimens which revealed that our results are coinciding with the previous study in Egypt (98%), Greece (96.9%) and India (100%) [35-37]. The present study illustrates the prevalence of MBL producing *A. baumannii* 74/150 (49.3%) that is consistent with previous studies from Pakistan as well as from other countries. A similar study from Lahore, Pakistan reported that 63/112 (56.25%) of *A. baumannii* were MBL producers [38]. A study conducted in Iran showed a high prevalence 86/108 (86.86%) of MBL producing carbapenem resistant *A. baumannii* [39]. In Asian countries the

incidence of MBL producing *A. baumannii* have been reported as 77% in 2019 in China [40]

In the present study *bla* OXA-51 was most prevailing 70/70(100%), followed by *bla*-VIM 62/70 (88%), *bla*-OXA 23 45/70 (64%), *bla*-OXA 58 7/70 (10%), *bla* NDM-1 6/70 (8.5%) and *bla* OXA-24 2/70 (2.8%). In this study None of isolate was positive for *bla* IMP gene. A study in Spain showed a 100% prevalence of *bla* OXA-51 in *A. baumannii* which is similar to our study [41].

In our study, these carbapenem resistant genes coexisted with each other in combinations as well. The most common combination was a 3 gene combination in 38(54%) isolates harboring *bla* VIM, *bla* OXA 23 and *bla* OXA 51 genes together. Majority of *A. baumannii* isolates were co-producers of class B and D carbapenemases harboring *bla* IMP, *bla* OXA 23 and *bla* NDM-1 in a study conducted in India [42]. Similar studies were previously reported in different countries [43-45]. In the current study a rise in resistance is seen among the *A. baumannii* isolates that is a major public health concern. As, the majority of isolates showed susceptibility to colistin indicating that colistin could play possible role in the control and treatment of carbapenem resistant *A. baumannii* [46].

This study possesses several shortcomings as well. The study was only done in Lahore, and the sample size was quite small. It is highly recommended that this research should be conducted on a broader scale and in other clinical settings throughout the country to obtain more of a valid susceptibility pattern against MDR and XDR *A. baumannii*, which will assist in controlling the transmission of infections that is caused by this deadly organism and will also be helpful for managing infectious diseases in a better way.

It is concluded that CRAB is major threat in hospitals of Pakistan and isolates exhibited a high rate of resistance to antimicrobials because of the presence of drug hydrolyzing enzymes, carbapenemases and MBLs. We also found out, *bla*OXA-24, *bla*OXA-58, *bla* OXA-51, *bla* OXA-23, *bla* NDM-1 & *bla* VIM, genes had not been widely reported in Pakistan. This study further concluded, CRAB isolates, co-harboring many resistant genes are very difficult to treat. There is a dire need to invent new antibiotics against resistant *A. baumannii* for reduction of its prevalence. Moreover, in order to prevent the development of resistance it is highly recommended that colistin treatment in the clinical settings should be continuously monitored. Understanding of genetic characterization and molecular epidemiology of *A. baumannii* to devise strategies for prevention and control of MDR infections are also necessary.

Author Contributions

Principal investigator, collection of data and Manuscript writing by Saadia Ijaz. Study conception and design by Farheen Ansari. Technical supervision and interpretation of results by Prof. Muhammad Nawaz. Sample and data collection by Karam Rasool. Technical guidance and final review by Prof. Aftab Anjum.

Conflicts of interest

The authors declare no conflict of interest.

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