

Complete Genome Sequence of *Pseudomonas aeruginosa* PA1, Isolated from a Patient with a Respiratory Tract Infection

Shuguang Lu, Shuai Le, Gang Li, Mengyu Shen, Yinling Tan, Xia Zhao, Jing Wang, Wei Shen, Keke Guo, Yuhui Yang, Hongbin Zhu, Shu Li, Ming Li, Junmin Zhu, Xiancai Rao, Fuquan Hu

Department of Microbiology, College of Basic Medical Science, Third Military Medical University, Chongqing, People's Republic of China

We report the 6,498,072-bp complete genome sequence of *Pseudomonas aeruginosa* PA1, which was isolated from a patient with a respiratory tract infection in Chongqing, People's Republic of China. Whole-genome sequencing was performed using single-molecule real-time (SMRT) technology, and *de novo* assembly revealed a single contig with 396-fold sequence coverage.

Received 20 October 2015 Accepted 23 October 2015 Published 10 December 2015

Citation Lu S, Le S, Li G, Shen M, Tan Y, Zhao X, Wang J, Shen W, Guo K, Yang Y, Zhu H, Li S, Li M, Zhu J, Rao X, Hu F. 2015. Complete genome sequence of *Pseudomonas aeruginosa* PA1, isolated from a patient with a respiratory tract infection. *Genome Announc* 3(6):e01453-15. doi:10.1128/genomeA.01453-15.

Copyright © 2015 Lu et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 3.0 Unported license](http://creativecommons.org/licenses/by/4.0/).

Address correspondence to Fuquan Hu, hufuquan2009@aliyun.com.

Pseudomonas aeruginosa is a Gram-negative rod-shaped gamma-proteobacterium that grows in a wide range of ecological niches, such as soil, marshes, and coastal marine habitats, as well as on plant and animal tissues (1–3). As an opportunistic pathogen, *P. aeruginosa* causes a wide range of syndromes in humans that can vary from local to systemic, and sometimes its infection is life-threatening (4). *P. aeruginosa* is a significant pathogen associated with infections of burn victims, urinary tract infections in catheterized patients, and respiratory tract infections (2, 5). When infecting immunocompromised or cystic fibrosis (CF) patients, *P. aeruginosa* can lead to deadly pneumonia (6, 7). Notably, intrinsic drug resistance of *P. aeruginosa* makes it difficult to treat *P. aeruginosa* infections with antibiotics (8, 9).

As of 20 October 2015, 27 complete genome sequences of different *P. aeruginosa* stains have been released from the GenBank database (10) (<http://www.ncbi.nlm.nih.gov/genome/genomes/187>). Different *P. aeruginosa* genomes share a remarkable amount of sequence similarity, despite having been isolated from various niches or different clinical origins (11–13). The *P. aeruginosa* pan-genome consists of at least 4,000 core genes, approximately 10,000 accessory genes, and 30,000 or more rare genes that are present in only a few strains or clonal complexes (4). These genome sequences have provided insight into virulence, drug resistance, and biofilm formation that are related to the pathogenicity of *P. aeruginosa* (2, 14, 15). However, hitherto the genomic information of *P. aeruginosa* is still very limited for researchers to analyze, compare, and evaluate the characteristics of the species. Thus, more *P. aeruginosa* genome sequences are required to explore potential ways to control this versatile opportunistic pathogen.

P. aeruginosa PA1 was originally isolated from a respiratory tract infection patient in Chongqing, China. It has a lytic bacteriophage that belongs to the PaP1-like phage genus (16). The genomic DNA of *P. aeruginosa* PA1 was extracted from the stationary-phase cultures grown in LB broth and purified using the TIANamp bacteria DNA kit (Tiangen Biotech, Beijing, China). PacBio single-molecule real-time (SMRT) sequencing of

the PA1 genome was carried out at the Institute of Medicinal Plant Development (IMPLAD) (Beijing, China) using the PacBio RS II Instrument (Pacific Biosciences, Menlo Park, CA, USA) (17, 18). Libraries of 5-kb were constructed and 4 SMRT cells of the libraries were sequenced with 180-min movies. *De novo* assembly was performed using RS_HGAP_Assembly v. 2.0 (19), revealing a single contig with an average sequence coverage of 396-fold. The length of the PA1 genome is 6,498,072 bp, with an average G+C content of 66.35%. Genome annotation of *P. aeruginosa* PA1 was performed through the NCBI Prokaryotic Genome Annotation Pipeline (20) (released 2013) (http://www.ncbi.nlm.nih.gov/genome/annotation_prok/).

Nucleotide sequence accession number. The complete genome sequence of *P. aeruginosa* PA1 has been deposited in GenBank under the accession number CP004054.

ACKNOWLEDGMENTS

This work was supported by grant 31400163 from the National Natural Science Foundation of China.

We thank Leigh A. Riley from the GenBank direct submission staff for helping us annotate the *P. aeruginosa* PA1 genome.

REFERENCES

1. Hardal C, Edberg SC. 1997. *Pseudomonas aeruginosa*: assessment of risk from drinking water. *Crit Rev Microbiol* 23:47–75. <http://dx.doi.org/10.3109/10408419709115130>.
2. Stover CK, Pham XQ, Erwin AL, Mizoguchi SD, Warrener P, Hickey MJ, Brinkman FS, Hufnagle WO, Kowalik DJ, Lagrou M, Garber RL, Goltry L, Tolentino E, Westbrock-Wadman S, Yuan Y, Brody LL, Coulter SN, Folger KR, Kas A, Larbig K, Lim R, Smith K, Spencer D, Wong GK, Wu Z, Paulsen IT, Reizer J, Saier MH, Hancock RE, Lory S, Olson MV. 2000. Complete genome sequence of *Pseudomonas aeruginosa* PAO1, an opportunistic pathogen. *Nature* 406:959–964. <http://dx.doi.org/10.1038/35023079>.
3. Nakano K, Terabayashi Y, Shiroma A, Shimoji M, Tamotsu H, Ashimine N, Ohki S, Shinzato M, Teruya K, Satou K, Hirano T. 2015. First complete genome sequence of *Pseudomonas aeruginosa* (Schroeter 1872) Migula 1900 (DSM 50071T), determined using PacBio single-molecule real-time technology. *Genome Announc* 3(4):e00932-15. <http://dx.doi.org/10.1128/genomeA.00932-15>.

4. Hilker R, Munder A, Klockgether J, Losada PM, Chouvarine P, Cramer N, Davenport CF, Dethlefsen S, Fischer S, Peng H, Schönfelder T, Türk O, Wiehlmann L, Wöbeling F, Gulbins E, Goesmann A, Tümmler B. 2015. Interclonal gradient of virulence in the *Pseudomonas aeruginosa* pan-genome from disease and environment. *Environ Microbiol* 17:29–46. <http://dx.doi.org/10.1111/1462-2920.12606>.
5. Wibberg D, Tielen P, Narten M, Schobert M, Blom J, Schatschneider S, Meyer A, Neubauer R, Albersmeier A, Albaum S, Jahn M, Goesmann A, Vorhölter F, Pühler A, Jahn D. 2015. Genome sequence of the urethral isolate *Pseudomonas aeruginosa* RN21. *Genome Announc* 3(4):e00788-15. <http://dx.doi.org/10.1128/genomeA.00788-15>.
6. Loré NI, Cigana C, De Fino I, Riva C, Juhas M, Schwager S, Eberl L, Bragonzi A. 2012. Cystic fibrosis-niche adaptation of *Pseudomonas aeruginosa* reduces virulence in multiple infection hosts. *PLoS One* 7:e35648. <http://dx.doi.org/10.1371/journal.pone.0035648>.
7. Stuart B, Lin JH, Mogayzel PJ, Jr. 2010. Early eradication of *Pseudomonas aeruginosa* in patients with cystic fibrosis. *Paediatr Respir Rev* 11:177–184. <http://dx.doi.org/10.1016/j.prrv.2010.05.003>.
8. Breidenstein EBM, de la Fuente-Núñez C, Hancock REW. 2011. *Pseudomonas aeruginosa*: all roads lead to resistance. *Trends Microbiol* 19: 419–426. <http://dx.doi.org/10.1016/j.tim.2011.04.005>.
9. Fernandez L, Hancock REW. 2012. Adaptive and mutational resistance: role of porins and efflux pumps in drug resistance. *Clin Microbiol Rev* 25:661–681. <http://dx.doi.org/10.1128/CMR.00043-12>.
10. Benson DA, Clark K, Karsch-Mizrachi I, Lipman DJ, Ostell J, Sayers EW. 2015. GenBank. *Nucleic Acids Res* 43:D30–D35. <http://dx.doi.org/10.1093/nar/gku1216>.
11. Lee DG, Urbach JM, Wu G, Liberati NT, Feinbaum RL, Miyata S, Diggins LT, He J, Saucier M, Deziel E, Friedman L, Li L, Grills G, Montgomery K, Kucherlapati R, Rahme LG, Ausubel FM. 2006. Genomic analysis reveals that *Pseudomonas aeruginosa* virulence is combinatorial. *Genome Biol* 7:R90. <http://dx.doi.org/10.1186/gb-2006-7-10-r90>.
12. Haenni M, Hocquet D, Ponsin C, Cholley P, Guyeux C, Madec J, Bertrand X. 2015. Population structure and antimicrobial susceptibility of *Pseudomonas aeruginosa* from animal infections in France. *BMC Vet Res* 11:9. <http://dx.doi.org/10.1186/s12917-015-0324-x>.
13. Wiehlmann L, Wagner G, Cramer N, Siebert B, Gudowius P, Morales G, Kohler T, van Delden C, Weinel C, Slickers P, Tummeler B. 2007. Population structure of *Pseudomonas aeruginosa*. *Proc Natl Acad Sci USA* 104:8101–8106. <http://dx.doi.org/10.1073/pnas.0609213104>.
14. Gao C, Hu C, Ma C, Su F, Yu H, Jiang T, Dou P, Wang Y, Qin T, Lv M, Xu P. 2012. Genome sequence of the lactate-utilizing *Pseudomonas aeruginosa* strain xmg. *J Bacteriol* 194:4751–4752. <http://dx.doi.org/10.1128/JB.00943-12>.
15. Roy PH, Tetu SG, Larouche A, Elbourne L, Tremblay S, Ren Q, Dodson R, Harkins D, Shay R, Watkins K, Mahamoud Y, Paulsen IT. 2010. Complete genome sequence of the multiresistant taxonomic outlier *Pseudomonas aeruginosa* PA7. *PLoS One* 5:e8842. <http://dx.doi.org/10.1371/journal.pone.0008842>.
16. Lu S, Le S, Tan Y, Zhu J, Li M, Rao X, Zou L, Li S, Wang J, Jin X, Huang G, Zhang L, Zhao X, Hu F. 2013. Genomic and proteomic analyses of the terminally redundant genome of the *Pseudomonas aeruginosa* phage PaP1: establishment of genus PaP1-like phages. *PLoS One* 8:e62933. <http://dx.doi.org/10.1371/journal.pone.0062933>.
17. Roberts RJ, Carneiro MO, Schatz MC. 2013. The advantages of SMRT sequencing. *Genome Biol* 14:405.
18. Gulati A, Swarnkar MK, Vyas P, Rahi P, Thakur R, Thakur N, Singh AK. 2015. Complete genome sequence of the rhizobacterium *Pseudomonas trivialis* strain IHBB745 with multiple plant growth-promoting activities and tolerance to desiccation and alkalinity. *Genome Announc* 3(5): e00943-15. <http://dx.doi.org/10.1128/genomeA.00943-15>.
19. Chin C, Alexander DH, Marks P, Klammer AA, Drake J, Heiner C, Clum A, Copeland A, Huddleston J, Eichler EE, Turner SW, Korlach J. 2013. Nonhybrid, finished microbial genome assemblies from long-read SMRT sequencing data. *Nat Methods* 10:563–569. <http://dx.doi.org/10.1038/nmeth.2474>.
20. Angiuoli SV, Gussman A, Klimke W, Cochrane G, Field D, Garrity GM, Kodira CD, Kyripides N, Madupu R, Markowitz V, Tatusova T, Thomson N, White O. 2008. Toward an online repository of standard operating procedures (SOPs) for (Meta) genomic annotation. *Oomics J Integr Biol* 12:137–141. <http://dx.doi.org/10.1089/omi.2008.0017>.