

Draft Genome Sequence of *Ralstonia solanacearum* Race 4 Biovar 4 Strain SD54

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Ralstonia solanacearum is an important etiological agent that can cause serious bacterial wilt in a very wide range of potential host plants, including ginger. Here, we report the complete genome sequence of *R. solanacearum* SD54, a race 4 biovar 4 (R4B4) strain from a diseased ginger plant in China.

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R*alstonia solanacearum* is an aerobic, non-spore-forming, Gram-negative plant pathogenic bacterium and is soil borne and motile with a polar flagellar tuft. It colonizes the xylem, causing bacterial wilt in a very wide range of potential host plants, including ginger. *R. solanacearum* is considered to be a species complex, a heterogeneous group of related but genetically distinct strains (1, 2). Traditionally, the group has been subdivided into 5 races and 4 biovars based on the differences in host range (3, 4) and the acidification of medium during the metabolism of six carbohydrates (maltose, lactose, cellobiose, mannitol, sorbitol, and dulcitol) (5, 6), respectively. An entirely new phylogenetic classification system was proposed recently, consisting of four phylotypes, each further divided into sequevars (1, 7).

One of the major strains that has caused serious bacterial wilt damage in ginger plants in China is *R. solanacearum* race 4 biovar 4 (R4B4) SD54, the draft genome sequence of which is reported here.

The nucleotide sequence was determined using a 454 GS FLX sequencer, and assembly was performed using the GS *de novo* assembler software version 2.3 (8). A total of 280,325 reads, including up to 100,016,604 bp, were obtained, which represents a 17.6-fold coverage of the genome. In addition, a 100-fold coverage of the 3-kb mate-pair library was constructed and sequenced using an Illumina HiSeq 2000. After scaffold construction and gap filling, we finally obtained the *R. solanacearum* SD54 draft genome sequence of 5,648,859 bp distributed in 175 contigs, with a G+C content of 66.9%.

Putative protein coding sequences (CDSs) were predicted using Glimmer (9) and GeneMark (10). Functional annotation was based on BLASTp results with the NR databases and the Kyoto Encyclopedia of Genes and Genomes (KEGG). tRNA genes were directly predicted with tRNAscan-SE (11). The draft genome sequence consists of one rRNA operon, 46 tRNA genes, and 5,112 CDSs with an average length of 942 bp. Among the CDS products, 3,720 proteins were assigned to Clusters of Orthologous Groups (COG) families (12) and 3,952 proteins were assigned biological functions. A total of 2,235 proteins of *R. solanacearum* SD54 had KEGG orthologs, and these proteins were involved in 177 pathways. In addition, 284 proteins have no known match to any proteins in the databases. In a comparison with the known proteins of the *Ralstonia* genus, 4,603 proteins of *R. solanacearum* SD54 have orthologs.

Nucleotide sequence accession numbers. The genome sequence of *R. solanacearum* strain SD54 has been deposited in the GenBank database under the accession no. ASQR00000000. The version described here is accession no. ASQR02000000.

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REFERENCES

- Fegan M, Prior P. 2005. How complex is the *Ralstonia solanacearum* species complex? p 449–461. *In* Allen C, Prior P, Hayward AC (ed), Bacterial wilt: the disease and the *Ralstonia solanacearum* species complex. American Phytopathological Society, St. Paul, MN.
- 2. Fegan M, Taghavi M, Sly LI, Hayward AC. 1998. Phylogeny, diversity, and molecular diagnostics of *Ralstonia solanacearum*, p 19–33. *In* Prior P, Allen C, Elphinstone J (ed), Bacterial wilt disease: molecular and ecological aspects. Springer-Verlag, Berlin, Germany.
- Buddenhagen I, Kelman A. 1964. Biological and physiological aspects of bacterial wilt caused by *Pseudomonas olanacearum*. Annu. Rev. Phytopathol. 2:203–230. doi:10.1146/annurev.py.02.090164.001223.
- 4. Buddenhagen I, Sequeira L, Kelman A. 1962. Designation of races in *Pseudomonas solanacearum*. Phytopathology 52:726.
- Hayward AC. 1964. Characteristics of *Pseudomonas solanacearum*. J. Appl. Bacteriol. 27:265–277. doi:10.1111/j.1365-2672.1964.tb04912.x.

- Hayward AC. 1994. Systematics and phylogeny of *Pseudomonas so-lanacearum* and related bacteria, p 123–135. *In* Hayward AC, Hartman GL (ed), Bacterial wilt: the disease and its causative agent, *Pseudomonas so-lanacearum*. CAB International, Wallingford, United Kingdom.
- Villa JE, Tsuchiya K, Horita M, Opina N, Hyakumachi M. 2005. Phylogenetic relationships of *Ralstonia solanacearum* species complex strains from Asia and other continents based on 16S rDNA, endoglucanase, and *hrpB* gene sequences. J. Gen. Plant Pathol. 71:39–46. doi:10.10 07/s10327-004-0156-1.
- 8. Margulies M, Egholm M, Altman WE, Attiya S, Bader JS, Bemben LA, Berka J, Braverman MS, Chen YJ, Chen Z, Dewell SB, Du L, Fierro JM, Gomes XV, Godwin BC, He W, Helgesen S, Ho CH, Irzyk GP, Jando SC, Alenquer MLI, Jarvie TP, Jirage KB, Kim JB, Knight JR, Lanza JR, Leamon JH, Lefkowitz SM, Lei M, Li J, Lohman KL, Lu H, Makhijani VB, McDade KE, McKenna MP, Myers EW, Nickerson E, Nobile JR, Plant R, Puc BP, Ronan MT, Roth GT, Sarkis GJ, Simons JF, Simpson

JW, Srinivasan M, Tartaro KR, Tomasz A, Vogt KA, Volkmer GA, Wang SH, Wang Y, Weiner MP, Yu P, Begley RF, Rothberg JM. 2005. Genome sequencing in microfabricated high-density picolitre reactors. Nature 437:376–380. doi:10.1038/nature03959.

- Delcher AL, Harmon D, Kasif S, White O, Salzberg SL. 1999. Improved microbial gene identification with Glimmer. Nucleic Acids Res. 27: 4636–4641. doi:10.1093/nar/27.23.4636.
- Besemer J, Borodovsky M. 2005. GeneMark: web software for gene finding in prokaryotes, eukaryotes and viruses. Nucleic Acids Res. 33: W451–W454. doi:10.1093/nar/gki487.
- Lowe TM, Eddy SR. 1997. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. Nucleic Acids Res. 25: 955–964.
- 12. Tatusov RL, Galperin MY, Natale DA, Koonin EV. 2000. The COG database: a tool for genome-scale analysis of protein functions and evolution. Nucleic Acids Res. 28:33–36. doi:10.1093/nar/28.1.33.