Complete Genome Sequence of the *Streptococcus* sp. Strain VT 162, Isolated from the Saliva of Pediatric Oncohematology Patients

Maria F. Vecherkovskaya, George V. Tetz, Victor V. Tetz

Institute of Human Microbiology LLC, Department of Microbiology, Virology and Immunology, First State I. P. Pavlov Medical University, Saint Petersburg, Russia

Streptococcus sp. strain VT 162 was isolated from the saliva of pediatric oncohematology patients. Its full genome is 2,045,418 bp. The availability of this genome will provide insights into the composition of microbial flora among pediatric oncohematology patients and the host interaction and pathogenicity of this species.

Received 17 June 2014 Accepted 23 June 2014 Published 10 July 2014

Citation Vecherkovskaya MF, Tetz GV, Tetz W. 2014. Complete genome sequence of the *Streptococcus* sp. strain VT 162, isolated from the saliva of pediatric oncohematology patients. Genome Announc. 2(4):e00647-14. doi:10.1128/genomeA.00647-14.

Copyright © 2014 Vecherkovskaya et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 3.0 Unported license.

Address correspondence to Maria F. Vecherkovskaya, mashavecher@yahoo.com.

S*treptococcus* species are Gram-positive facultatively anaerobic cocci. Streptococci are widely present in the oral cavity and are established part of commensal flora (1). In pediatric cancer patients, and in oncohematology patients in particular, streptococci are known to cause a wide variety of conditions ranging from local (caries, mucositis, stomatitis, and gingivitis) (2, 3) to generalized (septicemia and sepsis) complications (4, 5).

Streptococcus sp. strain VT162 was isolated from the saliva of several patients with oncohematological malignancies in remission during their follow-up visits. Biochemical and matrix-assisted laser desorption ionization-time of flight mass spectrometry (MALDI-TOF MS) identifications gave divergent low-discrimination results. Phylogenetic analysis of the 16S rRNA gene sequence had placed this isolate among strepto-cocci, but identification to the species level was still inconclusive. Thus, whole-genome sequencing was performed.

The *Streptococcus* sp. strain VT162 was sequenced using the Illumina HiSeq 2500 system sequencing technology (Illumina GA IIx; Illumina, CA). The library preparation, sequencing reaction, and the sequencing run were carried out according to Illumina's instructions. A total of 16.79 million high-quality 51-bp single-end reads were produced, resulting in approximate coverage of $345 \times$. Assembly was performed using Velvet assembler version 1.2.10 (6) and resulted in 340 contigs. The contig N_{50} is 42,108 bp, and the largest assembled contig is 115,694 bp. To produce the chromosome assembly, the contigs were oriented and ordered by alignment with the genome of *Streptococcus oralis* Uo5 (7) using the Mauve aligner version 2.3.1 (8).

Annotation was added by the NCBI Prokaryotic Genome Annotation Pipeline (2013 release). The VT 162 chromosome is 2,045,418 bp in length, with a G+C content of 41.1%. There are 1,935 predicted protein-coding sequences (CDSs), with an average length of 909 bp, and 34 tRNAs, 3 rRNAs, and 1 noncoding RNA (ncRNA) gene were identified.

The genome contains genes for several competence proteins and multidrug resistance transporters of the ABC, MATE, MFS, and DMT families, the *vanZ* gene that confers resistance to teicoplanin, and genes for hemolysin III and capsular and capsid proteins.

The average nucleotide identity (ANI) (9) demonstrates that *Streptococcus* sp. VT 162 differs significantly from the mitis phylogenetic group of streptococcus, showing ANI values of <95% compared to the genomes of the closest species, such as *S. oralis* Uo5 (7).

The complete genome sequence of *Streptococcus* sp. VT 162 will help in establishing a new species of *Streptococcus* and will provide insights into the composition of microbial flora among pediatric oncohematology patients, as well as the host interaction and pathogenicity of this species.

Nucleotide sequence accession number. The complete genome sequence has been deposited in the NCBI database under accession no. CP007628.

ACKNOWLEDGMENT

No financial support was received for this work.

REFERENCES

- Norder Grusell E, Dahlén G, Ruth M, Ny L, Quiding-Järbrink M, Bergquist H, Bove M. 2013. Bacterial flora of the human oral cavity, and the upper and lower esophagus. Dis. Esophagus 26:84–90. http:// dx.doi.org/10.1111/j.1442-2050.2012.01328.x.
- Stringer AM, Logan RM. 4 February 2014. The role of oral flora in the development of chemotherapy-induced oral mucositis. J. Oral Pathol. Med. http://dx.doi.org/10.1111/jop.12152.
- Chaveli-López B. 2014. Oral toxicity produced by chemotherapy: a systematic review. J. Clin. Exp. Dent. 6:e81–e90. http://dx.doi.org/10.4317/jced.51337.
- Freifeld AG, Razonable RR. 21 April 2014. Editorial commentary: viridans group streptococci in febrile neutropenic cancer patients: what should we fear? Clin. Infect. Dis. 59:231–233. http://dx.doi.org/10.1093/cid/ciu264.
- Han SB, Bae EY, Lee JW, Lee DG, Chung NG, Jeong DC, Cho B, Kang JH, Kim HK. 2013. Clinical characteristics and antimicrobial susceptibilities of viridans streptococcal bacteremia during febrile neutropenia in patients with hematologic malignancies: a comparison between adults and children. BMC Infect. Dis. 13:273. http://dx.doi.org/10.1186/1471-2334-13 -273.
- 6. Zerbino DR. 2010. Using the Velvet de novo assembler for short-read

sequencing technologies. Curr. Protoc. Bioinformatics Chapter 11:Unit 11.5. http://dx.doi.org/10.1002/0471250953.bi1105s31.
7. Reichmann P, Nuhn M, Denapaite D, Brückner R, Henrich B, Maurer P, Rieger

- Reichmann P, Nuhn M, Denapaite D, Brückner R, Henrich B, Maurer P, Rieger M, Klages S, Reinhard R, Hakenbeck R. 2011. Genome of *Streptococcus oralis* strain Uo5. J. Bacteriol. 193:2888–2889. http://dx.doi.org/10.1128/JB.00321-11.
- 8. Darling AE, Mau B, Perna NT. 2010. progressiveMauve: multiple genome

alignment with gene gain, loss and rearrangement. PLoS One 5:e11147. http://dx.doi.org/10.1371/journal.pone.0011147.
9. Kim M, Oh HS, Park SC, Chun J. 2014. Towards a taxonomic coherence

 Kim M, Oh HS, Park SC, Chun J. 2014. Towards a taxonomic coherence between average nucleotide identity and 16S rRNA gene sequence similarity for species demarcation of prokaryotes. Int. J. Syst. Evol. Microbiol. 64:346–351. http://dx.doi.org/10.1099/ijs.0.059774-0.