RND efflux pump systems in *Acinetobacter*, with special emphasis on their role in quorum sensing

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*Acinetobacter* is an important opportunistic, multidrug resistant pathogen causing majority of nosocomial infections worldwide. The multidrug resistance is attributed by a plethora of efflux pumps and the overexpression of the same mediates export of antimicrobial agents. Quorum sensing (QS) is the cell-to-cell communication system in which bacteria produces specific signaling molecules which are transported out to the surrounding environment to communicate with other bacterial cells. It has been noticed that multidrug efflux pumps like resistance-nodulation-cell division (RND) efflux pumps play an important role in QS by exporting these signaling molecules. This review discusses various RND efflux pumps and the current understanding of the interrelationship of RND efflux pumps and QS in *Acinetobacter* spp. Studies demonstrate that RND efflux pumps could be considered as potential targets to block QS thereby reducing pathogenesis and antibiotic resistance. The known RND efflux pump-mediated quorum quenching strategies for *Acinetobacter* and other bacterial strains are discussed in detail. Finally, the prospective quorum quenching strategies targeting the transcriptional regulators of RND efflux pumps to inhibit multidrug efflux pumps are addressed.

Key Words: *Acinetobacter*, RND efflux pumps, quorum sensing, quorum quenching

INTRODUCTION

Nowadays, the applicability of antibiotics is undermined by the evolution of multidrug resistant bacteria. Bacteria has developed various resistance strategies to fight against antibiotic stress and the most known ones are modification of the antibiotic by hydrolysis, acetylation or adenylation of the specific drugs, phosphorylation, modification of the antibiotic target by mutation or methylation, isolating the toxic compound by non-essential proteins in the cell or altering the membrane permeability of drugs (1). The pathogenic bacteria can transfer the resistant determinants to other organisms by horizontal gene transfer as many of these resistant gene elements are located on plasmids, transposons and integrons (2, 3). The first line of defense for many bacteria is to prevent the entry of toxic compounds via the cell membrane. Though the cell membrane of bacteria acts as an effective barrier to prevent the entry of many toxic compounds because of its amphipathic nature, the compounds may find their way into cells through pores and porins in the outer membrane (4). Thus, the downregulation of the expression of porins is an effective way to block the entry of toxic compounds into the cells (4).
In addition, bacteria express a plethora of multidrug efflux pumps and the overexpression of these pumps alone is enough to cause multidrug resistance (MDR), without additional resistance factors (5).

Multidrug efflux pumps are classified in six families: major facilitator superfamily (MFS), small multidrug resistance family (SMR), ATP-binding cassette superfamily (ABC), multidrug and toxic compound extrusion family (MATE), proteobacterial antimicrobial compound extrusion family (PACE) and resistance-nodulation-division superfamily (RND) (5). The SMR, MATE and MFS families form the main efflux pump systems in Gram-positive bacteria, while RND efflux pumps are widely distributed in Gram-negative bacteria. Depending on the specific family the efflux pumps belong to, they can be either single component transporters or multicomponent systems with an inner membrane transporter, outer membrane channel and a periplasmic protein, such as RND efflux pumps (6). The RND pumps are majorly associated with clinically relevant antibiotic resistance, such as AcrAB-TolC efflux pump of Escherichia coli (E. coli) and Salmonella enterica serovar Typhimurium (S. typhimurium) and MexAB-OprM of Pseudomonas aeruginosa (P. aeruginosa) (7). Just like other RND efflux pumps, AcrAB-ToIC efflux pumps are tripartite efflux pumps having three components, a transporter (efflux) protein, AcrB in the inner membrane, an outer membrane protein channel, ToIC and a periplasmic accessory protein, AcrA (8). The RND efflux pumps are usually expressed at a basal expression level in bacteria which helps them to survive in the presence of toxic compounds (9). The AcrAB efflux pump is reported to recognize and transport a wide range of structurally unrelated compounds including antibiotics, bile salts, dyes and detergents (10). The substrate profile of AcrAB-ToIC efflux pump of E. coli includes chloramphenicol, lipophilic β-lactams, fluoroquinolones, tetracycline, rifampin, novobiocin, fusidic acid, nalidixic acid, ethidium bromide, acriflavine, bile salts, short-chain fatty acids, SDS, Triton X-100, and triclosan (11-15).

Apart from their involvement in antibiotic resistance, RND efflux pumps are reported to play a role in bacterial pathogenicity by contributing to colonization and persistence of bacteria in their ecological niche (9). These efflux pumps extrude various host-derived antimicrobial compounds such as bile salts, fatty acids and detergents promoting the adaptation and survival of the bacterium in their ecological and physiological niches (16). The efflux of bile salts via AcrAB efflux pumps or its homologs has been reported in E. coli, P. aeruginosa, Neisseria gonorrhoeae (N. gonorrhoeae) and S. typhimurium (16-20). The defective mutations in these efflux pumps caused reduced virulence in several pathogens. In S. typhimurium, the inactivation of acrAB impaired intestinal colonization in murine model, indicating that the AcrAB efflux pump is required for full virulence (21). In addition, Buckley et al. reported that acrB and tolC gene mutants colonize poorly in the avian gut, pointing that AcrAB-ToIC system is essential for colonization of S. typhimurium in chickens (22). Similarly, In N. gonorrhoeae, a bacterial pathogen of the human genital mucosa, the deletion of mtrD or mtrE gene, the product of which constitutes the MtrCDE efflux system lead to poor bacterial colonization in genito-urinary tract of female mice (23). The CmeABC efflux pump of Campylobacter jejuni confers resistance to a wide range of antimicrobials and the functional inhibition of this efflux pump could prevent bacterial host colonization (24). Recently, the role of AcrAB-ToIC in virulence has also been reported in Klebsiella pneumoniae and Enterobacter cloacae in which the efflux pump defective mutants displayed reduced capability to infect mouse model (25, 26). In P. aeruginosa, the mutant lacking the MexAB-OprM, a homolog of AcrAB-ToIC couldn’t invade Madin–Darby canine kidney (MDCK) cells and it was suggested that the MexAB-OprM efflux system could export virulent determinants important for bacterial pathogenesis (27). The tolC mutant of S. typhimurium poorly adhered to both human embryonic intestinal cells and mouse monocyte macrophages, showing that the efflux pump system has a role in mediating bacterial adherence (22).

In addition to their direct role in bacterial pathogenesis, the efflux pumps affect bacterial virulence in a more indirect manner by altering the cell to cell communication (quorum sensing, QS) response in bacteria. QS is mediated by the release of chemical signaling molecules called autoinducers which are synthesized in vivo and needs to be transported across the cell membrane. The first report of autoinducers as substrate of RND family efflux pumps was made in studies of P. aeruginosa, in which the QS signals, acyl homoserine lactones (AHLs) are exported out by MexAB-OprM system (28, 29). P. aeruginosa overexpressing Mex pumps displayed reduced virulence due to the increased efflux of AHLs and the inability of the cells to accumulate QS signals. In addition, the overexpression of MexCD-OprD and MexEF-OprN are associated with the reduced expression of genes encoding type III secretion in P. aeruginosa (30). In E. coli, the overexpression of the QS regulator SdiA
lead to the overexpression of AcrAB efflux pump, suggesting a potential role of this pump in QS (31). In this review, we focus on the RND efflux pump systems in the nosocomial pathogen, *Acinetobacter* spp. with special emphasis on their role in QS. In addition, we discuss the current state of knowledge on quorum quenching strategies by inhibiting the RND efflux pumps and future perspectives to effectively tackle this nosocomial pathogen.

**RND efflux pump systems in *Acinetobacter* spp**

In *Acinetobacter baumannii* (*A. baumannii*), three RND efflux systems, AdeABC (32), AdeFGH (33), and AdeIJK (34) are reported to be primarily associated with MDR. AdeABC is the first characterized RND efflux pump in *A. baumannii* and is comprised of the major fusion protein AdeA, a multidrug transporter AdeB, and the outer membrane factor OMF (32). The AdeA and AdeB shares similarity to AcrA (55%) and MexA (58%) and to AcrB (68%) and MexB (67%), respectively, of *E. coli* and *P. aeruginosa*. The expression of AdeABC is under the tight control of the two-component regulatory system AdeR-AdeS, encoded by adeRS operon located upstream of adeABC operon (35). AdeS is a sensor kinase which monitors the environmental conditions and activates or inactivates the response regulator, AdeR which regulates the expression of the efflux pump (35). In addition, the AdeABC and AdeIJK RND pumps are indirectly regulated by the two-component regulatory system BaeSR in *A. baumannii* (36, 37). The AdeABC efflux system is not expressed in natural isolates of *A. baumannii* and the overexpression of the pump confers MDR by extruding aminoglycosides, β-lactams, fluoroquinolones, tetracyclines, tigecycline, macrolides, chloramphenicol, and trimethoprim (32). The AdeFGH efflux pump, encoded by the adeFGH operon provides high-level resistance to fluoroquinolones, chloramphenicol, trimethoprim, and clindamycin and decreased susceptibility to tetracyclines, tigecycline, and sulfamethoxazole without affecting β-lactams and aminoglycosides (33). As in the case AdeABC efflux pump, AdeFGH pump is also not constitutively expressed in wild type strains. A putative LysR-type transcriptional regulator, named AdeL, encoded by adeL located upstream of the adeFGH operon controls the expression of the AdeFGH efflux pump in *A. baumannii* (33). The AdeIJK efflux pump, encoded by the adeIJK operon is specific for the species (38, 39) where it confers intrinsic resistance to β-lactams, such as ticarcillin, cephalexin, and aztreonam, fluoroquinolones, tetracyclines, tigecycline, lincosamides, rifampin, chloramphenicol, cotrimoxazole, novobiocin, and fusidic acid (34). It has been noticed that AdeIJK acts in a synergistic fashion with AdeABC to export compounds such as tigecycline (34). The AdeXYZ efflux pump, encoded by the adeXYZ has 97% identity with AdeIJK (34) and is found in *Acinetobacter* GDG3, *Acinetobacter* GDG13TU and *Acinetobacter* GDG 17 (40). However, the functional aspect of this efflux pump with respect to antimicrobial resistance has not well understood. Another efflux pump system, AdeDE confers resistance to aminoglycosides, fluoroquinolones, erythromycin, tetracycline and chloramphenicol in *Acinetobacter* spp. belonging to *Acinetobacter* genomic DNA group 3 (41). AdeDE is encoded by the membrane fusion protein gene, adeD and the RND transporter gene, adeE, and the outer membrane protein of AdeDE has not been identified.

**Quorum sensing in *Acinetobacter* spp**

Quorum sensing is the regulatory mechanism by which bacterial cells communicate each other producing signaling molecules called autoinducers. At a specific density in the environment, the autoinducers specifically bind to transcriptional regulators thereby altering the expression of various genes in a population (42). It has been known that QS plays a major role in the production of virulence factors, motility, nodulation, plasmid transfer, antibiotic production, bioemulsion production, bioluminescence and biofilm formation (43-45).

*A. baumannii* has one QS system which is mediated by the two-component system, Abal/AbaR which is homologous to the typical LuxI/LuxR system in *E. coli*. Abal is the autoinducer synthase that synthesizes AHLs, the signaling molecules which interact directly with the AbaR and this complex binds to specific promoter sequences (*lux-box*) regulating the expression of QS target genes (46). It has been reported that Abal is responsible for the production of N-(3-hydroxydodecanoyl)-L-HSL (3-hydroxy-C12-HSL) in *A. baumannii* strain M2 (46). The complete genome analysis of *A. baumannii* ATCC 17978
suggested that the autoinducer synthase AbaI and acyltransferase are the sole enzymes responsible for the synthesis of AHLs of varying chain lengths by this organism (46). The comparative analysis of the autoinducer synthase from Gram negative bacteria revealed that AbaI is 45% identical to the autoinducer synthases from environmental isolates such as Halothiobacillus neapolitanus, Acidithiobacillus ferrooxidans ATCC 23270, Thiobacillus ferrooxidans, Pseudomonas spp. (pmt), Rašlonia solanaearum and Burkholderia ambifaria MC40-6 and 47% identical to the one from Pseudomonas spp. RW105 (47). The communication among bacterial cells depending on the cell density plays a crucial role in the maturation of biofilms (46, 48). In A. baumannii, mutation in abai which is responsible for the production of AHLs lead to 30-40% reduction in biofilm formation compared to the parental strain (49). In addition, Niu et al. revealed that the exogenous addition of AHLs restored the biofilm maturation in abai mutant of A. baumannii M2 (46). In a different study, the supplementation of AHL exogenously produced biofilm in a biofilm-negative clinical isolate of Acinetobacter (50). Recently, a homologue of the Abal/Abar system, referred as AnoI/AnoR was characterized in A. nosocomialis (51). The AnoI/AnoR system shares 94% identity with the Abal/Abar of A. baumannii. The A. nosocomialis strain produces N-(3-hydroxy-dodecanoyl)-L-homoserine lactone (OH-dDHL) as the signaling molecule and the anoI mutant was not able to synthesize OH-dDHL pointing that AnoI is important for the production of AHLs (51). In addition, the expression of anoI was derepressed in the anoR mutant, suggesting the role of AnoR as activator of anoI in A. nosocomialis. The deletion of anoR contributed to impaired biofilm formation and surface motility and the complementation of anoR in the anoR deletion mutant restored both the characteristics to that of wild type, indicating that AnoR is important for biofilm formation and motility (51).

It has been known that 63% of Acinetobacter strains identified so far produce more than one AHL (52). Though majority of the A. baumannii clinical isolates produce more than six AHLs, only one type is detected abundantly among them (50). Also, the AHL production is dependent on the culture conditions and four AHLs were identified when grown on minimal media and three when grown on minimal media with 0.1% tryptone in A. calcoaceticus BD413 (53). It is interesting to see that although multiple AHLs have been identified in Acinetobacter, only one autoinducer synthase is identified so far (46). Thus, it can be assumed that the AHL synthase has low specificity and it might be producing other AHLs as well. The quorum sensing signals are not homogenously distributed in Acinetobacter strains and thus it is difficult to distinguish the virulent and non-virulent strains based on the type of AHLs (52). However, one particular sensor, the Rf1-type sensor is widely distributed in strains belonging to A. calcoaceticus - A. baumannii complex (52).

Interrelationship of RND efflux pumps and quorum sensing in Acinetobacter spp

The QS system is mediated by the synthesis of signaling molecules and these molecules need to be exported into the surrounding media either by diffusion or active efflux (48). The AHLs vary in their carbon chain length and the short chain AHLs (4-8 carbon atoms) can easily diffuse through the cell membrane while the long chain AHLs (10-12 carbon atoms) need active transport across the membrane (48). The secretion of quorum sensing signals has been associated with multidrug efflux pumps (54, 55). In A. baumannii, the AHLs are exported into the extracellular environment through the AdeFGH efflux pump and the increase in concentration of AHLs in the extracellular environment accelerates the entry of AHLs into the intracellular environment to form Abar-AHL complexes (54). The AHLs exported out are sensed by nearby A. baumannii cells and the increased interaction between cells through AHLs results in the acceleration of biofilm formation. In another study, the overexpression of AdeABC efflux pump in A. baumannii displayed increased biofilm formation and virulence phenotype though very little is known on various genes associated with mechanisms related to QS in this nosocomial pathogen (56). In a clinical isolate of A. baumannii strain S, the expression of multidrug efflux pump genes adeA and adeB were induced by the production of AHLs (57). The loss of AHL production in the mutant strain lead to the decreased mRNA expression level of these efflux pump genes while the addition of AHLs restored their expression. Thus, it can be hypothesized that the AHL-mediated induction of AdeA and AdeB expression might be contributing to multidrug resistance in A. baumannii. Similarly, in a recent study, it was reported that the expression of acrA and acrB efflux pump genes encoding the AcrAB multidrug efflux system in A. nosocomialis is downregulated in the absence of the quorum
sensing regulator, AnoR, suggesting that the synthesis of AHLs is important for the proper functioning of the AcrAB efflux pump (58).

RND efflux pump-mediated quorum quenching strategies

Since QS allows bacteria to interact with the external environment and to adapt with the environmental changes thereby boosting its survival, inhibiting the QS system (quorum quenching) is a promising strategy to control infections (59-61). Multidrug efflux pumps have been reported to play a central role in the secretion of QS signals in *Acinetobacter* strains apart from drug resistance and pathogenicity. It has been known that the AdeABC, AdeFGHI and AcrAB efflux pumps are correlated to the QS system (54, 57, 58) in *Acinetobacter*. In *A. baumannii* S strain, the AHLs promoted the expression of AdeA and AdeB genes in the presence of the antibiotic meropenem, suggesting that the strain produces AHLs to increase antibiotic resistance (57). In addition, the AdeFGH efflux pump of *A. baumannii* plays a crucial role in biofilm formation during infection by secreting AHLs (54). In *A. nosocomialis*, the deletion mutant of the transcriptional regulator, AcrR which negatively controls the expression of AcrAB efflux pump displayed increased biofilm formation, motility and virulence (58). It was suggested that the enhanced expression of the AcrAB efflux pump in the *acrR* deletion mutant might be attributing well to the AHLs-mediated cell-to-cell signaling, contributing to increased virulence phenotypes. Thus, the development of inhibitors which target these multidrug efflux pumps would be a potential strategy to disrupt QS thereby controlling infections (Fig. 1). The already reported multidrug efflux pump inhibitors in *A. baumannii* include PAβN (phenylalanine-arginine β-naphthylamide, also called MC-207 110), CCCP (carbonyl cyanide-m-chlorophenylhydrazone), omeprazole, reserpine, verapamil, NMP(1-(1-naphthylmethyl)-piperazine), N-tert-butyl-2-(1-tert-butyltetrazol-5-yl) sulfanyl acetamide, (E)-4-((4-chlorobenzylidene)amino) benzene sulfonamide, etc. (62-66).

![RND efflux pump](image1.png)

**Figure 1.** Schematic representation of quorum quenching strategies targeting multidrug efflux pumps, considering *A. nosocomialis* as a representative strain. *anoR* and *anoR* genes encode AHL signal synthase and AHL receptor/activator protein respectively. The efflux pump inhibitors binding to the efflux pumps inhibit the export of AHLs to the external environment blocking the quorum sensing system. The nearby bacterial cells are deprived of AHLs to be transported into the cells disrupting the formation of AnoR-AHL complex, thus preventing the binding of AnoR-AHL to the promoter regions of target genes including those for biofilm formation and virulence. The *acrA* and *acrB* efflux pump genes are shown as examples of multidrug efflux pump genes in *A. nosocomialis*. The biomolecules which target the transcriptional regulators of efflux pump genes inhibit the expression of these genes leading to the disruption of the quorum sensing system.
It has been reported that metallic nanoparticles represent a potential candidate to block multidrug efflux pumps in bacteria (67). In *Staphylococcus aureus* (*S. aureus*), zinc oxide nanoparticles displayed inhibitory role on efflux pumps (67). Similarly, the synergistic use of polyacrylic acid-coated iron oxide (magnetite) nanoparticles (PAA-MNP) with rifampicin against *Mycobacterium smegmatis* resulted in four-fold higher growth inhibition than that of rifampicin alone (68). Christena et al. reported the applicability of copper nanoparticles as efflux pump inhibitors in *S. aureus* and *P. aeruginosa* (69). The above studies prove the potential of nanoparticles as multidrug efflux pump inhibitors and the utility of these nanoparticles as efflux pump inhibitors could be investigated in *Acinetobacter* spp. as well. In addition, trifluoromethyl ketones (TFs) are reported to inhibit the QS response through their inhibition of efflux pumps in *Chromobacterium violaceum* 026 and *E. coli* (70). Since TFs inhibit efflux systems and QS, and also have significant antibacterial property, they could be exploited to treat infections that rely on QS and efflux-pump mediated multidrug-resistant phenotypes like *Acinetobacter* infections.

Apart from targeting the multidrug efflux pumps directly as a quorum quenching strategy, the efflux pump-mediated QS could also be inhibited indirectly by targeting the transcriptional regulators which controls the efflux pump expression (Fig. 1). In *A. baumannii*, the overexpression of the AdeABC efflux pump is mediated by the regulator AdeR (71). Also, the AdeABC and AdelIk pumps are positively regulated by the BaeSR regulon in *A. baumannii* (36). Thus, the modulators which target AdeR or BaeR would be a promising tool to regulate the expression of the corresponding multidrug efflux pumps in *A. nosocomialis*. In *Yersinia enterocolitica*, the AcrAB-TolC efflux pump is positively regulated by the regulator protein, OmpR (72). In our lab, *in silico* analysis and the electrophoretic mobility shift assay revealed that OmpR can bind to the *acrAB* promoter region in *A. nosocomialis*, suggesting that OmpR might be controlling the expression of AcrAB efflux pump (data not published). However, further studies are of importance to elucidate the regulatory mechanism of OmpR in controlling the expression of AcrAB efflux pump in *A. nosocomialis*. In addition, screening for potential inhibitors of OmpR would be a desirable strategy to regulate the expression of OmpR thereby controlling the AcrAB efflux pump-mediated QS in *A. nosocomialis*.

**CONCLUSION**

Significant advances have been made in elucidating the functional aspects of various RND multidrug efflux pumps in *Acinetobacter* spp. RND efflux pumps are associated with MDR in *Acinetobacter* strains by extruding out antibiotics. Apart from this, these efflux pumps play an important role in the efflux of QS biomolecules thereby increasing bacterial virulence and biofilm formation. Thus, the inhibition of the RND efflux pumps would be a potential strategy to disrupt QS and to reduce the severity of infections. It is worth noticing that there are some promising RND efflux pump inhibitors like nanoparticles reported in other bacterial strains and the efficacy of the same could be tested in *Acinetobacter* as well. In addition, targeting the regulators of RND efflux pumps to inhibit multidrug efflux pumps would serve as a promising prospective quorum quenching strategy. Further studies are desired in this direction to completely understand the mechanism of transcriptional regulation of various efflux pumps and to screen the modulators of these regulators. The inhibition of RND efflux pumps not only increases the sensitivity of bacterial cells to antimicrobial agents but also blocks the QS system thereby reducing pathogenesis. Thus, the studies which focus on screening new biomolecules targeting transcriptional regulators of multidrug efflux pumps would provide new therapeutic options to effectively control *Acinetobacter* infections.

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