

Draft Genome Sequence of the Antibiotic-Producing Cystic Fibrosis Isolate *Pantoea agglomerans* Tx10

Derek D. N. Smith, Morgan W. B. Kirzinger, John Stavrinides

Department of Biology, University of Regina, Regina, Saskatchewan, Canada

Pantoea agglomerans is an enteric bacterium that is capable of causing both plant and human disease. Here, we report the genome sequence of a cystic fibrosis isolate, *P. agglomerans* Tx10, which produces an antibiotic that is effective against *Staphylococcus aureus*.

Received 26 September 2013 Accepted 2 October 2013 Published 31 October 2013

Citation Smith DDN, Kirzinger MWB, Stavrinides J. 2013. Draft genome sequence of the antibiotic-producing cystic fibrosis isolate Pantoea agglomerans Tx10. Genome Announc. 1(5):e00904-13. doi:10.1128/genomeA.00904-13.

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Address correspondence to John Stavrinides, john.stavrinides@uregina.ca.

Pantoea agglomerans (formerly Enterobacter agglomerans and Erwinia herbicola) (1) is a member of the Enterobacteriaceae that has been noted to cause plant disease, as well as opportunistic infections in humans (2–4). *P. agglomerans* has also been shown to have unique metabolic capabilities, including antibiotic biosynthesis (4–10). Among these antibiotics are pantocins (11–14), herbicolins (15, 16), microcins (17–19), and phenazines (20), several of which target amino acid biosynthesis in the fire blight pathogen, *Erwinia amylovora*, and have been developed into biocontrol agents. Here, we report the complete genome sequence of the clinical isolate *P. agglomerans* Tx10, which was isolated from the sputum of a cystic fibrosis patient. This isolate produces multiple antibiotics that target *E. amylovora* and clinically relevant pathogens, including *Staphylococcus aureus*, *Streptococcus epidermidis*, and *Escherichia coli*.

Total DNA was sequenced using Illumina HiSeq 2000, 100-bp paired-end sequencing, resulting in 17,035,538 reads, with an average Phred quality score of 31. ABySS version 1.3.5 (21) was used for *de novo* paired-end assembly using the default parameters and an optimized *k*-mer value of 83. This resulted in 38 contigs with an N₅₀ of 586,961 bp and an estimated genome size of 4,856,603 bp at 347× coverage. Contigs of ≥200 bp (25 total) were submitted to the NCBI Prokaryotic Genome Automatic Annotation Pipeline version 2.0, resulting in 4,627 predicted genes. Of these, there are 4,500 predicted coding genes, 32 pseudogenes, 24 rRNAs, and 71 tRNAs. Two contigs are predicted to represent plasmids of 173,724 bp and 681,148 bp.

The *P. agglomerans* Tx10 genome provides the means not only for identifying antibiotic biosynthetic clusters but also for evaluating the contribution of natural products to polymicrobial infections in cystic fibrosis.

Nucleotide sequence accession numbers. This whole-genome shotgun project has been deposited at GenBank under the accession no. ASJI000000000. The version described in this paper is the first version, ASJI01000000.

ACKNOWLEDGMENTS

Genome sequencing was carried out by Genome Quebec.

This work was supported by grants from the Natural Sciences and

Engineering Research Council of Canada, the Rx&D Health Research Foundation, and the Canada Foundation for Innovation. D.D.N.S. is supported by a Canadian Institutes of Health Research Master's Award.

REFERENCES

- Gavini F, Mergaert J, Beji A, Mielcarek C, Izard D, Kersters K, Deley J. 1989. Transfer of *Enterobacter agglomerans* (Beijerinck 1888) Ewing and Fife 1972 to *Pantoea* gen. nov. as *Pantoea agglomerans* comb. nov. and description of *Pantoea dispersa* sp. nov. Int. J. Syst. Bacteriol. 39:337–345.
- Aly NY, Salmeen HN, Lila RA, Nagaraja PA. 2008. Pantoea agglomerans bloodstream infection in preterm neonates. Med. Princ. Pract. 17: 500-503.
- Astagneau P, Gottot S, Gobin Y, Bocquet P, Gatignol C, Jouvet P, Brücker G. 1994. Nosocomial outbreak of *Enterobacter agglomerans* pseudobacteraemia associated with non-sterile blood collection tubes. J. Hosp. Infect. 27:73–75.
- Cruz AT, Cazacu AC, Allen CH. 2007. Pantoea agglomerans, a plant pathogen causing human disease. J. Clin. Microbiol. 45:1989–1992.
- Yaman M, Aslan I, Calmasur O, Sahin F. 2005. Two bacterial pathogens of *Helicoverpa armigera* (Hubner) (*Lepidoptera*: *Noctuidae*). Proc. Entomol. Soc. Wash. 107:623–626.
- Khleifat KM. 2006. Biodegradation of linear alkylbenzene sulfonate by a two-member facultative anaerobic bacterial consortium. Enzyme Microb. Technol. 39:1030–1035.
- Sergeeva E, Hirkala DLM, Nelson LM. 2007. Production of indole-3acetic acid, aromatic amino acid aminotransferase activities and plant growth promotion by *Pantoea agglomerans* rhizosphere isolates. Plant Soil 297:1–13.
- 8. Sundin GW, Werner NA, Yoder KS, Aldwinckle HS. 2009. Field evaluation of biological control of fire blight in the eastern United States. Plant Dis. 93:386–394.
- Delétoile A, Decré D, Courant S, Passet V, Audo J, Grimont P, Arlet G, Brisse S. 2009. Phylogeny and identification of *Pantoea* species and typing of *Pantoea agglomerans* strains by multilocus gene sequencing. J. Clin. Microbiol. 47:300–310.
- Brady C, Cleenwerck I, Venter S, Vancanneyt M, Swings J, Coutinho T. 2008. Phylogeny and identification of *Pantoea* species associated with plants, humans and the natural environment based on multilocus sequence analysis (MLSA). Syst. Appl. Microbiol. 31:447–460.
- Wright SA, Beer SV. 1996. The role of antibiotics in biological control of fire blight by *Erwinia herbicola* strain EH318. Acta Hortic. 411:309–311.
- 12. Wright SAI, Beer SV. 2002. Genes for biosynthesis of pantocin A and B by *Pantoea agglomerans* Eh318. Acta Hortic. **590**:237–241.
- 13. Wright SAI, Jin M, Clardy J, Beer SV. 2006. The biosynthetic genes of

pantocin A and pantocin B of *Pantoea agglomerans* Eh318. Acta Hortic. **704:**313–320.

- 14. Wright SA, Zumoff CH, Schneider L, Beer SV. 2001. Pantoea agglomerans strain EH318 produces two antibiotics that inhibit *Erwinia amylo*vora in vitro. Appl. Environ. Microbiol. 67:284–292.
- Smits TH, Rezzonico F, Kamber T, Blom J, Goesmann A, Ishimaru CA, Frey JE, Stockwell VO, Duffy B. 2011. Metabolic versatility and antibacterial metabolite biosynthesis are distinguishing genomic features of the fire blight antagonist *Pantoea vagans* C9-1. PLoS One 6:e22247. doi:10.13 71/journal.pone.0022247.
- Smits TH, Rezzonico F, Kamber T, Goesmann A, Ishimaru CA, Stockwell VO, Frey JE, Duffy B. 2010. Genome sequence of the biocontrol agent *Pantoea vagans* strain C9-1. J. Bacteriol. 192:6486–6487.
- 17. Vanneste JL, Cornish DA, Yu J, Voyle MD. 2002. The peptide antibiotic

produced by *Pantoea agglomerans* Eh252 is a microcin. Acta Hortic. **590**: 285–290.

- Vanneste JL, Cornish DA, Yu J, Voyle MD. 2000. A microcin produced by a strain of *Erwinia herbicola* is involved in biological control of fire blight and soft rot caused by *Erwinia* sp. Acta Hortic. 513:39–46.
- Vanneste JL, Yu J, Cornish DA. 2008. Presence of genes homologous to those necessary for synthesis of microcin MccEh252 in strains of *Pantoea* agglomerans. Acta Hortic. 793:391–396.
- Giddens SR, Feng Y, Mahanty HK. 2002. Characterization of a novel phenazine antibiotic gene cluster in *Erwinia herbicola* Eh1087. Mol. Microbiol. 45:769–783.
- Simpson JT, Wong K, Jackman SD, Schein JE, Jones SJ, Birol I. 2009. ABySS: A parallel assembler for short read sequence data. Genome Res. 19:1117–1123.