



Whole-Genome Sequence of a *Brucella melitensis* Strain Isolated from Sheep in Saudi Arabia

Majed F. Alghoribi,^{a,b} Kamal H. Zidan,^d Abdulrahman A. Alswaji,^a Ali N. Alhafufi,^d Abdalla Ahmed,^e Hanan H. Balkhy^{a,b,c}

^aKing Abdullah International Medical Research Center, Infectious Diseases Research Department, Riyadh, Saudi Arabia

^bKing Saud bin Abdulaziz University for Health Sciences, Riyadh, Saudi Arabia

^cInfection Prevention and Control Department, King Abdulaziz Medical City, Riyadh, Saudi Arabia

^dRiyadh Veterinary Diagnostic Laboratory, Ministry of Environment, Water and Agriculture, Riyadh, Saudi Arabia

^eDepartment of Microbiology, College of Medicine, Umm Al-Qura University, Makkah, Saudi Arabia

ABSTRACT The intracellular Gram-negative bacterium *Brucella melitensis* causes a zoonotic disease in humans originating from animals. Here, we report the whole-genome sequence (WGS) of *Brucella melitensis* strain KSA_BM_07, isolated from sheep in March 2017 in Huraymila, Kingdom of Saudi Arabia.

Brucella species are facultative Gram-negative intracellular pathogens that cause brucellosis in both animals and humans. Brucellosis is a contagious zoonosis that affects a wide range of animals, including sheep, goats, cattle, camels, horses, pigs, and dogs, and leads to negative health and economic impacts in wild and domestic livestock. Transmission of *Brucella* infection from animals to humans remains a serious public health problem worldwide, especially across low-income countries (1). *Brucella melitensis* is one of the most common causes of human brucellosis through direct contact with infected animals, products of conception, or animal discharge or consumption of raw dairy or meat products of infected animals (2). Brucellosis is a major agricultural and public health issue in Saudi Arabia with a high number of cases reported every year (2–8). In this report, the whole-genome sequence (WGS) was determined for *B. melitensis* strain KSA_BM_07, isolated from the aborted fetus of a sheep at the late stage of pregnancy (at approximately 4 months of gestation) in March 2017 in Huraymila (Riyadh Region, Kingdom of Saudi Arabia). Isolation, identification, and sequencing of the isolate was done at the Riyadh Veterinary Diagnostic Laboratory at the Ministry of Environment, Water and Agriculture. The isolate was cultured from aborted fetal stomach contents on *Brucella* selective agar and blood agar incubated under aerobic and microaerophilic conditions using a CO₂-generating kit (Oxoid, UK) for 48 h at 37°C. Isolate identification was confirmed as *B. melitensis* by using real-time PCR (TIB MOLBIOL, Berlin, Germany).

The genomic DNA of *B. melitensis* strain KSA_BM_07 was extracted using a MagNA Pure compact nucleic acid isolation kit I from fresh culture. DNA libraries were constructed using the Nextera XT DNA library preparation kit (Illumina, San Diego, CA) using a 250-bp paired-end library.

A total of 3,658,132 paired-end reads were generated using an Illumina MiSeq sequencing platform. The quality of the paired-end sequencing reads was assessed using FastQC (version 0.11.5) (9) and low-quality reads were removed using Trimmomatic (version 0.38) (10). The *de novo* assembly was performed using SPAdes (version 3.11.1) (11) and Unicycler (version 3.0) (12). The assembly was compared with available WGS of *B. melitensis* isolates in the NCBI BioProject database (BioProject number [PRJNA347914](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA347914)), which were isolated in Germany from patients of Middle Eastern origin

Received 30 August 2018 Accepted 31 October 2018 Published 29 November 2018

Citation Alghoribi MF, Zidan KH, Alswaji AA, Alhafufi AN, Ahmed A, Balkhy HH. 2018. Whole-genome sequence of a *Brucella melitensis* strain isolated from sheep in Saudi Arabia. *Microbiol Resour Announc* 7:e01189-18. <https://doi.org/10.1128/MRA.01189-18>.

Editor John J. Dennehy, Queens College

Copyright © 2018 Alghoribi 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 Majed F. Alghoribi, Alghoribima@nha.med.sa.

(13). The WGS of the BwIM_SYR_04 isolate (GenBank accession numbers [CP018512](#) and [CP018513](#)) showed ~99% identity to those of our isolates.

The contigs of *B. melitensis* strain KSA_BM_07 have been constructed using Mauve Contig Mover software (14) to relatively reorder the contigs that make up a whole genome based on comparison to the close reference genome of strain BwIM_SYR_04 (13). The WGS of *B. melitensis* strain KSA_BM_07 yielded 21 contigs (with an N_{50} length of 251,053 bp) which are composed of 2 chromosomes, chromosome I (14 contigs) and chromosome II (7 contigs). The total genome size was 3,290,234 bp for chromosome I (2,111,182 bp) and chromosome II (1,179,052 bp). The G+C content of *B. melitensis* strain KSA_BM_07 is 57%, and the average sequence coverage depth is 247.279. Multilocus sequence typing (MLST) indicated that the strain belonged to sequence type 8 (ST8) based on 9 loci (MLST9) and ST38 based on 21 loci (MLST21).

In summary, only one WGS of human *B. melitensis* from Saudi Arabia has been reported previously (13). Here, we report the first WGS of animal *B. melitensis* isolated from sheep in Huraymila (Riyadh Region, Kingdom of Saudi Arabia).

Data availability. The genome sequence of *B. melitensis* strain KSA_BM_07 was deposited in GenBank under the accession number [QVMF00000000](#).

REFERENCES

- Mcdermott JJ, Grace D, Zinsstag J. 2013. Economics of brucellosis impact and control in low-income countries. *Rev Sci Tech* 32:249–261. <https://doi.org/10.20506/rst.32.1.2197>.
- Musallam II, Abo-Shehada MN, Hegazy YM, Holt HR, Guitian FJ. 2016. Systematic review of brucellosis in the Middle East: disease frequency in ruminants and humans and risk factors for human infection. *Epidemiol Infect* 144:671–685. <https://doi.org/10.1017/S0950268815002575>.
- Aloufi AD, Memish ZA, Assiri AM, McNabb SJN. 2016. Trends of reported human cases of brucellosis, Kingdom of Saudi Arabia, 2004–2012. *J Epidemiol Glob Health* 6:11–18. <https://doi.org/10.1016/j.jegh.2015.09.001>.
- Al Ali AM, Alluwaimi AM. 2009. The incidents of human brucellosis in Al-Ahsaa area, Saudi Arabia. *Sci J King Faisal Univ* 10:115–121.
- Ismaeil A, Alrheam AA, Saad Z, Shehri A, Ibrahim A, Elneam A, Cruz CP, Ali AI. 2015. Human brucellosis incidence trends in central Saudi Arabia. *Int J Adv Res* 3:1580–1586.
- Ageely H, Bani I, Gaffar A, Eltigani M, Yassin AO, Said B, Mahfouz MS. 2016. Prevalence and risk factors for brucellosis in Jazan Province, Saudi Arabia. *Trop J Pharm Res* 15:189–194. <https://doi.org/10.4314/tjpr.v15i1.26>.
- Pappas G, Papadimitriou P, Akritidis N, Christou L, Tsianos EV. 2006. The new global map of human brucellosis. *Lancet Infect Dis* 6:91–99. [https://doi.org/10.1016/S1473-3099\(06\)70382-6](https://doi.org/10.1016/S1473-3099(06)70382-6).
- Memish ZA, Venkatesh S. 2001. Brucellar epididymo-orchitis in Saudi Arabia: a retrospective study of 26 cases and review of the literature. *BJU Int* 88:72–76. <https://doi.org/10.1046/j.1464-410x.2001.02243.x>.
- Andrews S. 2010. FastQC: a quality control tool for high throughput sequence data. Babraham Bioinformatics, Cambridge, United Kingdom.
- Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. *Bioinformatics* 30:2114–2120. <https://doi.org/10.1093/bioinformatics/btu170>.
- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Pribelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 19:455–477. <https://doi.org/10.1089/cmb.2012.0021>.
- Wick RR, Judd LM, Gorrie CL, Holt KE. 2017. Unicycler: resolving bacterial genome assemblies from short and long sequencing reads. *PLoS Comput Biol* 13:e1005595. <https://doi.org/10.1371/journal.pcbi.1005595>.
- Georgi E, Walter MC, Pfalzgraf MT, Northoff BH, Holdt LM, Scholz HC, Zoeller L, Zange S, Antwerpen MH. 2017. Whole genome sequencing of *Brucella melitensis* isolated from 57 patients in Germany reveals high diversity in strains from Middle East. *PLoS One* 12:e0175425. <https://doi.org/10.1371/journal.pone.0175425>.
- Rissman AI, Mau B, Biehl BS, Darling AE, Glasner JD, Perna NT. 2009. Reordering contigs of draft genomes using the Mauve aligner. *Bioinformatics* 25:2071–2073. <https://doi.org/10.1093/bioinformatics/btp356>.