



# Whole-Genome Sequence of *Mycoplasma bovis* Strain Ningxia-1

Peng Sun,<sup>a,b</sup> Haifeng Luo,<sup>a</sup> Xin Zhang,<sup>a</sup> Jingyi Xu,<sup>a</sup> Yanan Guo,<sup>a</sup> Shenghu He<sup>a</sup>

<sup>a</sup>School of Agriculture, Ningxia University, Yinchuan, China

<sup>b</sup>Virginia-Maryland Regional College of Veterinary Medicine, University of Maryland–College Park, College Park, Maryland, USA

**ABSTRACT** A genome sequence of the *Mycoplasma bovis* Ningxia-1 strain was tested by Pacific Biosciences (PacBio) single-molecule real-time (SMRT) sequencing technology. The strain was isolated from a lesioned calf lung in 2013 in Pengyang, Ningxia, China. The single circular chromosome of 1,033,629 bp shows differences between complete *Mycoplasma bovis* genome in insertion-like sequences (ISs), integrative conjugative elements (ICEs), lipoproteins (LPs), variable surface lipoproteins (VSPs), pathogenicity islands (PAIs), etc.

*Mycoplasma bovis* is the main cause of bovine respiratory disease syndrome. At present, the pathogen is prevalent worldwide, which causes huge economic losses to national cattle industries (1). In 2008, *M. bovis* was isolated from beef cattle in Hubei, China (2). Other places in China, such as Xinjiang, Ningxia, Chongqing, Guizhou, and Qingdao, later reported *M. bovis* isolated from beef cattle and dairy cows (3).

The whole-genome sequence of the *M. bovis* Ningxia-1 strain was isolated in 2013 from the lesioned lung of a beef calf, which was tested using single-molecule real-time (SMRT) technology (4), resulting in approximately 1,262-fold final sequence coverage. The genome of *M. bovis* Ningxia-1 contains a single circular chromosome of 1,033,629 bp, with a GC content of 29.33%. A total of 754 open reading frames (ORFs) were identified, totaling 845,928 bp (maximum length, 9,981 bp; minimum length, 114 bp), and occupied 81.84% of the whole genome, with an average length of 1121 bp and a mean GC content of 29.77%. A total of 577 coding sequence (CDS) genes could be classified into clusters of orthologous groups (COG) families, which have 19 functional categories. Seventy-four pseudogenes were predicted by GeneMarkS+ (5). The genome encodes 6 rRNA and 34 tRNA genes, representing all 20 amino acids.

We found 60 insertion-like sequence (IS) elements that comprised three distinct categories (<https://www-is.biotoul.fr>). A cluster of 8 variable surface protein (VSP)-related ORFs were found in the genome. Sixty-six lipoproteins (LPs) revealed signatures denoting distinct mutation-based mechanisms of phase variation (6). Of three predicted pathogenicity islands (PAIs) (7) with a lower GC content (8), PAI-1 (463,228 to 485,927) encoded one single-stranded DNA-binding protein and one conjugal transfer protein, TraE, and contained a transposase at the 5' end. PAI-2 (541731 to 556809) presented an IS4 and IS30 family transposase at the 3' end and contained an IS1634 family transposase gene. A total of two nucleotidyl transferase AbiEii/AbiGii toxin family proteins and abortive phage infection proteins in PAI-3 (968,698 to 977,477) were without an IS element but still denoted a pathogenicity island. These characteristics may contribute to the emergence of bacterial pathogens with new virulence properties (9).

**Accession number(s).** This whole-genome sequence assembly has been deposited at GenBank under the accession no. [CP023663](https://www.ncbi.nlm.nih.gov/nuclseq/CP023663).

Received 29 November 2017 Accepted 5 December 2017 Published 25 January 2018

**Citation** Sun P, Luo H, Zhang X, Xu J, Guo Y, He S. 2018. Whole-genome sequence of *Mycoplasma bovis* strain Ningxia-1. Genome Announc 6:e01367-17. <https://doi.org/10.1128/genomeA.01367-17>.

**Copyright** © 2018 Sun et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Shenghu He, [heshenghu308@163.com](mailto:heshenghu308@163.com).

## ACKNOWLEDGMENTS

This work was sponsored by the China Science and Technology of Ningxia support program (grant 412-0164).

We thank Wei Yangyang from GeneDenovo, Guangzhou, China, for his assistance with data analysis.

## REFERENCES

1. Tamang MD, Nam HM, Jang GC, Kim SR, Chae MH, Jung SC, Byun JW, Park YH, Lim SK. 2012. Molecular characterization of extended-spectrum-beta-lactamase-producing and plasmid-mediated AmpC beta-lactamase-producing *Escherichia coli* isolated from stray dogs in South Korea. *Antimicrob Agents Chemother* 56:2705–2712. <https://doi.org/10.1128/AAC.05598-11>.
2. Qi J, Guo A, Cui P, Chen Y, Mustafa R, Ba X, Hu C, Bai Z, Chen X, Shi L, Chen H. 2012. Comparative geno-plasticity analysis of *Mycoplasma bovis* HB0801 (Chinese isolate). *PLoS One* 7:e38239. <https://doi.org/10.1371/journal.pone.0038239>.
3. Zhou W, Li N, Wei X, Hu J, Xia T, Wang Z, Peng Y. 2012. The diagnosis of mycoplasma pneumoniae in a beef cattle farm in Chongqing, China. *Prev Vet Med* 34:326–328.
4. Bleidorn C. 2016. Third generation sequencing: technology and its potential impact on evolutionary biodiversity research. *Syst Biodivers* 14: 1–8. <https://doi.org/10.1080/14772000.2015.1099575>.
5. Besemer J, Lomsadze A, Borodovsky M. 2001. GeneMarkS: a self-training method for prediction of gene starts in microbial genomes. Implications for finding sequence motifs in regulatory regions. *Nucleic Acids Res* 29:2607–2618. <https://doi.org/10.1093/nar/29.12.2607>.
6. Wise KS, Calcutt MJ, Foecking MF, Röske K, Madupu R, Methé BA. 2011. Complete genome sequence of *Mycoplasma bovis* type strain PG45 (ATCC 25523). *Infect Immun* 79:982–983. <https://doi.org/10.1128/IAI.00726-10>.
7. Schmidt H, Hensel M. 2004. Pathogenicity islands in bacterial pathogenesis. *Clin Microbiol Rev* 17:14–56. <https://doi.org/10.1128/CMR.17.1.14-56.2004>.
8. Li Y, Zheng H, Liu Y, Jiang Y, Xin J, Chen W, Song Z. 2011. The complete genome sequence of *Mycoplasma bovis* strain Hubei-1. *PLoS One* 6:e20999. <https://doi.org/10.1371/journal.pone.0020999>.
9. Lee CA. 1996. Pathogenicity islands and the evolution of bacterial pathogens. *Infect Agents Dis* 5:1–7.