Contribtion of the Ade Resistance-Nodulation-Cell Division-Type Efflux Pumps to Fitness and Pathogenesis of Acinetobacter baumannii

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ABSTRACT Overexpression of chromosomal resistance-nodulation-cell division (RND)-type efflux systems with broad substrate specificity contributes to multidrug resistance (MDR) in Acinetobacter baumannii. We have shown that modulation of expression of the structural genes for the efflux systems AdeABC and AdeIJK confers MDR and results in numerous alterations of membrane-associated cellular functions, in particular biofilm formation. However, the contribution of these RND pumps to cell fitness and virulence has not yet been studied. The biological cost of an antibiotic resistance mechanism is a key parameter in determining its stability and dissemination. From an entirely sequenced susceptible clinical isolate, we have generated a set of isogenic derivatives having single point mutations resulting in overexpression of each efflux system or with every pump deleted by allelic replacement. We found that overproduction of the pumps results in a significant decrease in fitness of the bacterial host when measured by competition experiments in vitro. Fitness and virulence were also evaluated in vivo both in systemic and pulmonary infection models in immunocompetent mice. A diminished competitiveness of the AdeABC-overexpressing mutant was observed only after intraperitoneal inoculation, but not after intranasal inoculation, the latter mimicking the most frequent type of human A. baumannii infection. However, in mice infected intranasally, this mutant was more virulent and stimulated an enhanced neutrophil activation in the lungs. Altogether, these data account for the observation that adeABC overexpression is common in MDR A. baumannii frequently found in ventilator-associated pneumonia.

IMPORTANCE Overproduction of the RND AdeABC efflux system is observed with a high incidence in multidrug-resistant Acinetobacter baumannii and results in increased resistance to several antibiotics of choice for the treatment of infections caused by this nosocomial pathogen. It was therefore important to study the biological cost of the overexpression of the adeABC structural operon which is normally tightly regulated. Fitness diminution of an overexpressing mutant detected in vitro in a model that mimics sepsis was not observed in a pulmonary infection model in which the mutant was more virulent. This points out that increased virulence can occur independently from prolonged persistence in the infected organ and can account for the elevated incidence of this resistance mechanism in clinical isolates. The study also indicates that transposon libraries will identify only virulence genes that are expressed under physiological conditions but not those that are tightly regulated.

Acinetobacter baumannii is a major nosocomial pathogen that is clinically and epidemiologically successful at least in part due to its ability to persist in the hospital environment (1). Multidrug-resistant (MDR) strains have recently emerged resulting from the high capacity of A. baumannii to acquire genetic determinants and to the overproduction of resistance-nodulation-cell division (RND) efflux systems with broad substrate specificities (2). Three Acinetobacter drug efflux (Ade) RND systems, AdeABC (3), AdeFGH (4), and AdeIJK (5), have been reported to contribute to MDR. Expression of each pump is tightly regulated, AdeABC by the two-component regulatory system AdeRS (6), AdeFGH by the LysR-type transcriptional regulator AdeL (4), and AdeIJK by the TetR transcriptional regulator AdeN (7). Overexpression of the adeABC operon secondary to mutations in AdeRS plays a major role in MDR of clinical isolates (6, 8–10), whereas overexpression of adeFGH is rarely observed (8, 10); the constitutive expression of adeIJK is associated with intrinsic resistance (5, 9).

The fitness cost of antibiotic resistance mechanisms is a key parameter in determining the success of resistant bacteria (11). It is well established that efflux systems extrude not only antibiotics but also intracellular metabolites and that they play a role in the pathogenicity of Gram-negative bacteria (12). Inactivation of genes for efflux pumps impairs colonization or virulence as reported for acrAB in Salmonella enterica serovar Typhimurium (13), mexAB-oprM in Pseudomonas aeruginosa (14), acrAB in Klebsiella pneumoniae (15), and acrAB-toIC disruption in Enterobacter cloacae reduces both bacterial fitness and host virulence (16). Overexpression of efflux pumps has various and contrasting

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consequences on cell fitness and virulence: MtrCDE overproduction in Neisseria gonorrhoeae leads to increased fitness in vitro and higher virulence in the mouse vaginal infection model (17), whereas in Stenotrophomonas maltophilia, SmeDEF overproduction impairs fitness and decrease virulence (18); MexEF-OprN overproduction does not alter the fitness of Pseudomonas aerugi-

 TABLE 1 Relative growth rates of A. baumannii BM4587 and derivatives overexpressing each pump or with each pump deleted

| A. baumannii strain | Mutationa | Pump overexpression rate (fold)b,c | Relative growth rate (min−1)d | Reference |
|-------------------|-----------|-----------------------------------|-----------------------------|-----------|
| BM4587            | Wild-type | 1                                 | 1.00                        | 33        |
| BM4688            | AdeS9352K | 38 for adeB                        | 0.96 ± 0.05                 | 9         |
| BM4689            | AdeR751V  | 222 for adeB                       | 0.91 ± 0.04                 | 9         |
| BM4717            | ΔadeB    | NA                                | 1.00 ± 0.03                 | 9         |
| BM4690            | ΔadeI2N3541 | 402 for adeG                  | 0.82 ± 0.02                 | 9         |
| BM4691            | ΔadeI2Q3240p | 636 for adeG                | 0.81 ± 0.03                 | 9         |
| BM4718            | ΔadeG    | NA                                | 1.00 ± 0.03                 | 9         |
| BM4666            | adeNAC584 | 6 for adeJ                        | 0.94 ± 0.03                 | 33        |
| BM4719            | ΔadeJ    | NA                                | 0.98 ± 0.01                 | 9         |

a Data taken from reference 9.

b The pump overexpression rate values are shown as the fold or number of times the strain overexpressed the indicated gene. For instance, strain BM4688 overexpressed adeB 38-fold compared to the wild-type or parental BM4587 strain. NA, not applicable.

c The growth rates of derivatives overexpressing each RND efflux system or with each efflux system deleted were determined at the beginning of the exponential phase. The relative growth rate represents the ratio of growth of an isogenic derivative to that of the parental (wild-type) strain taken as 1.00. Every isolate was tested in duplicate in four to six independent experiments, and the results are presented as means ± SD. The mean values that are significantly different (P < 0.05 by the Student t test) from the value for the parental strain are indicated by an asterisk.

d The growth rate (per generation, respectively) associated with very high levels (402- and 636-fold) of adeG expression. Deletion of the adeG gene did not affect the host fitness.

The two AdeABC-overproducing isogenic mutants, strains BM4688 and BM4689, presented a moderately reduced relative growth rate which paralleled the level of overproduction of the pump (0.96 for 38-fold and 0.91 for 222-fold increased adeB expression, respectively). No relative growth rate reduction was observed for deletion mutant derivative BM4717.

Two AdeFGH-overproducing mutants, BM4690 and BM4691, had the largest fitness disadvantage (0.82 ± 0.02 and 0.81 ± 0.03, respectively) associated with very high levels (402- and 636-fold) of adeG expression. Deletion of the adeG gene did not affect the host fitness.

In this study, the contribution of the three Ade RND systems on the fitness and virulence of A. baumannii was determined using an entirely sequenced susceptible clinical isolate and a set of isogenic derivatives having single point mutations resulting in overexpression of each of the efflux systems or with deletion of every pump by allelic replacement (9). We found that overexpression of the pumps results in a significant decrease in fitness of the bacterial host when measured by in vitro competition experiments. However, in vivo in systemic and pulmonary infection models in immunocompetent mice, there were different patterns of fitness and virulence. Altogether, these data account for the observation that MDR A. baumannii with adeABC overexpression are frequent in clinical settings.

RESULTS

Impact of overexpression or deletion of RND efflux systems on in vitro fitness of A. baumannii. Growth rates were determined in monolayers at the beginning of the exponential phase in the absence of antibiotics, and the relative growth rates of the pump mutants are represented relative to that of the A. baumannii BM4587 parental strain (Table 1).

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For a more sensitive fitness evaluation, in vitro competition experiments were performed by mixing each derivative overexpressing a pump with the parent at an initial ratio of 1:1 for ca. 150 generations (Fig. 1). Strains BM4689 [adeABC] (BM4689 overexpressing adeABC) and BM4688 [adeABC] had competitive disadvantages of 3.1% and 2.2% per generation, respectively. Strains BM4691 [adeFGH], BM4690 [adeFGH], and BM4666 [adeIJK] had competitive disadvantages of 4.3%, 2.7%, and 2.2% per generation, respectively.

Comparison of A. baumannii virulence in mice after i.p. or i.n. inoculation. The i.p. 50% lethal dose (LD50) of A. baumannii parental strain BM4587 was 4.9 × 10^7 CFU/mouse, which indicates moderate virulence compared to other strains of A. baumannii with i.p. LD50 values in immunocompetent C57BL/6 mice ranging from 1.9 × 10^4 to 7.8 × 10^5 CFU (24, 25).

Groups of mice were sacrificed at various time points after i.p. inoculation with ca. 5 × 10^5 CFU/mouse of the parental strain and mutant strains, and bacterial loads in blood, lungs, liver, and spleen were quantified (Fig. 2A, top panels). In mice inoculated with parental strain BM4587, bacteria disseminated by 4 h from the peritoneum to blood (8.1 × 10^5 CFU/ml [median values
Growth competition between the parental strain and RND pump-overexpressing strains. (A) For adeABC overexpression, parental A. baumannii BM4587 strain and BM4688[ adeABC] or BM4689[ adeABC] were used. (B) For adeFGH overexpression, strains BM4587 and BM4690[ adeFGH] or BM4691[ adeFGH] were used. (C) For adeIJK overexpression, strains BM4587 and BM4666[ adeIJK] were used. (A to C) For all experiments shown, the strains were mixed at a 1:1 ratio at an initial inoculum of 5 × 10^4 CFU and transferred every 12 h (corresponding to approximately 20 generations) in fresh medium for up to six passages. The competition index (CI) was calculated as the CFU ratio of the resistant and susceptible strains (R/S) at time t, divided by the same R/S at time t_0, and the selection coefficient s was then calculated as the slope of the following linear regression model s = \ln (CI)/(t \times \ln (2)), where t is the number of generations. Every experiment was carried out in duplicate in four independent experiments, and the results are presented as means ± SD (error bars).

shown in parentheses), lung (4 × 10^4 CFU/lung), liver (3.2 × 10^5 CFU/liver), and spleen (6.5 × 10^4 CFU/spleen). Bacterial proliferation, of approximately 1 log_{10} unit, continued by 8 h with median values of CFU/organ of 2.5 × 10^9 per ml of blood, 2.6 × 10^5 per lung, 2.0 × 10^6 per liver, and 5.1 × 10^5 per spleen. Finally, by 24 h, mice either succumbed (less than 10% per experiment) or survived, and organ bacterial counts were very scattered, with median values of CFU/organ of 1.2 × 10^4 per ml of blood, 3.2 × 10^4 in lung, 4.0 × 10^6 in liver, and 1.9 × 10^5 in spleen.

By 4 h after i.p. injection of strain BM4689[ adeABC], the bacterial load per organ increased between 2.3- and 8.4-fold compared to the bacterial load in the parent (Fig. 2A, middle panels). Bacterial multiplication then lagged by 8 h, and counts were significantly lower than those of strain BM4587. At 24 h post-inoculation (p.i.), mean values were similar to those of the group infected with BM4587, as were the numbers of dead mice by 24 h. To further assess the early proliferative attenuation of BM4689[ adeABC], randomly selected colonies obtained ex vivo at each time point were studied, but only nonsignificant differences in in vitro susceptibility tests and in growth rates were observed.

In the mice inoculated with strain BM4666[ adeIJK], initial bacterial dissemination by 4 h to the liver and spleen was not as efficient as that of parental BM4587 with smaller bacterial burden in the spleen and liver. However, bacterial counts increased rapidly to reach similar levels by 24 h (Fig. 2A, bottom panels).

After i.n. inoculation of ca. 5 × 10^7 CFU of strain BM4587, 1.2 × 10^8 CFU/lung was recovered by 4 h and the CFU/lung gradually dropped with median values of 3.0 × 10^7 by 24 h and 1.3 × 10^5 by 48 h (Fig. 2B, top panel). The mice infected with BM4689[ adeABC] or BM4666[ adeIJK] displayed similar bacterial loads in the lungs, and the bacterial loads decreased at the same pace as in mice infected with parental BM4587 (Fig. 2B, middle and bottom panels). Extrapulmonary dissemination, except for occasional recovery in blood, was not observed at any time point.

AdeABC overproduction induces a moderate competitive disadvantage in a systemic infection model. Groups of six mice were inoculated with a 1:1 mixture of strains BM4587 and BM4689[ adeABC] with ca. 5 × 10^7 CFU/mouse for i.p. injection (Fig. 3A) and ca. 5 × 10^7 CFU/mouse for i.n. inoculation (Fig. 3B), and bacterial loads in organs were monitored for 48 h and 72 h, respectively. After i.p. injection, the parental and mutant mixture kept the initial ratio by 8 h p.i. (Fig. 3A). By 24 h, slightly lower loads of BM4689[ adeABC] were observed in the lungs and spleen (5.1 × 10^5 CFU/lung and 6.7 × 10^4 CFU/spleen, respectively) compared to BM4587 (2.1 × 10^6 CFU/lung and 8.3 × 10^4 CFU/spleen, respectively), and a continuous decrease in the relative loads was observed by 48 h (<10 CFU/lung, <10 CFU/liver, and 5.3 × 10^6 CFU/spleen) compared to BM4587 (3.2 × 10^5 CFU/lung, 2.0 × 10^6 CFU/liver, and 6.2 × 10^5 CFU/spleen). Strain BM4689[ adeABC] was moderately less competitive than the parent during the course of systemic infection in agreement with the in vitro reduced growth rate and competitive disadvantage. In the lungs of mice infected by i.n. inoculation with a 1:1 mixture of BM4587 and BM4689[ adeABC], the initial ratio remained constant during the entire experiment, up to 72 h (Fig. 3B), indicating no competitive disadvantage in the pulmonary infection model.

Overproduction of AdeABC resulted in enhanced neutrophil activation after i.n. infection. To study the host immune response induced by pulmonary infections with strains BM4587 and BM4689[ adeABC], bronchoalveolar lavage fluid (BALF) samples were collected from mice 4, 24, and 48 h after i.n. inoculation. Bacterial loads in BALF decreased steadily: from 1.2 × 10^5 to...
Bacterial loads in organs of mice infected with *A. baumannii* BM4587 and its derivatives. (A and B) C57BL/6 female mice were infected with *A. baumannii* strains given i.p. (A) or i.n. (B). (A) Mice infected i.p. were given $5 \times 10^7$ CFU/mouse. Mice infected i.n. were given $5 \times 10^5$ CFU/mouse. In the graphs in the top row in panels A and B, each symbol represents the value for an individual animal infected with strain BM4587 (17 mice infected i.p. and 16 mice infected i.n.), and the gray bar represents the median value of the group of mice. In the graphs in the middle and bottom rows in panels A and B, the means ± SE (error bars) of bacterial loads in organs of mice infected i.p. with either strain BM4689/adeABC (13 mice infected i.p. and 6 mice infected i.n. [middle row]) or strain BM4666/adeIJK (5 mice infected i.p. and 8 mice infected i.n. [bottom row]) and with parental strain BM4587 are indicated. The inoculating doses are indicated by small black arrows on the $y$ axis. Values that are significantly different by two-tailed Mann-Whitney U test are indicated as follows: *, $P < 0.05$; **, $P < 0.01$.

Before infection, the number of total BALF cells was $8.2 \times 10^4 \pm 1.8 \times 10^4$, composed of only macrophages. After i.n. inoculation with strain BM4587, by 4 h the BALF cell number remained constant ($7.4 \times 10^4 \pm 2.3 \times 10^4$ cells/ml) and was mainly composed of neutrophils ($83\%$ of total BALF cells). The number of cells increased from $6.6 \times 10^6 \pm 1.5 \times 10^6$ cells/ml by 24 h to $2.9 \times 10^6 \pm 2.0 \times 10^6$ by 48 h (more than $93\%$ being neutrophils). Infection with BM4689/adeABC induced similar changes in BALF composition. Despite the small difference in neutrophil populations between the groups infected with BM4587 or BM4689/adeABC, significant differences ($P < 0.01$) in the optical density at 450 nm (OD$_{450}$) of myeloperoxidase (MPO), an enzyme with antimicrobial activity, were observed in particular at 48 h i.p., $0.9 \pm 0.4$ and $1.5 \pm 0.3$, respectively (Fig. 4E), indicating more-effective activation of neutrophils by BM4689/adeABC.

To further characterize the immune response induced by infection with the parental or mutant strains, we determined the levels of proinflammatory cytokines in the BALF samples. (Fig. 4F to H). We observed substantial release of all cytokines by bacterial inoculation. In the BALF samples from mice infected with strain BM4587, the keratinocyte chemoattractant protein (KC)/chemokine (C-X-C motif) ligand 1 (CXCL1) peaked by 4 h ($2.4 \times 10^4 \pm 3.1 \times 10^3$ pg/ml) and rapidly decreased by 24 h. The concentration of the released tumor necrosis factor alpha (TNF-$\alpha$) also peaked at 4 h p.i. ($6.4 \times 10^3 \pm 3.6 \times 10^2$ pg/ml) and decreased to $9.4 \times 10^2$ pg/ml by 24 h. In the case of interleukin-6 (IL-6), the production of IL-6 was lower by 4 h but rapidly increased to concentrations higher than those elicited by BM4587 (Fig. 4F and G). The production of IL-6 was lower by 4 h but rapidly increased to concentrations higher than those obtained with the parent, although not reaching statistical significance (Fig. 4H).

After i.p. inoculation, BALF cells were composed of $99\%$ pulmonary macrophages, and the levels of chemokines (the same panel of chemokines) were similar to what is found in noninfected
animals. Bacterial loads in the BALF were under the detection threshold, whereas from 1.1×10^1 to 2.0×10^3 CFU/lung of bacteria was recovered. These results indicate that i.p. inoculation did not result in pulmonary infection and that the bacteria recovered in lungs after systemic infection were mostly from blood.

**Histopathological observations.** In mice infected i.n. with strain BM4587, minimal neutrophil infiltration in alveolar lumens was observed by 4 h (data not shown). The severity of inflammatory lesions increased with time: multifocal mild neutrophil infiltrates were observed 24 h p.i. (Fig. 5), whereas moderate neutrophilic inflammation was noted after 48 h (data not shown). These microscopic changes were parallel and showed a similar time course in both groups inoculated either with strain BM4587 (Fig. 5A and B) or with strain BM4689[adeABC] (Fig. 5C and D). In addition, to assess the location of bacteria in the lesions, lung tissue sections were stained with toluidine blue. Very few bacteria were present in lungs 4 h p.i. with BM4689[adeABC] or BM4587. They were observed with both strains within alveolar macrophages and, very occasionally, in monocytes in blood vessel lumen or free in the alveolar lumen (Fig. 6A and B). Twenty-four hours p.i., bacteria were observed within macrophages and recruited neutrophils (Fig. 6C and D). More infected neutrophils were found in lungs of mice inoculated with BM4689[adeABC] compared to those inoculated with BM4587, corresponding to the elevated level of MPO in the BALF samples from mice infected with BM4689[adeABC]. After 48 h, bacteria were still present in macrophages and/or neutrophils in alveoli following inoculation with BM4587 or BM4689[adeABC]. No microscopic changes or bacteria were observed in the lung at any time point after i.p. inoculation of BM4689[adeABC] or BM4587. By inoculating by this route, we observed bacteria only in blood, and the bacteria were free or within vascular macrophages, in alveoli, or on the alveolar wall.

**DISCUSSION**

Overexpression of chromosomal RND-type efflux systems with broad substrate specificity contributes to MDR in *A. baumannii* (9). Since such strains have emerged as a main concern in clinical settings, particularly in intensive care units, the role of RND pumps in antimicrobial resistance has been studied extensively but not their implications for virulence. We reported that overproduction of AdeABC is mainly responsible for MDR by efflux in clinical isolates (8). We have also generated, from an entirely sequenced susceptible *A. baumannii* clinical strain, a complete set of isogenic mutants overexpressing each of the three main RND systems or with each of the three main RND systems deleted and systematically evaluated the contribution of each Ade pump to antibiotic resistance and biofilm formation (9). We have shown that modulation of expression of the structural genes for these efflux systems led to several alterations in membrane-associated cellular functions, in particular overproduction of AdeABC and AdeIJK resulted in a decrease in biofilm formation (9). Therefore, a biological cost and an impact on virulence could be anticipated.

The fitness cost of a resistance determinant is best studied by comparison of isogenic strains differing only by this determinant (11). Of note, the levels of adeB expression in the mutants were in the same range as those in the 13 MDR clinical isolates (8). The biological cost observed for AdeABC overexpression was significant (2.2 to 3.1%; Fig. 1) but moderate compared to that associated with other resistance mechanisms, such as to colistin (8 to 20%) within the same species (27). The relatively low fitness burden associated with adeABC overexpression could be responsible,
at least in part, for its high prevalence in MDR clinical isolates; in our study, 10 out of 13 MDR clinical isolates overexpressed adeB more than 20-fold (8). Among 444 clinical A. baumannii isolates, adeB overexpression was also the most common, with similar elevated levels, and was generally associated with an MDR phenotype (10). The adeJK-overexpressing mutant had a relatively high fitness decrease (2.2%; Fig. 1). Such strains are found only occasionally: in a panel of 34 A. baumannii strains (28), a 3- to 4-fold increase in adeJ expression due to a truncated AdeN regulator was observed in three strains (unpublished data). In the study of 444 clinical A. baumannii isolates, the levels of adeJ expression in nine strains were ca. 10 times higher than the levels in the susceptible ATCC 17978 strain (10). Interestingly, strain BM4719[ΔadeJ] displayed a slightly diminished growth rate (0.98 ± 0.01), while the growth rates of the mutants with either adeB or adeG deleted were similar to that of the parent (Table 1), suggesting that constitutive expression of adeJK is critical for A. baumannii physiology. The adeFGH-overexpressing mutant displayed the highest fitness cost (4.3%), but the expression rate of the system in this in vitro mutant was very high (Table 1), a level that has not been observed in clinical isolates (from 2- to 15-fold) (8, 10). Overall, there was good agreement between the burden imposed on the host by overexpression of an efflux system and its relative occurrence in clinical isolates. In the mutants, as well as in clinical strains, overexpression of the structural genes for the efflux pumps resulted from regulatory mutations. Thus, the biological cost could be due to

FIG 4 Bronchoalveolar lavage fluid analysis. BALF samples from groups of five C57BL/6 mice infected i.n. with 5 × 10^7 CFU of strain BM4587 (black) or strain BM4689[ΔadeABC] (white) were analyzed. (A) Bacteria in the BALF samples were quantified. (B to D) Total BALF cells/ml (B), macrophages (C), and neutrophils (D) were quantified by examining Hema 3-stained cytospin slides. (E to H) Myeloperoxidase was measured by enzyme immunometric assay (E), and proinflammatory cytokine/chemokine production was determined for KC/CXCL1 (F), TNF-α (G), and IL-6 (H) by using sandwich ELISA. Data are presented as means plus SD. Values that are significantly different by one-way ANOVA are indicated by bars and asterisks as follows: *, P < 0.05; **, P < 0.01.

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excessive energy consumption by the pumps or too efficient export of molecules beneficial to the host. Alternatively, the regulatory genes may also control other genes that could be responsible for the fitness burden.

A recent approach to study fitness and virulence of pathogens is the use of saturated transposon (Tn) libraries to identify genes that contribute to optimal fitness and therefore virulence in vivo. As many as 157 A. baumannii genes required for persistence in a mouse pneumonia model were identified after i.n. injection of a transposon insertion library of strain ATCC 17978 (23). In the majority of cases, insertion led to a decrease in persistence. Insertion in two genes (A1S_1649 and A1S_1801) predicted to encode RND efflux pump determinants resulted in a decrease in the fitness of the mutants in mice (23). A similar result was obtained after i.p. injection of isogenic mutants of virulent strain AB5075 with Tn insertions in the same genes (26). However, none of these genes were related to the genes encoding the three components of the three efflux systems of our study. Growth analysis of a transposon library of AB5057 in the Galleria mellonella larva model detected 300 genes required for bacterial survival and identified the genes for AdeIJK (29). Of note, the adef deletion mutant was the only one with a slightly diminished growth rate, but unfortunately, it was not studied in vivo. Proteomic analysis of bacterial membranes indicated that, in susceptible strains, the AdeABC system is detected at very low levels, whereas all three proteins, AdeI, AdeJ, and AdeK, are easily detected in agreement with their basal constitutive synthesis (9). This observation points to the notion that the study of virulence using knockout transposons will identify only the genes that are expressed under the assay conditions. Genes with expression under tight regulation, such as adeABC, and the majority of the antibiotic resistance determinants will not be detected unless they are negatively regulated or when their induced expression results in a selective advantage under the experimental conditions used (30), which was not the case in this study.

The mouse pulmonary model is widely used to study A. baumannii virulence, although most isolates can induce only a self-limiting pneumonia with slow local bacterial multiplication. We have used both intraperitoneal and intranasal routes of infection with the same set of strains and observed a diminished competition of strain BM4689[adeABC] after i.p. inoculation, but not after i.n. inoculation (Fig. 3). However, the pneumonia model allows us to study the most frequent mode of infection in humans and to distinguish strains of various degrees of virulence and has revealed the important role of neutrophils, which are rapidly recruited for control of the respiratory infection (22). Only minor differences in bacterial growth kinetics were observed between parental and RND pump overexpression mutants (Fig. 2). However, BALF analysis and histopathological observations indicated an increased host defense response to BM4689[adeABC]. In BALF samples from mice infected with the mutant, a slight increase of neutrophils was observed 48 h p.i. (Fig. 4). TNF-α and KC/CLCX-1 contribute to neutrophil migration within the lung, while elevated IL-6 levels are a response to excessive inflammation (31). Increased MPO activity recovered in the BALF samples 48 h p.i. is the sign of neutrophil degranulation following activation (32). Histopathological observations showed that there were more
infected neutrophils at 24 h after mutant infection (Fig. 5 and 6). Taken together, pulmonary infection by the adeABC-overexpressing mutant resulted in a small delay in the onset but in a more severe inflammation in the lung secondary to a higher activation of recruited neutrophils. These results indicate that persistence in the lung is dissociated from virulence, the latter being best appreciated by immunological responses of the host and histological analysis. This observation may explain at least in part, the success of adeABC-overexpressing MDR A. baumannii in ventilator-associated pneumonia.

MATERIALS AND METHODS

Bacterial strains and growth conditions. Drug-susceptible parental A. baumannii BM4587 (33) and its mutant derivatives (9) are described in Table 1. Bacteria were grown in brain heart infusion (BHI), Luria-Bertani (LB), or Mueller-Hinton (MH) broth or agar at 37°C. Antibiotic susceptibility was assessed by disk diffusion on MH agar according to the CLSI guidelines (34).

Determination of growth rates and growth competitions. Growth rates were determined as described previously (35) in microplates coupled to a Multiscan spectrophotometer (Thermo Scientific). The strains were grown overnight, and the cultures were diluted to inoculate \( \sim 1 \times 10^8 \) bacteria into 200 \( \mu l \) of LB in a 96-well microplate which was incubated with shaking. Absorbance was measured at 595 nm every 3 min. Each culture was replicated three times in the same microplate. Growth rates, measured in five independent experiments, were determined at the beginning of the exponential phase, and relative growth rates were calculated as the ratio of the growth rate of the strains versus that of the parent. Competition experiments were performed in cocultures with \( 5 \times 10^6 \) of the susceptible strain and the resistant strain mixed in 10 ml of LB at an initial ratio of 1:1 and grown for 12 h (about 20 generations). The mixed culture was transferred to fresh LB by 10^6-fold serial dilution every 12 h for up to six passages. The total number of viable cells was determined at the end of every transfer on nonselective plates, and the proportion of resistant strains was determined by replica plating 500 to 1,000 colonies on MH agar and MH agar supplemented with gentamicin (4 \( \mu g/ml \) and 12 \( \mu g/ml \)), clindamycin (200 \( \mu g/ml \)), or tetracycline (2 \( \mu g/ml \)), depending on the pair of strains studied. Each experiment was carried out in duplicate and performed four times independently.

Mice. Specific-pathogen-free female C57BL/6J mice were purchased from the Centre d’Elevage R. Janvier (Le Genest Saint-Isle, France) and were used at 6 to 8 weeks of age with a body weight of 17 to 19 g. The animals were housed under specific-pathogen-free conditions in a small animal containment and fed ad libitum with sterile water and certified mouse chow. All animal studies were approved by the Pasteur Institute Safety Committee (protocol 12.062) in accordance with French and European guidelines, and mice were cared for in accordance with the Pasteur Institute guidelines in compliance with the European Animal Welfare regulations.

Intraperitoneal and intranasal inoculations. Bacteria grown to the exponential phase (optical density at 600 nm [OD_{600}] of 0.8 to 0.9) in BHI broth were centrifuged, resuspended in phosphate-buffered saline (PBS), and used immediately. To determine the intraperitoneal (i.p.) 50% lethal
doe (LD₅₀), five 3-fold dilutions from 2.0 × 10⁶ to 5.5 × 10⁹ CFU of strain BM4587 were injected i.p. to groups of six 7-week-old C57BL/6 female mice, and the mice were monitored for 7 days. All survivors showed a normal clinical score by 4 days postinoculation (p.i.) (see Fig. S1 in the supplemental material). Mice given the highest inoculum died in 1 day, and mice given the two lowest inocula survived until the end of the experiment. Survival rates of the mice given two intermediate inocula were 83.3% and 16.7%, respectively. The BM4587 i.p. LD₅₀ value, calculated by probit analysis, was 4.9 × 10⁵ CFU/mouse. Unless otherwise specified, mice were inoculated i.p. with ca. 5 × 10⁶ A. baumannii in 100 μl of PBS.

For the intranasal (i.n.) route, groups of mice anesthetized by i.p. injection of a mixture of ketamine (Imalgene 1000; Merial) and xylazine (Rompun; Bayer) were inoculated with six inocula from 2.5 × 10⁶ to 1 × 10⁹ CFU in 50 μl of PBS by dropping the inoculum on the tip of the nose. The mice were clinically monitored and sacrificed by 4, 24, and 48 h. By 12 h, infected mice had ruffled fur but they were all alive, and all mice had recovered by 24 h. Bacteria were not recovered from the blood, spleen, or liver of mice at any time point, and bacterial load in the lungs declined by more than 2 log₅ units between 4 h and 48 h. Five log₅ units of CFU/lung were harvested by 48 h after an inoculum of 1 × 10⁶ CFU. Therefore, ca. 5 × 10⁶ of A. baumannii was used for i.n. inoculation.

For bacterial growth kinetics, mice in each group were euthanized by i.p. administration of a lethal dose of pentobarbital at the indicated times. Systemic blood was collected by cardiac puncture, and the lungs, liver, and spleen were aseptically removed, weighed, and homogenized in sterile PBS using a Glas-Col homogenizer. Aliquots of 10-fold serial dilutions of the homogenates were cultured on BH agar to quantify the number of viable A. baumannii in the respective organs.

For in vivo competition, mice were inoculated with a 1:1 mixture of parental strain BM4587 and mutant strain BM4689[AdeABC] (BM4689 overexpressing adeABC] at a total of ca. 5 × 10⁶ and 5 × 10⁵ bacteria for i.p. and i.n. inoculation, respectively. The proportion of resistant strains was determined by replica plating on plates containing gentamicin (12 μg/ml).

Bronchoalveolar lavage fluid analysis. The trachea of euthanized mice were cannulated, and the lungs were washed four times with 0.3 ml PBS, yielding ca. 2 ml of bronchoalveolar lavage fluid (BALF). Cytospin slides of ca. 1 × 10⁶ BALF cells were prepared using a cytospin centrifuge and stained with Hema 3 Stat (Fisher, Pittsburgh, PA). The total and differential numbers of neutrophils and macrophages were determined by examining 200 cells. Cell-free BALF obtained after centrifugation (1,500 rpm or 5,000 rpm for 10 min) was used for chemokine measurement. The level of myeloperoxidase (MPO) in BALF was measured by enzyme immunomodified assay, and murine interleukin-6 (IL-6), tumor necrosis factor alpha (TNF-α), and keratinocyte chemotactant protein (KC) in BALF were quantified by a sandwich enzyme-linked immunosorbent assay (ELISA) with DuoSet ELISA development kits (R&D Systems, Minneapolis, MN) according to the manufacturer’s instructions.

Histopathological examination. Groups of five mice were inoculated i.n. or i.p. with A. baumannii BM4587 or BM4689[AdeABC]. Mice were sacrificed by i.p. administration of a lethal dose of pentobarbital on 4, 24, and 48 h after inoculation. The lungs were perfused by intratracheal injection of 0.5 ml of 4% neutral buffered formalin. The lungs were retrieved and routinely embedded in paraffin. The sections were 4 μm thick and stained with hematoxylin and eosin (H&E) or toluidine blue. All sections were examined by light microscopy. A semiquantitative scoring system was applied to evaluate the microscopic changes. The scores were as follows: 1 for minimal, 2 for mild, 3 for moderate, 4 for marked, and 5 for severe. Microphotographs (Fig. 5 and 6) are representative of the histological findings in each group.

Statistical analysis. Data are presented as means ± standard deviations (SD) or standard errors (SE) for each group as specified. Differences in quantitative measurement were assessed by Student’s t test or by two-way Mann-Whitney analysis of variance (ANOVA) followed by the Shapiro-Wilk test. A P value of <0.05 was considered significant.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at http://mbio.asm.org/lookup/suppl/doi:10.1128/mBio.00697-16/-/DCSupplemental.

Figure S1, DOCX file, 0.1 MB.

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