




First Complete *Providencia rettgeri* Genome Sequence, the NDM-1-Producing Clinical Strain RB151

R. Alejandro Marquez-Ortiz,^{a,b} Leanne Haggerty,^b Eby M. Sim,^b Carolina Duarte,^c Betsy E. Castro-Cardozo,^a Mauricio Beltran,^c Sandra Saavedra,^c Natasha Vanegas,^{a,b} Javier Escobar-Perez,^a  Nicola K. Petty^b

Bacterial Molecular Genetics Laboratory, Universidad El Bosque, Bogotá, Colombia^a; The ithree Institute, University of Technology Sydney, Sydney, Australia^b; Grupo de Microbiología, Instituto Nacional de Salud, Bogotá, Colombia^c

ABSTRACT *Providencia rettgeri* is an opportunistic bacterial pathogen of clinical significance due to its association with urinary tract infections and multidrug resistance. Here, we report the first complete genome sequence of *P. rettgeri*. The genome of strain RB151 consists of a 4.8-Mbp chromosome and a 108-kbp *bla*_{NDM-1}-positive plasmid.

Providencia rettgeri is an opportunistic human pathogen mainly associated with urinary tract infections (1, 2). A Gram-negative member of the *Enterobacteriaceae*, *P. rettgeri* is also known to cause diarrhea, meningitis, eye infections, and bacteremia in both hospital and community settings (1–5). *P. rettgeri* is intrinsically resistant to several antibiotics (5, 6), but notably, many recent independent isolates have been found to be carbapenemase producers carrying the New Delhi metallo- β -lactamase (NDM) gene *bla*_{NDM-1} (7–9). To date, there are only a few draft genomes of *P. rettgeri* available in the public databases. Here, we report the first complete genome sequence of *P. rettgeri*, that of a multidrug-resistant clinical isolate carrying the *bla*_{NDM-1} gene.

P. rettgeri RB151 was isolated in 2013 from a urine sample from a 58-year-old female patient, diagnosed with a urinary tract infection in the emergency department of a tertiary hospital in Bucaramanga, Colombia (7). Total genomic DNA was extracted using the UltraClean microbial DNA isolation kit (Mo Bio Laboratories, Inc.). A 20-kb BluePip-pin (Sage Science) size-selected SMRTbell library was constructed and sequenced using one single-molecule real-time (SMRT) cell with P6-C4 chemistry on the PacBio RSII platform (Pacific Biosciences, CA). The resulting 167,518 reads, which had an *N*₅₀ read length of 10,167 bp, were *de novo* assembled using the RS_HGAP_Assembly.3 protocol implemented in SMRT Analysis version 2.3 (10), into two contigs with a total genome length of 4.9 Mbp. The filtered subreads were then mapped to the assembly using BWA-MEM (11), revealing an average coverage of 176 \times . The assembly was manually checked using Tablet (12), and low-coverage misassembled terminal repeat sequences, a known artifact of HGAP assembly (10, 13), were manually trimmed and removed from each contig. The final sequences were manually reordered so that the linear representation of each circular contig started at *dnaA* (chromosome) and *repA* (plasmid). The final assembly was verified using Circlator version 1.4.0 (13) and Artemis Comparison Tool version 13 (14). The genome was annotated using Prokka version 1.11 (15), and the antibiotic resistance genes were identified using ARIBA (<https://github.com/sanger-pathogens/ariba>).

The complete genome of *P. rettgeri* RB151 has an average G+C content of 41.7% and consists of a 4,780,676-bp chromosome and a 108,417-bp NDM-1-encoding plasmid (pRB151-NDM). The automated genome annotation predicted 4,497 coding se-

Received 1 November 2016 Accepted 14 November 2016 Published 19 January 2017

Citation Marquez-Ortiz RA, Haggerty L, Sim EM, Duarte C, Castro-Cardozo BE, Beltran M, Saavedra S, Vanegas N, Escobar-Perez J, Petty NK. 2017. First complete *Providencia rettgeri* genome sequence, the NDM-1-producing clinical strain RB151. *Genome Announc* 5: e01472-16. <https://doi.org/10.1128/genomeA.01472-16>.

Copyright © 2017 Marquez-Ortiz et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to R. Alejandro Marquez-Ortiz, rmarquez@unbosque.edu.co, or Nicola K. Petty, nicola.petty@uts.edu.au.

quences (CDSs), 22 rRNAs, 77 tRNAs, and one transfer-messenger RNA (tmRNA). The antimicrobial resistome of the RB151 chromosome evaluated using ARIBA included resistance genes to aminoglycosides [*aac(3)-lia*, *armA*, and *aacA4*], β -lactams (*bla*_{TEM-1B} and *bla*_{OXA-2}), fluoroquinolones [*aac(6')/lb-cr*], sulfonamides (*sul1* and *sul2*), and trimethoprim (*dfrA31*). The plasmid pRB151-NDM only contained the *bla*_{NDM-1} gene, which confers resistance to β -lactams.

As the first complete genome of *P. rettgeri*, this genome sequence will be a useful reference genome and could be utilized to contribute further insights into this species.

Accession number(s). The complete genome of *Providencia rettgeri* RB151 has been deposited in DDBJ/EMBL/GenBank under the GenBank accession numbers [CP017671](#) (chromosome) and [CP017672](#) (plasmid pRB151-NDM). The version described in this paper is the first version.

ACKNOWLEDGMENTS

R.A.M.-O. was supported by Colciencias Estudios de Doctorado en Colombia Fellowship 567-2012. This work was funded by Colciencias (grant FP44842-175-2016) and Vicerrectoría de Investigaciones, Universidad El Bosque (grant PCI-2012-330). The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

R.A.M.-O., N.V., J.E.-P., and N.K.P. designed the study; R.A.M.-O. and N.K.P. performed the research and analyzed the data; L.H., E.M.S., C.D., B.E.C.-C., M.B., and S.S. contributed new methods/analytical tools; and R.A.M.-O. and N.K.P. wrote the paper.

REFERENCES

- O'Hara CM, Brenner FW, Miller JM. 2000. Classification, identification, and clinical significance of *Proteus*, *Providencia*, and *Morganella*. Clin Microbiol Rev 13:534–546. <https://doi.org/10.1128/CMR.13.4.534-546.2000>.
- Wie SH. 2015. Clinical significance of *Providencia* bacteremia or bacteriuria. Korean J Intern Med 30:167–169. <https://doi.org/10.3904/kjim.2015.30.2.167>.
- Yoh M, Matsuyama J, Ohnishi M, Takagi K, Miyagi H, Mori K, Park KS, Ono T, Honda T. 2005. Importance of *Providencia* species as a major cause of travellers' diarrhoea. J Med Microbiol 54:1077–1082. <https://doi.org/10.1099/jmm.0.45846-0>.
- Maiti TK, Singh VK, Pandey P. 2013. *Providencia rettgeri*: an unusual cause of central nervous system infections. Am J Med Sci 346:158–159. <https://doi.org/10.1097/MAJ.0b013e318294f998>.
- Kim BN, Kim NJ, Kim MN, Kim YS, Woo JH, Ryu J. 2003. Bacteraemia due to tribe *Proteeae*: a review of 132 cases during a decade (1991–2000). Scand J Infect Dis 35:98–103. <https://doi.org/10.1080/0036554021000027015>.
- Stock I, Wiedemann B. 1998. Natural antibiotic susceptibility of *Providencia stuartii*, *P. rettgeri*, *P. alcalifaciens* and *P. rustigianii* strains. J Med Microbiol 47:629–642. <https://doi.org/10.1099/00222615-47-7-629>.
- Saavedra-Rojas SY, Duarte-Valderrama C, González-de-Arias MN, Ovalle-Guerro MV. 2015. Emergence of *Providencia rettgeri* NDM-1 in two departments of Colombia, 2012–2013 [Epub ahead of print]. Enferm Infecc Microbiol Clin pii:S0213-005X(15)00198-6. <https://doi.org/10.1016/j.eimc.2015.05.011>. (In Spanish.)
- Gefen-Halevi S, Hindiyeh MY, Ben-David D, Smollan G, Gal-Mor O, Azar R, Castanheira M, Belasov N, Rahav G, Tal I, Mendelson E, Keller N. 2013. Isolation of genetically unrelated *bla*_{NDM-1}-positive *Providencia rettgeri* strains in Israel. J Clin Microbiol 51:1642–1643. <https://doi.org/10.1128/JCM.00381-13>.
- Tada T, Miyoshi-Akiyama T, Dahal RK, Sah MK, Ohara H, Shimada K, Kirikae T, Pokhrel BM. 2014. NDM-1 metallo-beta-lactamase and ArmA 16S rRNA methylase producing *Providencia rettgeri* clinical isolates in Nepal. BMC Infect Dis 14:56. <https://doi.org/10.1186/1471-2334-14-56>.
- Chin CS, Alexander DH, Marks P, Klammer AA, Drake J, Heiner C, Clum A, Copeland A, Huddleston J, Eichler EE, Turner SW, Korlach J. 2013. Non-hybrid, finished microbial genome assemblies from long-read SMRT sequencing data. Nat Methods 10:563–569. <https://doi.org/10.1038/nmeth.2474>.
- Li H. 2013. Aligning sequence reads, clone sequences and assembly contigs with BWA-MEM. arXiv arXiv:1303.3997. <https://arxiv.org/abs/1303.3997>.
- Milne I, Stephen G, Bayer M, Cock PJ, Pritchard L, Cardle L, Shaw PD, Marshall D. 2013. Using Tablet for visual exploration of second-generation sequencing data. Brief Bioinform 14:193–202. <https://doi.org/10.1093/bib/bbs012>.
- Hunt M, Silva ND, Otto TD, Parkhill J, Keane JA, Harris SR. 2015. Circlator: automated circularization of genome assemblies using long sequencing reads. Genome Biol 16:294. <https://doi.org/10.1186/s13059-015-0849-0>.
- Carver TJ, Rutherford KM, Berriman M, Rajandream MA, Barrell BG, Parkhill J. 2005. ACT: the Artemis Comparison Tool. Bioinformatics 21:3422–3423. <https://doi.org/10.1093/bioinformatics/bti553>.
- Seemann T. 2014. Prokka: rapid prokaryotic genome annotation. Bioinformatics 30:2068–2069. <https://doi.org/10.1093/bioinformatics/btu153>.