



# Whole-Genome Sequence of a *Mycobacterium goodii* Isolate from a Pediatric Patient in South Africa

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**ABSTRACT** We describe here the draft genome sequence of a *Mycobacterium goodii* isolate from a pediatric patient in Western Cape, South Africa. To our knowledge, this is the second reported genome of this rapidly growing nontuberculous mycobacterial species.

Nine specimens from a pediatric patient were submitted for culture, including specimens obtained from bronchoalveolar lavage fluid, tissue, gastric washing fluid, and a pus swab for tuberculosis investigation. No acid-fast bacilli were observed on direct auramine staining and microscopy. Cultivation of mycobacteria was successful for the gastric washing fluid, pus swab, and bronchoalveolar lavage fluid specimens within a median time of 4 days (range, 2 to 14 days) using the MGIT 960 instrument (BD, Sparks, MD, USA). The presence of noncorded acid-fast bacilli was observed when we performed Ziehl-Neelsen staining on these cultures. The tuberculosis antigen MPT64 rapid test (SD Biotline) for detection of the presence of *Mycobacterium tuberculosis* complex was negative. Thereafter, species identification was performed using the GenoType Mycobacterium CM version 2.0 (Hain Lifescience GH, Nehren, Germany) on an isolate obtained from a gastric washing fluid specimen which was identified as a *Mycobacterium* species. The isolate, which was designated strain ST0139456, was then subjected to 16S rRNA sequencing using the forward primer 5'-AGTTTGATCMTGGCTCAG-3' and reverse primer 5'-GGACTACHAGGGTATCTAAT-3', and the resulting BLAST search (<https://blast.ncbi.nlm.nih.gov>) confirmed 99% homology to *Mycobacterium goodii*, a nontuberculous mycobacterium species of the *Mycobacterium smegmatis* group. The species identification as *Mycobacterium goodii* was further confirmed using the GenoType Mycobacterium AS version 1.0 kit (Hain Lifescience GH) and whole-genome sequencing.

Paired-end libraries were prepared using the Nextera XT DNA library kit, followed by 2 × 300-bp sequencing on a MiSeq instrument (Illumina, San Diego, CA, USA). The sequenced reads were quality trimmed using Sickle version 1.33 (<https://github.com/najoshi/sickle>) and *de novo* assembled using SPAdes genome assembler version 3.5 (1). The assembly contains 156 contig sequences of longer than 200 bp and covers 6,621,508 bp, with a G+C content of 67.07% and an  $N_{50}$  of 116,855 bp. Genome annotation was performed via the NCBI Prokaryotic Genome Annotation Pipeline

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(PGAP) (2). The total number of 6,514 genes predicted by PGAP includes 6,281 protein-coding genes, 175 pseudogenes, and 58 RNA genes.

**Accession number(s).** The draft genome sequence has been deposited at NCBI under the BioProject number PRJNA415539, BioSample number SAMN07828250, and GenBank accession number [PEBB00000000](https://www.ncbi.nlm.nih.gov/nuccore/PEBB00000000).

## ACKNOWLEDGMENT

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