

Genome Sequences of Multidrug-Resistant Acinetobacter baumannii Strains from Nosocomial Outbreaks in Japan

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Acinetobacter baumannii has emerged worldwide as an important nosocomial pathogen in medical institutions. Here, we present the draft genome sequences of *A. baumannii* strains MRY09-0642, MRY10-0558, and MRY12-0277 that were isolated from nosocomial outbreaks in Japan between 2008 and 2012 and that are resistant to antimicrobial agents, including carbapenems, fluoroquinolones, and aminoglycosides.

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cinetobacter baumannii often causes infections in hospitalized immunocompromised patients (1). A. baumannii strains belonging to international clone II (IC2)/sequence type 2 (ST2), the most prevalent epidemic lineage, are associated with multidrug resistance and often cause nosocomial outbreaks (2). Since 2000, carbapenem-resistant and multidrug-resistant A. baumannii strains have emerged and their prevalence has increased worldwide. In some countries, carbapenem-resistant strains have reached a prevalence of >50%, becoming a serious public health threat (3). According to national surveillance data from Japan Nosocomial Infections Surveillance (JANIS) conducted by the Ministry of Health, Labour and Welfare (http://www.nih-janis.jp /english/), carbapenem resistance among Acinetobacter spp. in Japan remains much lower than in other countries (approximately 2% of imipenem resistance in Japan in 2012) (S. Suzuki, unpublished data). However, the prevalence tends to increase gradually, and nosocomial outbreaks of A. baumannii IC2 infections have occasionally occurred in Japan.

To date, whole-genome sequences of *A. baumannii* strains isolated in Japan have not been available in GenBank. In this report, we announce the availability of the draft genome sequences of *A. baumannii* MRY09-0642, MRY10-0558, and MRY12-0277, which caused nosocomial outbreaks in geographically different medical institutions in Japan in 2008, 2010, and 2012, respectively. These isolates were classified as IC2 by multilocus sequence typing (4), whereas they showed distinct ApaI fragment patterns in pulsed-field gel electrophoresis. Whole-genome shotgun (WGS) sequencing of the *A. baumannii* strains was performed using the Roche 454 pyrosequencing platform (500-bp insert size). Reads were assembled with Newbler assembler version 2.3 (Roche), using *A. baumannii* Taiwanese strain MDR-TJ (5) as the reference.

The draft genome sequences of *A. baumannii* MRY09-0642, MRY10-0558, and MRY12-0277 consist of 147, 77, and 106 contigs, respectively, yielding total sequences for each strain of

3,746,543, 3,782,742, and 3,829,745 bp, with N_{50} contig sizes of 64,650, 164,570, and 91,207 bp, respectively. Their mean G+C content is 39.0% \pm 0.1%. A total of 3,645, 3,604, and 3,735 coding genes for MRY09-0642, MRY10-0558, and MRY12-0277, respectively, were detected by the RAST server (http://rast.nmpdr.org) (6). Acquired antimicrobial resistance genes in the WGS data were identified using a Web-based tool, ResFinder version 1.3 (http://cge.cbs.dtu.dk/services/ResFinder/) (7). *A. baumannii* MRY09-0642, MRY10-0558, and MRY12-0277 carry OXA-51-like β -lactamase genes, which predominantly confer carbapenem resistance in *A. baumannii* (8), found as bla_{OXA-82} in contig 00034, and bla_{OXA-66} in contig 00056, respectively.

A more-detailed report of the drug resistance and virulence phenotypes of these three *A. baumannii* strains will be included in a future publication. Access to these genome sequences and their comparative analyses with other epidemic and nonepidemic strains will facilitate additional comprehensive bioinformatics and phylogenetic analyses, thus expanding our understanding of the global public health problem caused by this nosocomial pathogen.

Nucleotide sequence accession numbers. These WGS projects have been deposited at DDBJ/EMBL/GenBank under the accession no. BASA00000000, BASB00000000, and BASC00000000. The versions described in this report are the first versions, accession no. BASA01000000, BASB01000000, and BASC01000000 for *A. baumannii* strains MRY09-0642, MRY10-0558, and MRY12-0277, respectively.

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REFERENCES

- 1. Dijkshoorn L, Nemec A, Seifert H. 2007. An increasing threat in hospitals: multidrug-resistant *Acinetobacter baumannii*. Nat. Rev. Microbiol. 5:939–951.
- Diancourt L, Passet V, Nemec A, Dijkshoorn L, Brisse S. 2010. The population structure of *Acinetobacter baumannii*: expanding multiresistant clones from an ancestral susceptible genetic pool. PLoS One 5:e10034. doi: 10.1371/journal.pone.0010034.
- Laxminarayan R, Klugman KP. 2011. Communicating trends in resistance using a drug resistance index. BMJ Open 1:e000135. doi:10.1136/bmjopen -2011-000135.
- 4. Matsui M, Suzuki S, Suzuki M, Arakawa Y, Shibayama K. 2013. Rapid discrimination of *Acinetobacter baumannii* international clone II lineage by

pyrosequencing SNP analyses of bla_{OXA} 51-like genes. J. Microbiol. Methods 94:121–124.

- Gao F, Wang Y, Liu YJ, Wu XM, Lv X, Gan YR, Song SD, Huang H. 2011. Genome sequence of *Acinetobacter baumannii* MDR-TJ. J. Bacteriol. 193:2365–2366.
- Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, Meyer F, Olsen GJ, Olson R, Osterman AL, Overbeek RA, McNeil LK, Paarmann D, Paczian T, Parrello B, Pusch GD, Reich C, Stevens R, Vassieva O, Vonstein V, Wilke A, Zagnitko O. 2008. The RAST server: rapid annotations using subsystems technology. BMC Genomics 9:75. doi:10.1186/1471-2164-9-75.
- Zankari E, Hasman H, Cosentino S, Vestergaard M, Rasmussen S, Lund O, Aarestrup FM, Larsen MV. 2012. Identification of acquired antimicrobial resistance genes. J. Antimicrob. Chemother. 67:2640–2644.
- Lee JH, Choi CH, Kang HY, Lee JY, Kim J, Lee YC, Seol SY, Cho DT, Kim KW, Song do Y, Lee JC. 2007. Differences in phenotypic and genotypic traits against antimicrobial agents between *Acinetobacter baumannii* and *Acinetobacter* genomic species 13TU. J. Antimicrob. Chemother. 59: 633–639.