Draft genome sequence of *Streptococcus constellatus* KCOM 1039 isolated from human postoperative maxillary cyst lesion

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사람 수술후 상악낭종 병소에서 분리된 Streptococcus constellatus KCOM 1039의 유전체 염기서열 완전 해독

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Streptococcus constellatus is Gram-stain-positive, facultative anaerobic, and non-spore forming coccus. It is a member of normal flora of human oral cavity. *S. constellatus* KCOM 1039 was isolated from human postoperative maxillary cyst lesion. In this report, we present the draft genome sequence of *S. constellatus* KCOM 1039.

Keywords: *Streptococcus constellatus*, human, postoperative maxillary cyst

Streptococcus constellatus is Gram-stain-positive, facultative anaerobic, and non-spore forming coccus which are round to slightly oval and occur in pairs and short chain (Whiley et al., 1999). It showed beta-hemolytic activity and produced a streptolysin S-like protein (Tabata et al., 2014). The bacterium is primarily inhabits the human oral cavity and is considered to be an opportunistic human pathogen (Tabata et al., 2014). S. constellatus KCOM 1039 (= ChDC B280) was isolated from

S. constellatus KCOM 1039 was grown in brain heart infusion (BHI, Difco Laboratories) medium in an anaerobic chamber (Model Bactron I) maintained using a gas mixture of 10% H₂, 5% CO₂, and 85% N₂. Genomic DNA of S. constellatus KCOM 1039 was prepared as previously described (Cho et al., 2015).

Genomic DNA of *S. constellatus* KCOM 1039 was sequenced using the Illumina Hiseq 2000 platform by Macrogen Inc. Three libraries of 350 bp paired-end, 5 kb mate-pair, and 8 kb mate-pair were sequenced which reached coverage of 657.0 ×, 646.2 ×, and 955.7 ×, respectively. The *de novo* assembly was performed by SPAdes (http://bioinf.spbau.ru/spades) (Bankevich *et al.*, 2012). All gaps among the scaffolds were filled by GapCloser (http://soap.genomics.org.cn/soapdenovo.html) (Luo *et al.*, 2012). Error correction was performed by Pilon (https://github.com/broadinstitute/pilon/wiki) (Walker *et al.*, 2014). Genome annotation was conducted by the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) (https://www.ncbi.nlm.nih.gov/genome/annotation_prok/) (Tatusova *et al.*, 2016).

The draft genome of S. constellatus KCOM 1039 was

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human periodontitis lesion. In this report, we presented the draft genome sequence of *S. constellatus* KCOM 1039.

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Table 1. Genome features of Streptococcus constellatus KCOM 1039

Attribute	Value
Genome size (bp)	1,885,802
GC content (%)	38.0
No. of contigs	2
Total genes	1,898
Protein-coding genes	1,730
tRNA	58
Complete rRNA (5S, 16S, 23S)	9 (3,3,3)
ncRNA	3
Pseudogene	97

composed of 2 contigs; 1,883,587 bp and 2,215 bp in length. The average G+C content of the genome was 38.0%. A total of 1,730 protein-coding sequences, 9 complete rRNA operons (3 5S rRNAs, 3 16S rRNAs, and 3 23 rRNAs), 58 tRNAs, and 3 ncRNAs were identified (Table 1).

In the genome, genes involved in proteinase were identified; putative protease YdcP/YdeA, protease HtpX, ATP-dependent zinc metalloprotease FtsH, putative zinc metalloprotease, Lon protease, rhomboid protease GluP, serine protease Do-like HtrA, and putative zinc protease AlbF. It also contained antibioticresistance-related genes; multiple antibiotic resistance protein MarA/MarR, daunorubicin/doxorubicin resistance ATP-binding protein DrrA, putative multidrug resistance ABC transporter ATP-binding/permease protein Yhel/YheH, multidrug resistance protein NorM, and tetracycline resistance protein TetM from transposon TnFO1. It also contained biofilm formation-related genes; toxin-antitoxin biofilm protein TabA, putative glycosyltransferase EpsH/EpsJ/CsbB, biofilm regulatory protein A, and D-inositol-3-phosphate glycosyltransferase. It also contained type II secretion system protein F, protein translocase subunit SecA/SecY, and ESAT-6 secretion accessory factor EsaA. The genome also contained the oxidative stress-response genes; glutathione reductase and thioredoxin reductase. Hemolysin A and diphtheria toxin repressor were also found in the genome sequence.

The *S. constellatus* KCOM 1039 strain was deposited into the Korean Collection for Oral Microbiology.

Nucleotide sequence accession number

This whole genome shotgun project has been deposited at DDBJ/ENA/GenBank under the accession QWKV00000000. The version described in this paper is version QWKV00000000.

적 요

Streptococcus constellatus는 그람 염색에 양성을 띄며, 조건 혐기성이면서 아포를 생성하지 않는 구균이다. S. constellatus 는 사람 구강의 정상 세균 총에 속한다. 수술후 상악낭종 병소에서 S. constellatus KCOM 1039 균주가 분리되었다. 여기에서 S. constellatus KCOM 1039 균주의 유전체 염기서열을 결정하여 보고한다.

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References

- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski AD, et al. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. J. Comput. Biol. 19, 455–477.
- Cho E, Park SN, Lim YK, Shin Y, Paek J, Hwang CH, Chang YH, and Kook JK. 2015. *Fusobacterium hwasookii* sp. nov., isolated from a human periodontitis lesion. *Curr. Microbiol.* **70**, 169–175.
- Luo R, Liu B, Xie Y, Li Z, Huang W, Yuan J, He G, Chen Y, Pan Q, Liu Y, et al. 2012. SOAPdenovo2: an empirically improved memory-efficient short-read de novo assembler. Gigascience 1, 18. Erratum in: Gigascience 2015. 4, 30.
- Tabata A, Sato Y, Maya K, Nakano K, Kikuchi K, Whiley RA, Ohkura K, Tomoyasu T, and Nagamune H. 2014. A streptolysin S homologue is essential for β-haemolytic *Streptococcus constellatus* subsp. constellatus cytotoxicity. *Microbiology* **160**, 980–991.
- Tatusova T, DiCuccio M, Badretdin A, Chetvernin V, Nawrocki EP, Zaslavsky L, Lomsadze A, Pruitt KD, Borodovsky M, and Ostell J. 2016. NCBI prokaryotic genome annotation pipeline. *Nucleic Acids Res.* 44, 6614–6624.
- Walker BJ, Abeel T, Shea T, Priest M, Abouelliel A, Sakthikumar S, Cuomo CA, Zeng Q, Wortman J, Young SK, et al. 2014. Pilon: an integrated tool for comprehensive microbial variant detection and genome assembly improvement. PLoS One 9, e112963.
- Whiley RA, Hall LM, Hardie JM, and Beighton D. 1999. A study of small-colony, beta-haemolytic, Lancefield group C streptococci within the anginosus group: description of *Streptococcus constellatus* subsp. *pharyngis* subsp. nov., associated with the human throat and pharyngitis. *Int. J. Syst. Bacteriol.* 49, 1443–1449.