

Distribution of the multidrug efflux pump genes *adeA*, *adeI*, *adeJ*, *adeY* and integrons in clinical isolates of *Acinetobacter baumannii* from Malaysian hospitals.

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Abstract

The objective of this study was to evaluate the incidence of multidrug resistance nodulation-cell division pump (RND pump) genes and integrons in multidrug-resistant *Acinetobacter baumannii* isolated from in-patients in Malaysian hospitals. A total of 43 clinical isolates of *A. baumannii* were examined. All samples were found to be 100% resistant to ampicillin (n=43), ceftriaxone (n=43) and cefuroxime (n=43), followed by augmentin (n=40) and chloramphenicol (n=40) (93%). Local isolates of *A. baumannii* were also found to develop resistance towards the new class of antibiotic-tigecycline (58.14%). The presence of *adeA*, *adeIJ*, *adeY* RND efflux system genes and integrase gene was assessed by polymerase chain reaction (PCR). *adeY* was not detected in any of the samples. However, majority of the isolates (62.7%, n=27) carried *adeA* and *adeIJ* genes, but not integrase gene. Interestingly, the isolates harboring these genes also showed the highest level of resistance toward all antibiotics. In addition, 13.9% of the isolates (n=6) did not carry any of the RND genes; while class 1 integrons were only detected in three isolates (7%, n=3), in which all harbored the same gene cassette *aacA4-catB8-aadA1*. This suggests the possibility of higher usage of RND efflux pump genes in antibiotic resistance mechanism for local *A. baumannii* compared to integron genes. Besides, the co-presence of *adeA* and *adeIJ* genes also reveals their possible relationship for antimicrobial resistance in *A. baumannii* isolates.

Keywords: *Acinetobacter baumannii*, RND efflux pump genes, Integrons, Multidrug resistance

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Introduction

Acinetobacter baumannii, a Gram-negative opportunistic pathogen, is the agent that frequently causes various types nosocomial infections, including urinary tract infections, meningitis and bacteremia in hospitalised patients [1-3]. It has been observed to be resistant to all existing classes of antibiotics [2], including glycolcyclines; a new class of modified tetracycline antibiotics [4].

Resistance-nodulation division (RND) family, a member of active efflux pump mechanism has been known to play an important role in multidrug resistance in Gram-negative bacteria. The efflux transporters are organized as multicomponent system where it is protected by an outer membrane and the efflux pump works in conjunction with a periplasmic fusion protein together with an outer membrane protein [5]. To date, five kinds of RND system has been described in *Acinetobacter* sp. [6-9]. Indeed, Ade

ABC efflux pump is well characterized. The increased expression of the *adeABC* genes has led to resistance to many antibiotics including aminoglycosides, beta-lactams, chloramphenicol, erythromycin, tetracyclines, and the dye ethidium bromide [6-10]. However, studies on the other two RND pump genes, *adeIJK* and *adeXYZ* are limited as compared to *adeABC*. Structural genes for *adeIJK* have been perceived to contribute to intrinsic low-level of resistance to several antibiotics in *A. baumannii* [11]. The role of *AdeXYZ* in antimicrobial resistance, on the other hand, has yet to be ascertained.

Integrons, first described by Stokes and Hall [12], have strong linkage with multidrug resistance in Gram-negative bacteria [13]. The association of the integron-driven capture system with mobile elements such as transposons and plasmids has facilitated the spread of antibiotic resistance genes between species significantly. To date, five distinct classes of integrons have been de-

scribed [14]. Class 1 integrons are commonly found in clinical isolates of Gram-negative bacilli, including *A. baumannii* [15-16]. In this study, we assessed the distribution of RND pump genes *adeA*, *adeI*, *adeJ* and *adeY*, and integrase genes in *A. baumannii* collected from hospitals around peninsular Malaysia, and analyzed their antibiotic susceptibility pattern. Such typing can be used to further characterize isolates and should prove to be very useful in epidemiological studies.

Materials and Methods

Bacteria strains and growth media

A total of 43 multidrug resistant (MDR) clinical *A. baumannii* strains were collected from various hospitals around Malaysia. These isolates were obtained from wound (n=10), pus (n=4), sputum (n=8), blood (n=1), swab (n=9) and others (n=11). The isolates were routinely grown at 37°C on Luria Bertani (LB) agar.

Antimicrobial susceptibility testing

Antimicrobial susceptibility was determined according to the Clinical and Laboratory Standards Institute (CLSI) using the Kirby-Bauer disk diffusion assay on Mueller Hinton agar (Oxoid, UK) in accordance to the described method in Cheong et al. [17]. The susceptibility profile was determined for 15 antibiotics: Amikacin (30 µg), Amoxicillin/Clavulanic acid (Augmentin, 3 µg), Ampicillin (10 µg), Ceftazidime (30 µg), Ceftriaxone (30 µg), Cefuroxime (30 µg), Chloramphenicol (30 µg), Ciprofloxacin (5 µg), Gentamicin (10 µg), Imipenem (10 µg), Meropenem (10 µg), Norfloxacin (10 µg), Tetracycline (30 µg), Tigecycline (15 µg), and Trimethoprim/Sulfamethoxazole (Co-trimoxazole, 25 µg). All antibiotic discs were purchased from Oxoid, United Kingdom.

Preparation of DNA template

Total DNA template was prepared using fast-boiling method. Overnight bacterial culture was centrifuged at 12,000 rpm for 5 min and pellet was re-suspended with 300 µl of sterile deionized water. The bacterial suspension was then boiled for 5 min and placed immediately on ice for another 2 min before centrifuging at 12,000 rpm for 2 min. The supernatant was aspirated and used for subsequent polymerase chain reaction.

Amplification of efflux pump genes

The presence of *adeA*, *adeI*, *adeJ* and *adeY* genes of RND efflux pump system was determined by PCR. The primers used and their references are listed in Table 1. PCR reaction was performed in a final volume of 25 µl containing PCR buffer, 2 mM of MgCl₂, 0.4 µM of each primers, 0.4 mM of dNTP mix, 1 U of *Taq* polymerase (Promega, USA) and 250 ng/µl of template DNA. PCR reaction was performed in thermal cycler (Biometra, Germany) with the parameter: initial denaturation 94°C for 5 min, 30 cycles of 94°C for 60 sec, 56°C for 60 sec and 72°C for 60 sec, and a final elongation at 72°C for 7 min. The pres-

ence and sizes of the amplicons were assessed by electrophoresis in 1.5% agarose gels stained with ethidium bromide.

Integrons, gene cassettes amplification and restriction fragment length polymorphism

Integrons were detected using degenerate primers, hep35 and hep36, which hybridize to the conserved region of integrase genes *intI1*, *intI2*, and *intI3*. Class of integrons was further determined by performing restriction fragment length polymorphism (RFLP) using *HinfI* and *RsaI* [18-19]. To detect for the inserted gene cassettes, 5'CS and 3'CS primers were used to amplify the variable regions of class 1 integrons [19-20]. The amplification protocol used was: denaturation step at 94°C for 5 min, 30 cycles of 94°C for 30 sec, 55°C for 30 sec and 72°C for 45 sec for integrase gene or 90 sec for amplification of gene cassette.

Results

Figure 1 shows the antimicrobial susceptibility profiles of the 43 clinical isolates of *A. baumannii*. As observed, all the isolates (100%) were resistant to ampicillin, ceftriaxone and cefuroxime. This was followed closely by augmentin and chloramphenicol (93%), tetracycline (79.1%), amikacin and ceftazidime (74.4%), and gentamicin and meropenem (72.1%). More than 58.1% of the isolates showed resistance to tigecycline.

In this study, four RND pump system genes (*adeA*, *adeI*, *adeJ*, *adeY*) and integrons were chosen as the target gene. The specificity of all primers used was confirmed with nucleotide sequencing of generated PCR product (data not shown). From the results obtained, the distribution of RND pump genes and integrons were classified into six groups (Table 2). All of the clinical isolates were screened negative for *adeY*, indicating its non-existence in *A. baumannii* samples. The PCR results also revealed that 29 samples (67.4%) carried *adeA*, *adeI* as well carried *adeJ* genes, indicating that AdeABC and AdeIJK could co-exist in *A. baumannii* [7, 21].

Out of these 29 samples, a total of 27 (93.1%) samples did not harbor any integrons (Group 1), but this group displayed high resistance towards the antibiotics tested especially ampicillin, ceftriaxone, cefuroxime, chloramphenicol. The isolates classified under Group 3 and 5, which carried only one type of RND efflux genes showed reduced resistance to most of the antibiotics as compared to isolates from Group 1. This proved that isolates with the co-existence of both RND efflux pumps were significantly more resistant to all of the tested antibiotics.

Group 4 consisted of the two samples, which were positive for all *adeA*, *adeI*, *adeJ* genes and integrons. As observed in Table 2, the presence of the genes in these samples significantly increased the resistance towards all antibiotics tested significantly. Both the isolates in this

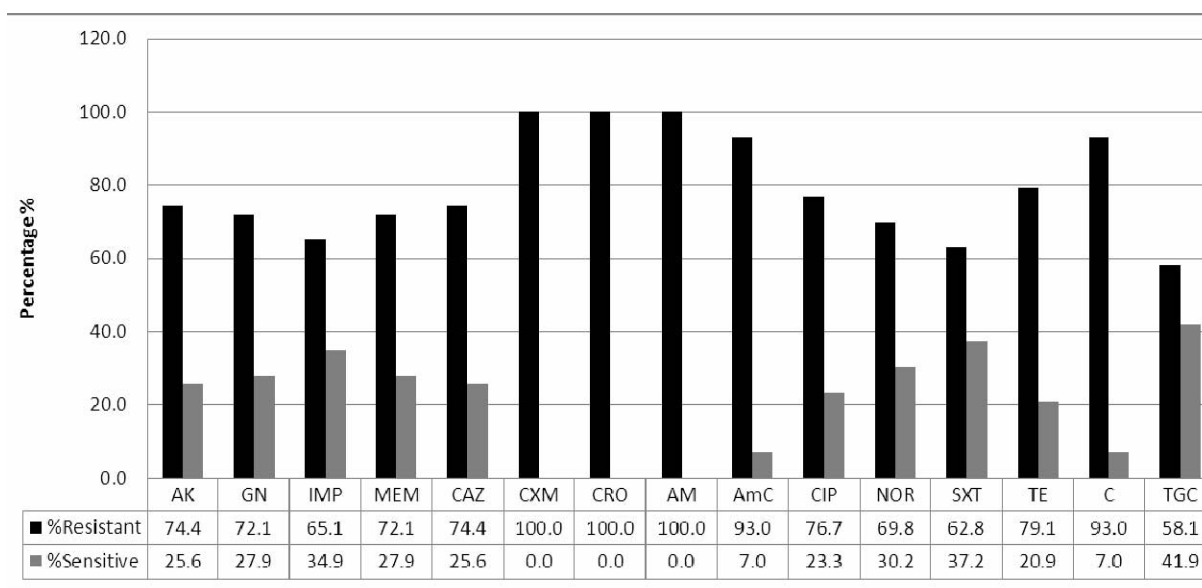


Figure 1. Analysis of antibiotics susceptibility profiles on 43 clinical isolates of *A. baumannii*. AK, amikacin; GN, gentamicin; IMP, imipenem; MEM, meropenem; CAZ, ceftazidime; CXM, cefuroxime; CRO, ceftriaxone; AM, ampicillin; AmC, amoxicillin/ clavulanic acid (Augmentin); CIP, ciprofloxacin; NOR, norfloxacin; SXT, trimethoprim/ Sulfamethoxazole; TE, tetracycline; C, chloramphenicol; and TGC, tigecycline

Table 1: Primers used in this study.

| Gene | Sequence | Expected Size (bp) | Sources |
|---------------|--|--------------------|---------|
| <i>adeA</i> | F: 5'- ATC TTC CTG CAC GTG TAC AT -3' R: 5'- GGC GTT CAT ACT CAC TAA CC -3' | 513 | [16] |
| <i>adeI</i> | F: 5'- ATC GCG CTT GTT GGT TGT AG -3' R: 5'- AAG CAC CAG CCG TTA CTG AA-3' | 541 | [16] |
| <i>adeJ</i> | F: 5'- ATT GCA CCA CCA ACC GTA AC -3' R: 5'- TAG CTG GAT CAA GCC AGA TA -3' | 453 | [16] |
| <i>adeY</i> | F: 5'- CAA TCT GCA ACT GCG CTT -3' R: 5'- TCA ACA GCT TCT GCG GTA -3' | 587 | [22] |
| integrase | F: 5'- TGC GGG TYA ARG ATB TKG ATT T -3' R: 5'- CAR CAC ATG CGT RTA RAT -3' | 491 | [18] |
| Gene cassette | 5'CS: 5'- GGC ATC CAA GCA GCA AG- 3' 3'CS: 5'- AAG CAG ACT TGA CCT GA- 3' | variable | [20] |
| 16S rRNA | F: 5'- GAC GTA CTC GCA GAA TAA GC -3' R: 5'- TTA GTC TTG CGA CCG TAC TC -3' | 426 | [16] |

Table 2. Antimicrobial susceptibility profiles in accordance to group distribution of *adeA*, *adeI*, *adeJ* and integron genes in clinical isolates of *A. baumannii*.

| Antimicrobial agents | Group 1 <i>adeA</i> ⁺ , <i>adeI</i> ⁺ , <i>int</i> ⁻ (n=27) | | Group 2 <i>adeA</i> ⁻ , <i>adeI</i> ⁻ , <i>int</i> ⁻ (n=5) | | Group 3 <i>adeA</i> ⁺ , <i>adeI</i> ⁻ , <i>int</i> ⁻ (n=6) | | Group 4 <i>adeA</i> ⁺ , <i>adeI</i> ⁺ , <i>int</i> ⁺ (n=2) | | Group 5 <i>adeA</i> ⁻ , <i>adeI</i> ⁺ , <i>int</i> ⁻ (n=2) | | Group 6 <i>adeA</i> ⁻ , <i>adeI</i> ⁻ , <i>int</i> ⁺ (n=1) | | |
|----------------------|--|--------------|---|-------------|---|--------------|---|------------|---|-------------|---|-------------|-----------|
| | R | S | R | S | R | S | R | S | R | S | R | S | |
| AK | 24 (88.9%) | 3 (11.1%) | 1 (20%) | 4 (80%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| AmC | 26 (96.3%) | 1 (3.7%) | 4 (80%) | 1 (20%) | 5 (83.3%) | 1 (16.7%) | 2 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 1 (100%) | 1 (100%) | 0 (0%) |
| AM | 27 (100%) | 0 (0%) | 5 (100%) | 0 (0%) | 6 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 1 (100%) | 1 (100%) | 0 (0%) |
| CAZ | 23 (85.2%) | 4 (14.8%) | 1 (20%) | 4 (80%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 1 (50%) | 1 (50%) | 1 (100%) | 1 (100%) | 0 (0%) |
| CRO | 27 (100%) | 0 (0%) | 5 (100%) | 0 (0%) | 6 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 1 (100%) | 1 (100%) | 0 (0%) |
| CXM | 27 (100%) | 0 (0%) | 5 (100%) | 0 (0%) | 6 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 1 (100%) | 1 (100%) | 0 (0%) |
| C | 27 (100%) | 0 (0%) | 2 (40%) | 3 (60%) | 6 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 2 (100%) | 0 (0%) | 1 (100%) | 1 (100%) | 0 (0%) |
| CIP | 25 (92.6%) | 2 (7.4%) | 2 (40%) | 3 (60%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 1 (50%) | 2 (100%) | 1 (100%) | 1 (100%) | 0 (0%) |
| GN | 23 (85.2%) | 4 (14.8%) | 1 (20%) | 4 (80%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| IMP | 21 (77.8%) | 6 (22.2%) | 1 (20%) | 4 (80%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| MEM | 24 (88.9%) | 3 (11.1%) | 0 (0%) | 5 (100%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| NOR | 23 (85.2%) | 4 (14.8%) | 0 (0%) | 5 (100%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| TE | 25 (92.6%) | 2 (7.4%) | 1 (20%) | 4 (80%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 1 (50%) | 1 (50%) | 1 (100%) | 1 (100%) | 0 (0%) |
| TGC | 18 (66.7%) | 9 (33.3%) | 2 (40%) | 3 (60%) | 2 (33.3%) | 4 (66.7%) | 1 (50%) | 1 (50%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |
| SXT | 21 (77.8%) | 6 (22.2%) | 0 (0%) | 5 (100%) | 4 (66.7%) | 2 (33.3%) | 2 (100%) | 0 (0%) | 0 (0%) | 2 (100%) | 2 (100%) | 1 (100%) | 0 (0%) |

group demonstrated resistance to all antibiotics, except for tigecycline. Our study revealed that *adeA* was the most prevalent RND pump, constituting to 81.4% (35/43) of the clinical isolates and this was followed closely by *adeIJ* which constituted 72.1% (31/43).

Only three out of 43 isolates (9.38%) was detected as class 1 integron positive: two isolates were placed in Group 4, while another was placed in Group 6. Interestingly, these samples exhibited much higher resistance to almost all of the antibiotics tested. PCR amplification of gene cassettes of class 1 integrons successfully generated amplicons with sizes of 2.3 kb. Nucleotide sequencing confirmed the presence of gene cassette *aacA4-catB8-aadA1* in all three integron-positive isolates of *A. baumannii*.

Group 2 comprises of 11.6% of the clinical isolates, which did not carry any of the *adeA*, *adeIJ* or integron genes. Other than the penicillins (i.e. augmentin and ampicillin) and the cephalosporins (i.e. ceftriaxone and cefuroxime), it was observed that more than 60% of the isolates in this group were sensitive to the antibiotics, which harbored at least one of the genes tested. The results here confirm the role played by RND genes and integrons in bacterial multidrug resistance.

Discussion

In this study, *adeA* gene is shown to be the most prevalent in our clinical isolates. *adeA* gene encodes for one of the proteins that make up the tripartite system of efflux pump AdeABC. The finding is consistent with several studies, which showed the high prevalence of AdeABC amongst *A. baumannii* strains [22-23]. Both *adeI* and *adeJ* genes also encode for component proteins of efflux pump AdeIJK, and they are co-transcribed simultaneously [11]. Here, *adeI* and *adeJ* were proven to be highly prevalent in *A. baumannii* isolated from Malaysian hospitals, and they were always detected together in the isolates. The organization of three adjacent *adeI*, *adeJ* and *adeK* genes suggests that AdeIJK formed an operon that is similar to AdeABC efflux system [7]. Therefore, the results conformed to previous studies [7, 11].

A high percentage of the isolates also harbored both *adeA* and *adeIJ* genes, which indicates the co-existence of AdeABC and AdeIJK efflux pumps system in *A. baumannii*. This is in line with other studies whereby majority of the isolates were positive for the presence of both groups of RND efflux pump and their roles in resistance development [16]. As expected, the co-existence of *adeA* and *adeIJ* genes greatly increased the resistance profile of the isolates, and thus, signifying their role in antibiotic resistance development [11]. This study also showed the increasing resistance toward tigecycline in this region, and thus conforming to recent studies [24-25].

Although not extensively distributed, the discovery of class 1 integrons in three of our clinical isolates is still an interesting find. The existence of the integron alone in the isolate (Group 6) rendered the isolate resistance to 14 out of 15 of the antibiotics tested. Similar to most studies,

only class 1 integrons were detected here. Class 1 integrons are by far the most common class of integrons in clinical isolates of Gram-negative bacteria [15, 19], including *A. baumannii* [26-29]. Gene cassette array showed the presence of a 2.3 kb cassette array, *aacA4-catB8-aadA1*, in all the integron-positive *A. baumannii* isolates. This array is particularly common in *A. baumannii* worldwide [29-31]. The high percentage of *aacA4-catB8-aadA1* existence in class 1 integrons implicates that this cassette is the prevalent one among the *A. baumannii* isolates and the epidemiological relationship to other *A. baumannii* strains worldwide. Further studies are required to determine the clonal relationship with the *A. baumannii* strains reported in other countries, and also to determine if there are mutations occurring inside the gene cassettes that affect cassette gene expression.

In conclusion, the present work gives a brief description on the distribution of RND efflux pumps, integrons and their associated cassette arrays in *A. baumannii* isolates in Malaysia. *A. baumannii* isolates in this study demonstrate the frequent use of RND efflux pump genes rather than the acquisition of mobile gene cassette using site-specific recombination integrons.

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Abbreviation: MDR, Multidrug resistant;
 RND pump, Resistance nodulation-cell division pump