

Draft Genome Assembly of *Pseudomonas aeruginosa* Quality Control Reference Strain Boston 41501

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We present the scaffolded genome assembly of *Pseudomonas aeruginosa* Boston 41501, now publicly available in GenBank (JOVK00000000) in 10 contigs placed into a single scaffold. The 6.82-Mbp genome contains 66.1% G+C content and 6,295 coding sequences, including type 4 pilus and type 3 secretion system production genes.

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Pseudomonas aeruginosa is a Gram-negative member of the *Gammaproteobacteria* found in a wide variety of environments (soil water, animals, humans, and hospitals). *P. aeruginosa* Boston 41501 (ATCC 27853) is a quality control strain often used in opportunistic pathogen research and originally isolated from a blood culture. Most human infections are hospital acquired and/or limited to immunocompromised persons. Recently the U.S. Centers for Disease Control and Prevention elevated the risk of antibiotic-resistant *P. aeruginosa* infection to “serious” due to estimates of 6,700 cases and 400 deaths annually (1).

High-quality genomic DNA was extracted from a stationary-phase purified isolate using a Qiagen Genomic-tip 500 at the USAMRIID Diagnostic Systems Division (DSD) per the manufacturer’s recommendations. Draft sequence data included a 100-bp Illumina library (128-fold genome coverage) and a separate long-insert paired-end library (8,792- ± 2,198-bp insert, 35-fold genome coverage) (Roche 454 Titanium platform). The two data sets were assembled together in Newbler (Roche) and the consensus sequences computationally shredded into 2-kbp overlapping fake reads (shreds). The raw reads were also assembled in Velvet and those consensus sequences computationally shredded into 1.5-kbp overlapping shreds (2). Draft data from all platforms were then assembled together with Allpaths and the consensus sequences computationally shredded into 10-kbp overlapping shreds (3). We then integrated the Newbler consensus shreds, Velvet consensus shreds, Allpaths consensus shreds, and a subset of the long-insert read pairs using parallel Phrap (High Performance Software, LLC). Possible misassemblies were corrected, and some gap closure was accomplished with manual editing in Consed (4–6).

Automatic annotation of the *P. aeruginosa* Boston 41501 genome utilized an Ergatis-based workflow at LANL with minor manual curation. The final annotated assembly (6,295 coding sequences, 31 rRNAs, and 66 tRNAs in the 6,819,384-bp genome sequence) is available in GenBank (accession number JOVK00000000), and raw data can be provided upon request.

Preliminary review of the annotations finds at least 28% of the virulence genes noted by Feinbaum et al. (7) and both type 4 pilus and type 3 secretion systems (8).

Nucleotide sequence accession number. This genome sequence is available in GenBank under the accession number JOVK00000000.

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REFERENCES

- Centers for Disease Control and Prevention (CDC). 2013. Antibiotic resistance threats in the United States. U.S. Centers for Disease Control and Prevention, Atlanta, GA. <http://www.cdc.gov/drugresistance/threat-report-2013/pdf/ar-threats-2013-508.pdf>.
- Zerbino DR, Birney E. 2008. Velvet: algorithms for *de novo* short read assembly using de Bruijn graphs. *Genome Res*. 18:821–829. <http://dx.doi.org/10.1101/gr.074492.107>.
- Butler J, MacCallum I, Kleber M, Shlyakhter IA, Belmonte MK, Lander ES, Nusbaum C, Jaffe DB. 2008. ALLPATHS: *de novo* assembly of whole-genome shotgun microreads. *Genome Res*. 18:810–820. <http://dx.doi.org/10.1101/gr.7337908>.
- Ewing B, Hillier L, Wendl MC, Green P. 1998. Base-calling of automated sequencer traces using phred. I. Accuracy assessment. *Genome Res*. 8:175–185. <http://dx.doi.org/10.1101/gr.8.3.175>.
- Ewing B, Green P. 1998. Base-calling of automated sequencer traces using phred. II. Error probabilities. *Genome Res*. 8:186–194.
- Gordon D, Abajian C, Green P. 1998. Consed: a graphical tool for sequence finishing. *Genome Res*. 8:195–202. <http://dx.doi.org/10.1101/gr.8.3.195>.
- Feinbaum RL, Urbach JM, Liberati NT, Djonovic S, Adonizio A, Carvunis AR, Ausubel FM. 2012. Genome-wide identification of *Pseudomonas aeruginosa* virulence-related genes using a *Caenorhabditis elegans* infection model. *PLOS Pathog*. 8:e1002813. <http://dx.doi.org/10.1371/journal.ppat.1002813>.
- Gellatly SL, Hancock RE. 2013. *Pseudomonas aeruginosa*: new insights into pathogenesis and host defenses. *Pathog. Dis*. 67:159–173. <http://dx.doi.org/10.1111/2049-632X.12033>.