A Comprehensive Study of the Relationship between the Production of $\beta$-Lactamase Enzymes and Iron/Siderophore Uptake Regulatory Genes in Clinical Isolates of Acinetobacter baumannii

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Background. The iron/siderophore uptake system (IUS) involved in the Acinetobacter baumannii pathogenicity. However, IUS’s role in antibiotic resistance and the production of $\beta$-lactamase enzymes of A. baumannii are unclear. This study aimed to investigate the relationship between the production of $\beta$-lactamase enzymes and IUS regulatory genes in clinical isolates of A. baumannii. Methods. A. baumannii isolates were collected from clinical isolates using biochemical tests. The antibiotic resistance patterns and $\beta$-lactamase-producing strains were identified using the disk diffusion method (DDM). Also, IUS genes were detected by the polymerase chain reaction (PCR) method. Results. Seventy-two (72) A. baumannii isolates were collected from a different clinical specimen. Gentamicin-resistant strains (43%) had the highest frequency, and aztreonam-resistant strains (12.5%) had the lowest frequency. Also, the distribution of AmpC and MBL producing isolates were 27.7% and 35%, respectively. Moreover, the frequencies of basD, bauA, pld, paaE, entA, feoB, hemO, and tonB genes were as follows: 12.5%, 15.2%, 11.1%, 15.2%, 19.4%, 16.6%, 23.6%, and 6.9%. Further, a strong correlation was observed between the abundance of $\beta$-lactamase-producing strains and IUS genes. Conclusions. Based on our knowledge from this study, the association between $\beta$-lactamase production and IUS genes in A. baumannii plays an essential role in the emergence of drug-resistant strains.

1. Introduction

Acinetobacter baumannii (A. baumannii) has become an increasingly important human pathogen connected with infections acquired in hospitals [1]. It has become more challenging to treat A. baumannii infections in the hospitalize portions because of their resistance to significant groups of antimicrobial agents such as $\beta$-lactams [2]. Various resistance mechanisms make A. baumannii a successful pathogen of the 21st century [3]. Extended-spectrum $\beta$-lactamas (ESBL) belong to the class A $\beta$-lactamas. Metallo $\beta$-lactamas (MBL), which have metal ions governing the active site, belong to the class B $\beta$-lactamas [4–6]. Class C $\beta$-lactamas are acinetobacter-derived cephalosporinases or the chromosomally encoded AmpC enzymes. Also, in class D $\beta$-lactamas, oxacillinases are involved in resistance to carbapenems. However, there is now a crucial need for new antibacterial strategies due to the spread of pathogen populations [7–9].

In A. baumannii, virulence factors and biofilm formation play a crucial role in disease progression in an infected host [10–12]. Another virulence factor studied in detail is A. baumannii to survive in an iron-limited environment, like the human host [13]. In response to low iron, A. baumannii produces the siderophore acinetobactin to acquire this essential micronutrient. A better understanding of the acinetobactin system could lead to new antimicrobial drugs to treat infections caused by multidrug-resistant (MDR) A. baumannii infections [10]. However, A. baumannii synthesizes siderophores which are relatively low-molecular-weight agents capable of
converting polymeric ferric oxy-hydroxides to soluble iron chelates under low iron stress. The ability of bacteria to assimilate iron is related to invasiveness [3, 14]. However, over the past decade, a need for new antimicrobial compounds has arisen with the advent of MDR bacteria, such as A. baumannii, plaguing our healthcare systems [13, 15]. Since iron is essential for pathogenic bacteria to cause a successful infection, siderophore-mediated iron acquisition systems have been exploited as potential therapeutic targets [13]. One strategy utilizes the siderophore uptake machinery to deliver an antibiotic conjugated to the siderophore [2]. Nevertheless, in this study, to determine the relationship between β-lactamase enzymes and iron uptake pathways, we investigated the iron uptake system’s role in the emergence of resistant strains.

2. Materials and Methods

2.1. Design of Study and Sample Collections. A cross-sectional study was performed on patients admitted to Hamedan’s Hospital, attached to Iran’s Hamedan Medical School. The study was conducted between November 2018 and April 2020. Six hundred ninety clinical samples were collected, including wound swab, urine, blood, ascetic fluid, endotracheal fluid, bronchoalveolar lavage—inclusion criteria for patients presenting with signs of bacterial infection from bacterial culture examination. However, patients without bacterial infection symptoms, incomplete medical record data, or those who underwent nonstandard bacterial culture screening procedures were excluded from the study. After considering the inclusion and exclusion criteria, samples are used for culture and further biochemical investigations. Samples are inoculated in blood agar, MacConkey agar, and nutrient agar.

2.2. Antibiotic Susceptibility Test (AST). The AST was detected by disk diffusion methods (DDM). Twelve antibiotic discs (trimethoprim/sulfamethoxazole, pipercillin/tazobactam, amikacin, ciprofloxacin, ofloxacin, gentamicin, cefotaxin, imipenem, cotrimoxazole, cefepime, ertapenem, and aztreonam) all owned by Mast, UK, were used to evaluate the AST. Interpretation of antimicrobial sensitive or resistance zone was done by CLSI 2019 Guidelines [16].

2.3. Detection of AmpC and MBL Producing Strains. The AmpC and MBL producing A. baumannii were detected by the DDM [7].

2.4. Detection of Biofilm-Forming Strains. Microtiter dish assay was used to identify the biofilm-forming A. baumannii [17].

2.5. Total DNA Extraction. Genomic DNA was extracted using the method of Tahmasebi et al. [7]. By loading of 5 μl of the extracted DNA in 1% agarose gel, the extracted quality was checked.

2.6. Determination of Iron/Siderophore Mediate Gene. According to Abdi et al. [3], the PCR method was used for the detection of the IUS gene. For partial gene, the PCR conditions used to amplify the iron/siderophore mediate gene were the same as described in Table 1. Moreover, PCR was performed using a C1001 Bio-Rad thermal cycler. Finally, 5 μl of amplified product is electrophoresed on 1.5% agarose gel at a constant of 80 V for 60 min [3].

2.7. Statistical Analysis. In this study, GraphPad Prism software 8 (GraphPad Software, Inc., San Diego, CA) was used to analyze the data. Chi-square, two-way ANOVA, Tokay, and t-test were also used to examine the relationship between different variables. A p value of 0.05, 0.01, and 0.001 was considered as the level of significance.

3. Results

Out of 690 isolates of different clinical isolates, 72 isolates of A. baumannii were collected. According to Table 2, most samples were obtained from wound 23 samples (31.9%), followed by urine 21 samples (29.1%), blood 18 (25%), and body fluids 10 (13.8%).

3.1. Antibiotic Susceptable Profile. Table 2 shows the percentage of sensitivity, intermediate, and resistance of the tested A. baumannii isolates. More than 30% of the isolates were resistant to gentamycin (43.0%), ofloxacin (30.5%), and ciprofloxacin (40.2%). Only nine isolates (12.5%) were resistant to aztreonam. Further, 22 isolates (30.5%) and 13 isolates (18.0%) were considered MDR and XDR, respectively.

3.2. Prevalence of MBL and AmpC Producing Strains. Table 2 and Figure 1 show that 27.7% and 36.1% of A. baumannii isolates were AmpC and MBL producers, respectively. Also, coexistence of AmpC and MBL enzymes was observed in 12 isolates (16.6%).

3.3. Prevalence of Biofilm-Forming A. baumannii. The frequency of biofilm-producing A. baumannii is shown in Table 2. Among the 72 isolates of A. baumannii, 39 biofilm-forming isolates (54.1%) and 33 nonbiofilm-forming isolates (45.8%) were detected.

3.4. Prevalence of Iron/Siderophore Mediate Gene. The distribution of siderophore genes is illustrated in Table 2 and Figure 2. Among 72 isolates of A. baumannii, 9 isolates (12.5%) carry BasD gene, 11 isolates (15.2%) carry BauA and paaE genes, 8 isolates (11.1%) carry Pld gene, 12 isolates (16.6%) carry feoB and entA genes, 17 isolates (23.6%) carry hemO gene, and 5 isolates (6.9%) carry tonB gene.

3.5. Statistical Analysis. Based on Figure 3, a statistical association was detected when the IUS had compared virulence factors. Using two-way ANOVA tests, t-test, and
Table 1: Oligonucleotide sequences used in this study and thermal cycling conditions.

| Gene | Sequence of primers | Thermal cycles | Product size (bp) | Ref. |
|------|---------------------|----------------|------------------|------|
| basD | F: CTCTTGCACTGGAACACAC<br>R: CCAACGAGACCGCCTATGTT | 95°C/5 min; (95°C/1 min, 57°C/60 sec, 72°C/45 sec) X30; 72°C/5 min | 868 [2] | |
| bauA | F: TGGCAAGGTTAAAAATGCAAG<br>R: GCCGCTATGGCCTCAACCTG | 95°C/5 min; (95°C/1 min, 58°C/45 sec, 72°C/45 sec) X35; 72°C/5 min | 224 [2] | |
| pld | F: CCGTCAGATCTGCTGCTTGG<br>R: CTGACGCTACCTGAGGTTT | 95°C/5 min; (95°C/1 min, 57°C/1 min, 72°C/45 sec) X30; 72°C/5 min | 662 [2] | |
| paaE | F: CTATTTAGGGTTGTCGCG<br>R: CCTTACAACGACAGGTCGCA | 95°C/5 min; (95°C/1 min, 59°C/1 min, 72°C/45 sec) X30; 72°C/5 min | 593 [2] | |
| feoB | F: AAGTCGCCAACTATGCGGTGT<br>R: AAGGCGCTGCCCATGCAAAAAC | 95°C/10 min; (95°C/1 min, 57°C/30 sec, 72°C/1 min) X30; 72°C/5 min | 636 [3] | |
| hemO | F: TCGTGGCCGCTCAAAACAAGCA<br>R: AGGCCGCTAAATTACGTGCAGC | 95°C/5 min; (95°C/1 min, 59°C/1 min, 72°C/45 sec) X30; 72°C/5 min | 249 [3] | |
| tonB | F: TTGTGGTGCCTCTGCAATCGGT<br>R: TCGTGTACCCAAACGAGCAGGA | 95°C/5 min; (95°C/1 min, 55°C/1 min, 72°C/45 sec) X30; 72°C/5 min | 1279 [3] | |

Table 2: The biofilm-forming capacity of A. baumannii and percentages of their iron/siderophore mediate gene.

| Biofilm | Antibiotic resistance profiles (NO) (%) | Total |
|---------|----------------------------------------|-------|
| Strong  | IMI: 47.3 52.3 44.4 50 52.3 55.0 100 58.5 55.0 64.7 100 50 84.6 55.0 42.3 15.2 | 26.3 29.1 25.0 43.0 30.5 29.1 27.7 12.5 40.2 27.7 23.6 25.0 30.5 18.0 27.7 36.1 |
| Moderate| IMI: 47.3 52.3 47.6 38.8 64.5 45.4 47.6 45 55.1 45.0 35.2 33.3 45.5 18.1 45.0 50.0 31.9 | 52.2 54.4 47.6 45.4 64.5 45.4 47.6 45 55.1 45.0 35.2 33.3 45.5 18.1 45.0 50.0 31.9 |
| Weak   | IMI: 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 | 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 |
| Nonbiofilm| IMI: 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 | 19 21 18 31 21 20 9 29 20 17 18 22 13 20 26 72 |

Iron/siderophore mediate gene

| BasD | IMI: 47.3 52.3 48.0 30.2 32 41.0 42.8 45.0 100 31.0 45.0 52.9 81.1 | 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 |
| BauA | IMI: 57.8 53.3 61.1 35.4 50.0 52.3 55.0 100 37.9 55.0 64.7 61.1 | 11 11 11 11 11 11 11 11 11 11 11 11 9 11 9 11 |
| Pld   | IMI: 42.1 38.0 44.4 25.8 36.6 38.0 40.0 88.9 27.5 40.0 77.7 24.1 | 8 8 8 8 8 8 8 8 8 8 8 8 7 8 | 7 8 |
| paaE  | IMI: 36.8 52.3 61.1 35.4 40.0 52.3 55.0 100 31.0 55.0 64.7 61.1 | 9 11 11 11 11 11 11 11 11 11 11 11 9 11 11 11 |
| feoB  | IMI: 47.3 52.3 52.3 38.7 57.1 | 11 11 12 12 11 12 11 12 11 12 11 12 4 5 9 12 |
| entA  | IMI: 73.6 61.9 77.7 38.7 52.3 60.0 100 37.9 60.0 64.7 66.7 | 14 13 14 12 12 14 9 14 13 14 14 14 13 14 13 14 |
| hemO  | IMI: 89.4 71.4 100 54.8 77.2 | 17 15 17 17 16 14 19 17 17 17 17 17 17 12 14 11 17 |
| tonB  | IMI: 26.3 23.8 27.7 12.9 22.7 | 5 5 5 4 5 5 3 5 5 0 0 4 5 5 5 4 5 |

Clinical samples

| Wound | IMI: 9 11 11 11 11 11 11 11 11 11 11 11 11 11 11 11 11 | 19 20 10 9 2 11 9 7 16 10 10 9 13 7 9 13 23 |
| Urine  | IMI: 100 95.2 55.5 29.0 9.0 52.3 45.0 77.7 55.1 50.0 58.8 50.0 | 0 0 0 0 20 18 1 2 0 3 2 3 2 2 2 2 4 21 |
| Blood  | IMI: 11.1 64.5 81.8 4.7 10 | 0 0 0 5 1 2 9 2 7 7 3 6 7 4 9 7 18 |
| Body fluids | IMI: 27.7 3.2 9.0 42.8 45.0 28.5 | 0 0 0 1 1 1 0 0 0 3 1 1 1 0 0 0 2 10 |

IMI: imipenem; ERT: ertapenem; PIP: piperacillin; GEN: gentamycin; CIP: ciprofloxacin; AMK: amikacin; FOX: cefoxitin; AZT: aztreonam; OFL: ofloxacin; PIP/TAZ: piperacillin/tazobactam; TMP/SMX: trimethoprim/sulfamethoxazole; CEP: cefepime; MDR: multidrug-resistant; XDR: extensively drug-resistant; AmpC: ampicillins C; MBL: metallo-β-lactamase.
Figure 1: Evaluations of antibiotic resistance pattern and AmpC and MBL producing strains by disk diffusion method. (a) Resistant to cefoxitin (FOX) disk indicates AmpC producing strain. (b) The MBL producing isolates were considered positive when the zone diameter difference between imipenem + EDTA and imipenem discs was more extensive than 7 mm.

Figure 2: The amplification result of iron/siderophore mediate gene in A. baumannii. (a) basD gene with 868bp and bauA gene with 224bp; well 1: positive control; wells 2 to 7: positive strains with basD gene; well 11: positive control; wells 8 to 10: positive strains with bauA gene. (b) pld gene with 662bp and paaE gene with 593bp; well 1: positive control; well 3 to 10: positive strains with paaE; well 13: positive control; wells 7 to 12: positive strains with pld. (c) fecB gene with 636bp and hemO gene with 215bp; wells 1: positive control; wells 2 to 5: positive strains with hemO; wells 10: positive control; wells 6 to 9: positive strains with pld. (d) entA gene with 504bp; wells 1: positive control; wells 2 to 9: positive strains with tonB gene. (M) Ladder 100bp.
Tukey, IUS genes’ high frequency was significantly reported in AmpC/MBL producing isolates. However, biofilm formation is positively associated with antibiotic resistance, siderophore, and $\beta$-lactamase productions ($p < 0.05$). In terms of MDR and XDR strains, harboring IUS genes and biofilm formation was strongly associated with $\beta$-lactamase production. In contrast, AmpC/MBL $\beta$-lactamase production was positively associated with siderophore production ($p < 0.001$).

4. Discussions

Acinetobacter’s hospital infection has developed as a serious threat to the healthcare system due to the emergence of pan resistance from multiresistance. In recent years, A. baumannii has become a worldwide concern due to severe nosocomial infection.

In the current study, A. baumannii isolates obtained from several clinical samples were collected. Most samples were obtained from wound samples (31.9%) followed by urine (29.1%), blood (25.0%), and body fluids (13.8), which is concordance with study Lukovic et al. and Sheikh et al., where there are maximum isolates wound samples [18, 19].

The current study revealed the lowest frequency of resistance to aztreonam (12.5%), followed by TMP/SMX (23%) and PIP/TAZ (25%). This finding was also reported by Armin et al. and Kamali et al. [20, 21]. In line with Jasemi et al.’s [22] study, 27.7% and 36.1% of isolates were considered XDR and MDR, respectively. However, these findings were in contrast to the data reported from India, Sweden, and Italy [4, 23, 24]. Also, 27.7% and 36.1% of A. baumannii produced AmpC and MBL enzymes, respectively. A similar pattern of results was obtained in El-
Bakyet al. and Kaur and Singh. They showed that about 35% of A. baumannii isolates produced AmpC and MBL enzymes [8, 25]. However, these findings contrast with the previous studies in India and Pakistan [26, 27]. Also, by the time A. baumannii isolates acquire resistance to β-lactams and carbapenems, they often acquire resistance to several other antimicrobial agents as well [26].

According to the present study, the frequencies of basD, basA, pld, paaE, feoB, hemO, tonB, and entA genes in A. baumannii clinical isolates were 12.5%, 15.2%, 11.1%, 15.2%, 16.6%, 23.6%, 6.9%, and 19.4%, respectively. However, this frequency has not previously been described. The levels observed in this investigation are far below those observed by Abdi et al. [3]. They reported that frequencies of tonB, barA, feoB, entA, and hemO genes were 85%, 97%, 99%, 99%, and 95%, respectively. A study conducted by Liu et al. [2] also showed a high prevalence of iron/siderophore genes in A. baumannii isolates. Unfortunately, few studies have surveyed the abundance of iron/siderophore mediate genes.

A significant relationship was observed between antibiotic resistance and IUS genes’ distribution based on our study (p < 0.05). However, there are very few nationwide studies conducted on IUS genes associated with antibiotic resistance and biofilm formation in Iran. Furthermore, we found that AmpC/MBL producing among A. baumannii isolates is significantly correlated with increased biofilm formation (p < 0.001). A correlation between IUS and antibiotic resistance patterns was also reported by Zeighami et al. Moreover, these authors reported a correlation between IUS and upregulation of the β-lactamase gene encoding the AmpC and MBL enzymes. They suggested that IUS played an essential role in the prevalence of antibiotic resistance in clinical isolates of A. baumannii [28]. Moreover, following the present results, Kröger et al. and Mohajerji et al. have demonstrated that iron uptake through siderophores’ production is another factor of virulence in this organism. However, this finding broadly supports the work of other studies in this area linking iron/siderophore mediate with drug resistance [13, 28, 29].

A. baumannii isolates by several mechanisms of resistance can also differentially express various virulence factors. These factors include siderophore production, biofilm formation, and hemolysis on blood agar [13, 30]. However, the association between virulence and antimicrobial resistance seems to be a highly complex one that is still unclear. Notably, the effect of harboring β-lactamase on virulence in A. baumannii is not well defined.

Finally, an association between IUS and biofilm formation on the one hand and biofilm formation and antibiotic resistance on the other was determined. The relationship between β-lactamase production and biofilm formation has been previously reported among MDR A. baumannii isolates [12, 28, 29]. Moreover, the association between motility and siderophore production is not surprising since the former factor was associated with biofilm production. Simultaneously, the latter allows for iron acquisition that is crucial for biofilm formation [14]. These associations reveal a highly complex interplay between the different virulence determinants in A. baumannii, especially multifactorial. Therefore, considering this relationship, drug-resistant strains can be regarded as more prone to invasion. On the other hand, A. baumannii reduces its sensitivity to antibiotics by using iron/siderophore pathway regulatory systems.

In the present study, IUS pathways in antibiotic resistance in clinical isolates of A. baumannii were identified. We demonstrated that iron/siderophore uptake plays an essential role in the prevalence of AmpC/MBL producing strains among A. baumannii isolated from clinical specimens. Besides, the presence of feoB, entA, and hemO genes was associated with increased virulence and antibiotic resistance compared to pld, and tonB was positively associated with AmpC/MBLand siderophore production. Nevertheless, our results confirmed that IUS pathways significantly affect the forming extracellular layers in A. baumannii. Thus, suppression of feoB, entA, and hemO genes is one of the best ways to control virulent strains. It also inhibits the iron/siderophore uptake pathway increased biofilm-forming strains’ sensitivity to the β-lactams. Further, these associations could help better understand the complex interactions between antimicrobial resistance and virulence.

Data Availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Conflicts of Interest

The authors declare no competing interest.

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