

Journal of Bacteriology

Genomic Analysis of *Mycobacterium abscessus* Strain M139, Which Has an Ambiguous Subspecies Taxonomic Position

Yun Fong Ngeow, Wei Yee Wee, Yan Ling Wong, Joon Liang Tan, Chia Su Ongi, Kee Peng Ng and Siew Woh Choo
J. Bacteriol. 2012, 194(21):6002. DOI: 10.1128/JB.01455-12.

Updated information and services can be found at:
<http://jb.asm.org/content/194/21/6002>

REFERENCES

These include:

This article cites 9 articles, 0 of which can be accessed free at:
<http://jb.asm.org/content/194/21/6002#ref-list-1>

CONTENT ALERTS

Receive: RSS Feeds, eTOCs, free email alerts (when new articles cite this article), [more»](#)

Information about commercial reprint orders: <http://journals.asm.org/site/misc/reprints.xhtml>
To subscribe to to another ASM Journal go to: <http://journals.asm.org/site/subscriptions/>

Journals.ASM.org

Genomic Analysis of *Mycobacterium abscessus* Strain M139, Which Has an Ambiguous Subspecies Taxonomic Position

Yun Fong Ngeow,^a Wei Yee Wee,^b Yan Ling Wong,^a Joon Liang Tan,^b Chia Su Ongi,^c Kee Peng Ng,^a and Siew Woh Choo^b

Department of Medical Microbiology, Faculty of Medicine, University of Malaya, Kuala Lumpur, Malaysia^a; Dental Research and Training Unit, Faculty of Dentistry, University of Malaya, Kuala Lumpur, Malaysia^b; and Faculty of Information Science and Technology, Multimedia University, Melaka, Malaysia^c

***Mycobacterium abscessus* is a ubiquitous, rapidly growing species of nontuberculous mycobacteria that colonizes organic surfaces and is frequently associated with opportunistic infections in humans. We report here the draft genome sequence of *Mycobacterium abscessus* strain M139, which shows genomic features reported to be characteristic of both *Mycobacterium abscessus* subsp. *abscessus* and *Mycobacterium abscessus* subsp. *massiliense*.**

The nontuberculous mycobacteria (NTM) are gaining importance as human pathogens, as they are being recovered from an increasingly wide spectrum of clinical material (2, 6, 9). Many of them show intrinsic resistance to antituberculosis drugs currently in use, and their identification beyond the species level is sometimes necessary to predict their response to antibiotic therapy (3). The species *Mycobacterium abscessus*, which has been associated with skin and soft tissue lesions, as well as pulmonary and other deep-seated sepsis (5), has been subdivided into subspecies by DNA sequence variation detected by molecular tests based on individual genes or a combination of genes (7). We had shotgun sequenced another genome for this species, from strain M139 isolated from the sputum sample of a 26-year-old Nepalese male presenting with hemoptysis using the Illumina GA 2X technology. With this technology, we generated 4,086,927 sequencing reads. These sequences were processed and assembled with Genomics Workbench 4.9, resulting in 42 contigs (36 contigs have a genomic size of >500 bp) with an N_{50} contig size of 645,701 bp. The M139 genome includes 5,046,600 bp with a GC content of 64%.

We compared the sequenced genome of M139 with the published reference genome of *M. abscessus* strain ATCC 19977 in order to search for common genomic regions (core regions) and strain-specific regions (accessory regions), using the Pan-genomic Sequence (Panseq) web server (4). Information from these regions is expected to provide insight into species diversity, genetic determinants of basic lifestyle, and strain-specific properties such as environmental survival (8). In our analysis, Panseq identified core regions with a total size of 4,857,651 bp and a GC content of 64% and accessory regions (M139 strain specific) of 481,453 bp with a GC content of 61%. Further annotation using the Rapid Annotation using Subsystems Technology (RAST) pipeline (1) revealed 4,675 coding sequences (CDS) and 43 RNAs in the core genome. There are 419 putative genes distributed in the subsystem of cofactors, vitamins, prosthetic groups, and pigments and 254 genes in the subsystem of carbohydrates. In the accessory genome, there are 525 CDS, including 10 putative genes in the subsystem of phages, prophages, transposable elements, and plasmids which might contribute to species diversity.

To determine the subspecies classification for M139, we performed a phylogenetic analysis first with *rpoB* and *hsp65* genes

separately and then with both genes concatenated. Both phylogenetic trees showed M139 clustering with *Mycobacterium abscessus* subsp. *massiliense*. This subspecies has been reported to be associated with *erm41* (erythromycin ribosome methyltransferase) gene deletions at positions 64 and 65, as well as a 276-bp deletion after position 158 (3). These features have been used to differentiate *M. abscessus* subsp. *massiliense* from the other *M. abscessus* subspecies of *M. abscessus sensu stricto* and *Mycobacterium bolletii*. However, we found no deletions at these positions in the *erm41* gene in strain M139. Conversely, the *erm41* gene sequence of M139 shows the nucleotide variations observed by Kim et al. (3) in the *erm41* gene of *M. abscessus sensu stricto*. These conflicting preliminary genomic observations will be further investigated with the use of other gene targets, and on a larger number of strains.

Nucleotide sequence accession numbers. The *M. abscessus* strain M139 genome sequence and annotation data have been deposited in NCBI GenBank under the accession number AKVR00000000. The version described in this paper is the first version, AKVR01000000.

ACKNOWLEDGMENTS

This work was supported by research grants UM.C/625/1/HIR/004 and UM.C/HIR/MOHE/08 from the University of Malaya, Kuala Lumpur, Malaysia.

REFERENCES

1. Aziz RK, et al. 2008. The RAST server: Rapid Annotations using Subsystem Technology. *BMC Genomics* 9:75.
2. Herdman AV, Steele JCH. 2004. The new mycobacterial species. Emerging or newly distinguished pathogens. *Clin. Lab. Med.* 24:651–690.
3. Kim H-Y, et al. 2010. *Mycobacterium massiliense* is differentiated from *Mycobacterium abscessus* and *Mycobacterium bolletii* by erythromycin ribosome methyltransferase gene (*erm*) and clarithromycin susceptibility patterns. *Microbiol. Immunol.* 54:347–353.

Received 12 August 2012 Accepted 22 August 2012

Address correspondence to Yun Fong Ngeow, yunngeow@um.edu.my, or Siew Woh Choo, lchoo@um.edu.my.

Copyright © 2012, American Society for Microbiology. All Rights Reserved.

doi:10.1128/JB.01455-12

4. Laing C, et al. 2010. Pan-genome sequence analysis using Panseq: an online tool for the rapid analysis of core and accessory genomic regions. *BMC Bioinformatics* 11:461.
5. Morales P, et al. 2007. Successful recovery after disseminated infection due to *Mycobacterium abscessus* in a lung transplant patient: subcutaneous nodule as first manifestation. *Transplant. Proc.* 39:2413–2415.
6. Piersimoni C. 2012. Nontuberculous mycobacteria infection in solid organ transplant recipients. *Eur. J. Clin. Microbiol. Infect. Dis.* 31:397–403.
7. Simmon KE, et al. 2011. *Mycobacterium chelonae-abscessus* complex associated with sinopulmonary disease, Northeastern U. S. A. *Emerg. Infect. Dis.* 17:1692–1700.
8. Tettelin H, Riley D, Cattuto C, Medini D. 2008. Comparative genomics: the bacterial pan-genome. *Curr. Opin. Microbiol.* 12:472–477.
9. Wallace RJ, Jr, Brown BA, Griffith DE. 1998. Nosocomial outbreaks/pseudo-outbreaks caused by nontuberculous mycobacteria. *Annu. Rev. Microbiol.* 52:453–490.