



Complete Genome Sequences of Three Fish-Associated *Streptococcus agalactiae* Isolates

Anita Jaglarz,^a Artur Gurgul,^b William J. Leigh,^a Janina Z. Costa,^a Kim D. Thompson^a

^aAquaculture Research Group, Moredun Research Institute, Pentlands Science Park, Penicuik, United Kingdom

^bDepartment of Genomics and Molecular Biology of Animals, The National Research Institute of Animal Production, Balice, Poland

ABSTRACT The whole-genome sequences are described here for three group B *Streptococcus* (GBS) (*S. agalactiae*) serotype Ib isolates obtained from tilapia (*Oreochromis niloticus*) farmed at sites in Honduras, Costa Rica, and the United States. The bacteria were isolated from the brains of fish displaying signs of streptococcosis.

Streptococcus agalactiae is a major cause of streptococcosis in farmed tilapia and has been associated with significant levels of mortality in diseased fish (1, 2). We have selected three group B *Streptococcus* (GBS) (*S. agalactiae*) serotype Ib isolates obtained from the brains of diseased tilapia for complete genome sequencing. The isolates we analyzed belong to clonal complex 552 (CC552) and sequence types 260 (isolates 14-98 and 14-104) and 261 (isolate 14-110) and were typed using molecular serotyping and multilocus sequence typing (MLST), as we previously described for serotype Ia (3). Serotype Ib isolates belonging to CC552 have a significantly reduced genome size compared to that of other *S. agalactiae* genomes (4) and are predominantly associated with fish and other poikilothermic species (5).

Sequencing of the isolates was performed using a MiSeq system (Illumina, San Diego, CA) with 250-bp paired-end reads. Libraries with an average 550-bp insert were prepared using a TruSeq Nano DNA library preparation kit (Illumina). *De novo* assembly of the reads was performed with the SPAdes version 3.5.0 software (6) and the quality of the assembly assessed with the QUAST software (7). The contigs (>200 nucleotides [nt]) were annotated using the NCBI Prokaryotic Genome Annotation Pipeline to predict protein-coding genes and other functional genome units, including structural RNAs, tRNAs, small RNAs, pseudogenes, control regions, direct and inverted repeats, insertion sequences, transposons, and other mobile elements.

The genomes of *S. agalactiae* Ib isolates 14-98, 14-104, and 14-110 were 1,842,528, 1,815,466, and 1,803,751 bp distributed in 99, 51, and 30 contigs and with average *de novo* assembly coverages of 193×, 227×, and 154×, respectively. The G+C genome content for each of the three isolates was 35.3%. The genome of isolate 14-98 was composed of a total of 1,980 genes, including 1,761 coding genes, 56 RNA genes (50 tRNA), and 163 pseudogenes, while the genome of isolate 14-104 contained 1,910 genes, of which 1,703 are coding genes, 48 are RNA genes (42 tRNAs), and 159 are pseudogenes. The genome of strain 14-110 was composed of a total of 1,883 genes, with 1,647 coding genes, 59 RNA genes containing 52 tRNAs, and 177 pseudogenes.

Accession number(s). The genome sequences of *S. agalactiae* strains 14-98, 14-104, and 14-110 have been deposited in the GenBank database under the accession numbers [NKQD00000000](#), [NKQC00000000](#), and [NKQB00000000](#), respectively.

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Address correspondence to Anita Jaglarz, anita.jaglarz@moredun.ac.uk.

REFERENCES

1. Hernández E, Figueroa J, Iregui C. 2009. Streptococcosis on a red tilapia, *Oreochromis* sp., farm: a case study. *J Fish Dis* 32:247–252. <https://doi.org/10.1111/j.1365-2761.2008.00981.x>.
2. Suebsing R, Kampeera J, Tookdee B, Withyachumnarnkul B, Turner W, Kiatpathomchai W. 2013. Evaluation of colorimetric loop-mediated isothermal amplification assay for visual detection of *Streptococcus agalactiae* and *Streptococcus iniae* in tilapia. *Lett Appl Microbiol* 57:317–324. <https://doi.org/10.1111/lam.12114>.
3. Jaglarz A, Gurgul A, Leigh WJ, Costa JZ, Thompson KD. 2017. Complete genome sequence of three *Streptococcus agalactiae* serotype Ia isolates obtained from disease outbreaks in Nile tilapia *Oreochromis niloticus*. *Genome Announc* 6:e01432-17. <https://doi.org/10.1128/genomeA.01432-17>
4. Rosinski-Chupin I, Sauvage E, Mairey B, Mangenot S, Ma L, Da Cunha V, Rusniok C, Bouchier C, Barbe V, Glaser P. 2013. Reductive evolution in *Streptococcus agalactiae* and the emergence of a host adapted lineage. *BMC Genomics* 14:252. <https://doi.org/10.1186/1471-2164-14-252>.
5. Delannoy CMJ, Zadoks RN, Crumlish M, Rodgers D, Lainson FA, Ferguson HW, Turnbull J, Fontaine MC. 2016. Genomic comparison of virulent and non-virulent *Streptococcus agalactiae* in fish. *J Fish Dis* 39:13–29. <https://doi.org/10.1111/jfd.12319>.
6. Nurk S, Bankevich A, Antipov D, Gurevich AA, Korobeynikov A, Lapidus A, Prjibelski AD, Pyshkin A, Sirotkin A, Sirotkin Y, Stepanauskas R, Clingenpeel SR, Woyke T, McLean JS, Lasken R, Tesler G, Alekseyev MA, Pevzner PA. 2013. Assembling single-cell genomes and mini-metagenomes from chimeric MDA products. *J Comput Biol* 20:714–737. <https://doi.org/10.1089/cmb.2013.0084>.
7. Gurevich A, Saveliev V, Vyahhi N, Tesler G. 2013. QUASt: quality assessment tool for genome assemblies. *Bioinformatics* 29:1072–1075. <https://doi.org/10.1093/bioinformatics/btt086>.