



# Molecular Characterization Identifies Upstream Presence of *ISAb<sub>a</sub>1* to OXA Carbapenemase Genes in Carbapenem-Resistant *Acinetobacter baumannii*

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## Abstract

**Background** Evaluating the expression pattern of oxacillinases (OXA) carbapenemases is essential to understand the prevalence and spread of carbapenem resistance *Acinetobacter baumannii*.

**Objectives** The aim of the study is to evaluate the presence of OXA carbapenemase genes and *ISAb<sub>a</sub>1* upstream to these genes in carbapenem-resistant *A. baumannii* clinical isolates.

**Materials and Methods** *A. baumannii* isolated from clinical samples were phenotypically identified and antibiotics sensitivity was performed. Multiplex polymerase chain reaction (PCR) was used to detect OXA51-like gene, OXA carbapenemases genes (OXA-23-like, OXA-24-like, and OXA-58-like), and *ISAb<sub>a</sub>1* in carbapenem-resistant isolates.

**Results** Out of 55 *Acinetobacter* isolates, 54 were confirmed as *A. baumannii* by PCR. *Bla*<sub>OXA-23</sub>-like gene was observed in 51 isolates of *A. baumannii* and none of the isolates showed the presence of *bla*<sub>OXA-24</sub>-like and *bla*<sub>OXA-58</sub>-like genes. Presence of *ISAb<sub>a</sub>1* upstream to OXA-23-like gene, OXA-51-like gene, and both OXA-51-like/OXA-23-like genes was observed in 51, 7, and 4 *A. baumannii* isolates, respectively.

**Conclusion** The genetic pattern of carbapenem-resistant *A. baumannii* isolated in this study was unique, which should be factored for clinical protocols to manage infections caused by emerging resistant strains of *A. baumannii*.

## Keywords

- ▶ CRAB
- ▶ *ISAb<sub>a</sub>1*
- ▶ OXA

## Introduction

Multidrug-resistant (MDR) bacterial infections are responsible for increased mortality and morbidity in intensive care units (ICUs). *Acinetobacter baumannii* is one of the MDR

pathogens which poses a significant risk to public health due to its high levels of resistance to currently used antibiotics.<sup>1</sup> Despite improvements in hospital infection control and antimicrobial stewardship, infections from *A. baumannii* continue to rise.<sup>2 3</sup> The ubiquitous nature of *Acinetobacter*

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poses a significant challenge to eradicate it from the hospital environments.<sup>3</sup> Due to the prevalence of variable antimicrobial susceptibility patterns in different hospitals, facility-specific surveillance studies are valuable in deciding tailored therapy to effectively control infections associated with *A. baumannii*. Further it is necessary to continuously isolate and monitor resistance patterns for adopting optimal clinical antimicrobial therapy protocols.

*A. baumannii* has acquired many genetic characters by horizontal gene transfer, which has led to increase in its virulence and resistance and making it a nosocomial pathogen of serious concern. *Acinetobacter* spp. are opportunistic pathogens that affect critical care units, where they are associated with pneumonia consequence to endotracheal tube intubations or tracheostomy procedures. They are also associated with bacteraemia's urinary tract infection and wound infections.<sup>3</sup> Multiple mechanisms of resistance such as  $\beta$  lactamases, modifications in porin proteins, and efflux pumps are reported in *Acinetobacter* spp. Enzymatic degradation by  $\beta$  lactamases is reported to be the most prevalent mechanism of resistance observed. Resistance to aminoglycosides and quinolones is reported to be due to mutations in *gyrA* and *parC* genes.<sup>3</sup> Carbapenems are the drugs of choice in the treatment of antibiotic-resistant Gram-negative organisms, however, increasing incidence of resistance to carbapenems is also reported.<sup>4</sup> Infections from carbapenem-resistant *Acinetobacter baumannii* (CRAB) are increasing globally and difficult to treat.<sup>5</sup>

*A. baumannii* contains intrinsic oxacillinases-51 (OXA-51) serine type oxacillinase which imparts natural resistance to  $\beta$ -lactams. Carbapenem resistance is reported to be due to serine oxacillinases (Ambler class D) and metallo  $\beta$  lactamases (MBLs) (Ambler class B). In the class D OXA-type beta lactamases, the five main groups (OXA-23-Like, OXA 24/40 like, OXA-58, OXA-143 like, and OXA-235 like) are identified. Resistance to carbapenem is reported to increase significantly when OXA genes are juxtaposed to insertion elements (IS*Aba1*) with promoter functions.<sup>6</sup> However, the prevalence of these resistance factors in *A. baumannii* isolates from all geographical regions is not known. Hence this study was designed to evaluate the distribution of resistance genes (OXA-51-like, OXA-58-like, OXA-23-like, and OXA 24-like) and associated insertion elements (IS*Aba1*) among clinical isolates of CRAB in a tertiary care hospital facility.

## Materials and Methods

### Study Design and Ethical Considerations

This study was a prospective observational study, which was reviewed and approved by the Institutional Ethical Committee. The study was conducted at the Department of Microbiology of a tertiary care hospital.

### Inclusion Criteria

Carbapenem-resistant *A. baumannii* isolates obtained from various samples (blood, urine, pus, respiratory tract, CSF, etc.) of patients admitted in the various wards and ICUs of hospital were collected between November 2018 and De-

cember 2019. A total of 55 clinical isolates of *A. baumannii* were included in the study.

### Exclusion Criteria

Carbapenem sensitive *A. baumannii* isolates obtained from various samples of patients were excluded.

### *Acinetobacter Baumannii* Identification

*A. baumannii* isolates were identified by phenotypic test scheme including nonlactose fermenting colonies on MacConkey agar and preliminary tests (Gram staining, motility, oxidase, and growth at 44°C). All isolates after phenotypic identification were genotypically confirmed by detection of intrinsic *bla*<sub>OXA-51</sub>-like gene by PCR.<sup>2,3</sup>

### Antibiotic Susceptibility Testing

Antibiotic susceptibility testing (ABST) was performed on all *A. baumannii* isolates by Kirby-Bauer disk diffusion method.<sup>3</sup> The drugs for disk diffusion testing were in the following concentrations: amikacin (30  $\mu$ g), gentamicin (10  $\mu$ g), cotrimoxazole (25  $\mu$ g), cefoxitin (30  $\mu$ g), cefotaxime (30  $\mu$ g), ceftazidime (30  $\mu$ g), ceftriaxone (30  $\mu$ g), imipenem (10  $\mu$ g), meropenem (10  $\mu$ g), and piperacillin-tazobactam (100/10  $\mu$ g). Interpretation of zone of inhibition was done as per the Clinical and Laboratory Standards Institute guidelines.<sup>7</sup> MBL production was detected phenotypically using imipenem and imipenem-ethylenediaminetetraacetic acid (EDTA) disk. Zone of inhibition  $\geq$  7 mm in imipenem-EDTA disk in comparison to imipenem alone was considered positive for MBL production.<sup>8</sup> Multidrug resistance (MDR) was considered if the isolates were resistant to at least one agent from three or more antimicrobial categories.<sup>9</sup>

### Multiplex PCR

Bacterial genomic DNA extraction was performed using DNeasy Tissue kit, (QIAGEN) from 55 *Acinetobacter* isolates which showed resistance to carbapenem. Multiplex PCR was performed on all isolates to detect the presence of *bla*<sub>OXA-51</sub>-like, *bla*<sub>OXA-23</sub>-like, *bla*<sub>OXA-24</sub>-like, and *bla*<sub>OXA-58</sub>-like genes, and IS*Aba1*. The primer sequences used for amplification, as shown in ► **Supplementary Table 1** (available online only), are previously reported.<sup>10,11</sup> The amplification conditions used were as follows: initial denaturation at 95°C for 5 minutes, 35 cycles of 95°C for 30 seconds, 50°C for 30 seconds, and 72°C for 30 seconds, and a final elongation cycle at 72°C for 7 minutes. The presence of IS*Aba1* and OXA target genes was further investigated for the upstream presence of IS*Aba1* element to each OXA gene. To check this downstream primer of each OXA gene was combined with the forward primer of IS*Aba1* in a separate multiplex PCR.

## Results

During the study period, 47 patients infected with CRAB were identified and 55 isolates of *A. baumannii* were obtained from these patients.

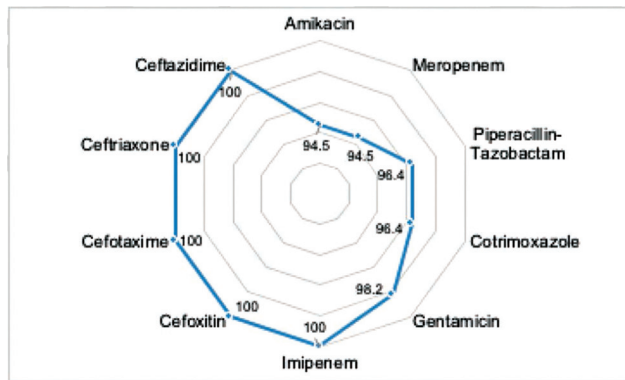
**Antibiotic Sensitivity Testing**

ABST revealed that all the isolates tested were resistant to imipenem, ceftazidime, ceftriaxone, cefotaxime, and ceftioxin. A higher degree of resistance to other antibiotics (gentamicin, cotrimoxazole, piperacillin-tazobactam, meropenem, and amikacin) was also observed (►Fig. 1). Most of the isolates (49/55, 89%) were observed to be MBL producers (►Fig. 1) and all the isolates were multidrug resistant (MDR).

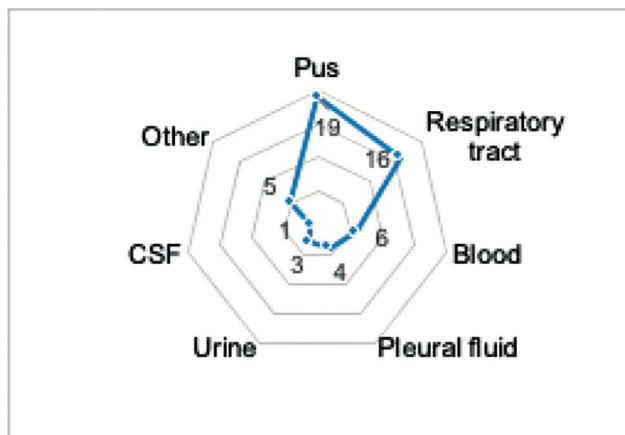
**Multiplex PCR**

Out of 55 *Acinetobacter* isolates included in the study after the phenotypic test, 54 were confirmed as *A. baumannii* by PCR based on the presence of *bla*<sub>OXA-51</sub> gene, which is intrinsic to this species. Of the 54 isolates, 19 were isolated from pus samples, 16 were isolated from the respiratory tract, 6 were isolated from blood samples, 4 were isolated from pleural fluid, 3 were isolated from urine samples, 1 was isolated from cerebrospinal fluid, and 5 were isolated from other clinical samples (►Fig. 2).

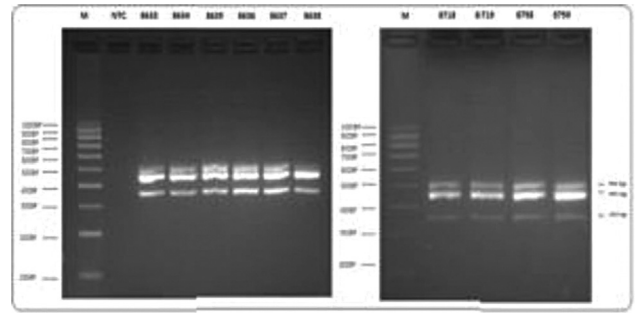
Agarose gel electrophoresis (►Fig. 3) followed by multiplex PCR analysis for the presence of three carbapenemase genes (*bla*<sub>OXA-23</sub>-like, *bla*<sub>OXA-24</sub>-like, *bla*<sub>OXA-58</sub>-like) showed the presence of *bla*<sub>OXA-23</sub>-like genes in 51 isolates of *A.*



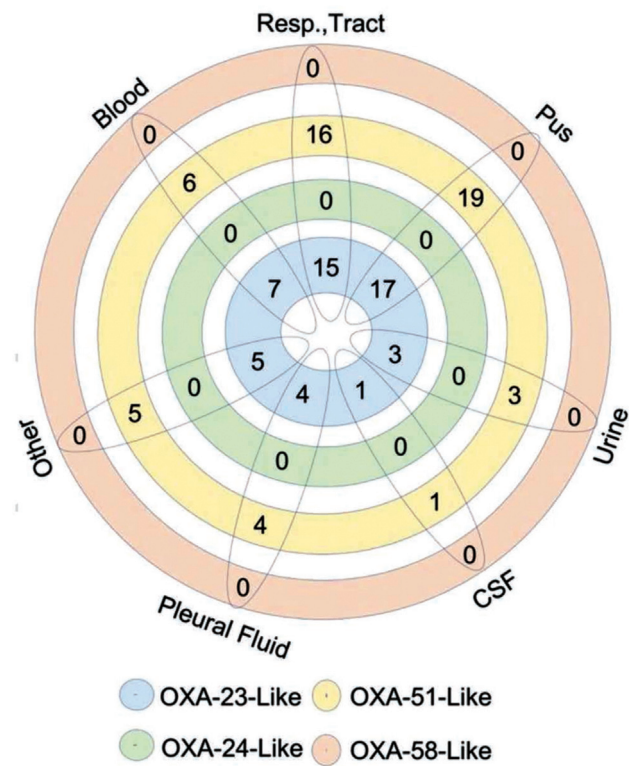
**Fig. 1** Percentage of *Acinetobacter baumannii* isolates resistant to antibiotics tested.



**Fig. 2** Number of various samples used for the isolation of *Acinetobacter baumannii*.



**Fig. 3** Detection of OXA-51-like, OXA-23-like genes and ISAb<sub>a</sub>1 by multiplex PCR. A representative image of agarose gel electrophoresis showing detection of genes encoding OXA-carbapenemases and ISAb<sub>a</sub>1. PCR, polymerase chain reaction.

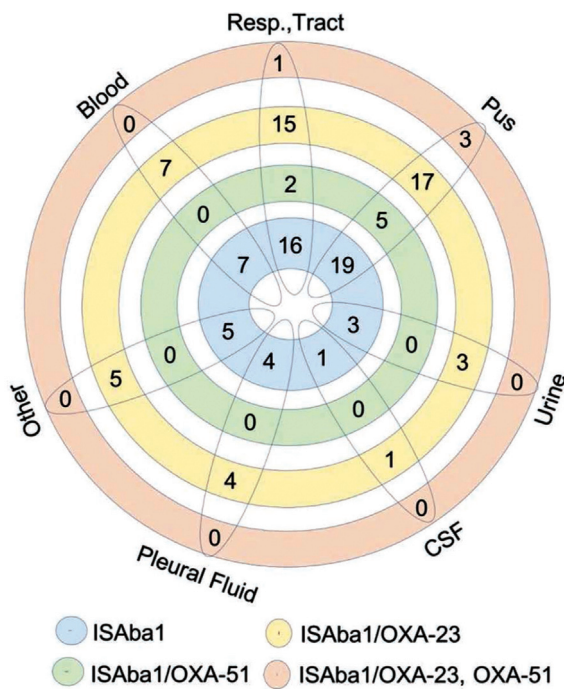


**Fig. 4** Prevalence of OXA-carbapenemase genes in *Acinetobacter baumannii* isolates from various clinical samples.

*baumannii*. However, none of the isolates showed the presence of *bla*<sub>OXA-24</sub>-like or *bla*<sub>OXA-58</sub>-like genes (►Fig. 4).

Presence of ISAb<sub>a</sub>1 sequence was observed in all 54 *A. baumannii* isolates. Fifty-one isolates showed the presence of ISAb<sub>a</sub>1 upstream to OXA-23-like gene, while seven isolates detected ISAb<sub>a</sub>1 upstream to OXA-51-like gene. Four *A. baumannii* isolates carried ISAb<sub>a</sub>1 upstream to both OXA-51-like and OXA-23-like genes and interestingly three of these isolates were from pus samples and one isolate was from ET secretion (►Fig. 5).

In one isolate, OXA-51-like gene was not detected but the presence of OXA-23 was observed. This isolate showed



**Fig. 5** Prevalence and upstream presence of ISAb<sub>a</sub>1 to Oxa-carbapenemase genes from various clinical samples.

ISAb<sub>a</sub>1 upstream to OXA-23-like gene and was further investigated by the Vitek-2 method. The organism was identified as *A. baumannii* complex with excellent identification features by this method (► Fig. 5).

## Discussion

Development of multidrug resistance in *A. baumannii* together with increasing resistance to carbapenems poses a major challenge in the treatment of patients infected with this microbe. Carbapenem resistance in *A. baumannii* is reported to be associated with Class D OXA carbapenemase (including intrinsic chromosomal *bla*<sub>OXA-51</sub>) and Class B metallo-β-lactamase production.<sup>12</sup> The expression of these resistance factors in addition to helping identification of *A. baumannii* species also contributes to increase in the minimum inhibitory concentration of carbapenem.<sup>13</sup> In this study a higher prevalence (94.4%) of *bla*<sub>OXA-23</sub>-like carrying genes was observed among the *A. baumannii* isolates. The OXA-23-like gene which is responsible for the development of resistance is present on the bacterial plasmid and can be transferred in conjugation between the *A. baumannii* strains facilitating rapid spread of resistance in any geographical regions.<sup>14</sup> In Asia<sup>15</sup> and various parts of India<sup>16,17</sup> the *bla*<sub>OXA-23</sub>-like is the dominant acquired gene detected among the CRAB. Hence our observations are consistent with these previous reports.

The *bla*<sub>OXA-24</sub>-like and *bla*<sub>OXA-58</sub>-like genes are associated with nosocomial outbreaks of MDR *A. baumannii*. Specifically, OXA-58 is widely reported from European Union and United States, while *bla*<sub>OXA-24</sub>-like is reported from United States, Spain, and China.<sup>18,19</sup> However, both *bla*<sub>OXA-24</sub>-like and *bla*<sub>OXA-58</sub>-like genes were not observed in any of the

isolates of *A. baumannii* in this study. Concurrent to our observations a previous study involving five centers across India also did not detect these genes.<sup>20</sup> In contrast a study from South India reported OXA-24 like and OXA-58-like genes in 22.9 and 4.2% *A. baumannii* isolates, respectively.<sup>21</sup> The regional selective identification of the resistance gene, although beyond the scope of this study to explain, merits detailed evaluation in future. Expression of *bla*<sub>OXA</sub>-like genes is reported to play a major role in the development of carbapenem resistance, which can be potentiated by insertion sequences such as ISAb<sub>a</sub>1, ISAb<sub>a</sub>2, ISAb<sub>a</sub>3, ISAb<sub>a</sub>4, and ISAb<sub>a</sub>10.<sup>21</sup> In this study presence of ISAb<sub>a</sub>1 sequence was observed in all *A. baumannii* isolates. The presence of ISAb<sub>a</sub>1 upstream to some OXA genes is associated with overexpression of these genes.<sup>21</sup> Most of the isolates in this study had upstream detection of ISAb<sub>a</sub>1 to OXA-23-like gene with few isolates having ISAb<sub>a</sub>1 either upstream to OXA-51-like gene or both OXA-23-like and OXA-51-like gene. Previously upstream presence of ISAb<sub>a</sub>1 to OXA-23-like in *Acinetobacter nosocomialis* was reported from Latin America and Iran.<sup>22,23</sup> However, to the best of our knowledge ISAb<sub>a</sub>1 expression upstream to OXA-23-like without OXA-51-like in *A. baumannii*/*A. nosocomialis* isolates is rarely reported from India. The *A. baumannii* isolate which did not express OXA-51-like gene but had ISAb<sub>a</sub>1 upstream to OXA-23-like gene was identified as *A. baumannii* complex by Vitek-2 method. The variable observation of the OXA-23-like and OXA-51-like genes in *A. baumannii* isolates paraps suggests intraspecies horizontal transfer of oxacillinase genes.

This study observed 100% resistance to ceftazidime, ceftriaxone, cefotaxime, and ceftioxin. The resistance pattern in this study is marginally higher than that reported by a previous study,<sup>24</sup> which perhaps suggests the progressive nature of resistance development by *A. baumannii*. Also, a majority (89%) of the *A. baumannii* isolates were MBL producers, indicating coexistence of both MBL and OXA-like genes in the evolving resistant strains of *A. baumannii*. This finding is a significant increase from 14% MBL as seen in a previous study by the same authors.<sup>25</sup> However, the characterization of the MBL *A. baumannii* isolates in this study was performed based on phenotypic and not genotypic methods, which remains the limitation of this study.

## Conclusion

This study reports *bla*<sub>OXA-23</sub>-like gene as the predominant cause of carbapenem resistance in *A. baumannii*. The upstream presence of ISAb<sub>a</sub>1 over *bla*<sub>OXA-23</sub>-like and *bla*<sub>OXA51</sub>-like genes, besides enhancing the development of carbapenem resistance also influences the intrastain mobility of resistant genes. The genetic pattern of carbapenem-resistant *A. baumannii* isolates observed in this study should be factored for clinical protocols to manage infections caused by emerging resistant strains.

## Conflict of Interest

None declared.

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