

Support from Phylogenomic Networks and Subspecies Signatures for Separation of *Mycobacterium massiliense* from *Mycobacterium bolletii*

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Mycobacterium abscessus subspecies classification has important clinical implications. We used phylogenomic network and amino acid analyses to provide evidence for the separation of *Mycobacterium bolletii* and *Mycobacterium massiliense* into two distinct subspecies which can potentially be differentiated rapidly by their protein signatures.

Mycobacterium abscessus has become one of the most frequently isolated nontuberculous mycobacterium (NTM) in clinical laboratories. It is associated with chronic, recurrent infections that are difficult to treat, partly because of its resistance to many of the usual medications for NTM infections. This species was previously divided into three subspecies (*M. abscessus*, *M. massiliense*, and *M. bolletii*) based on biological and genetic differences (1–3). Currently, however, only two subspecies are recognized; while *M. abscessus* is retained as *Mycobacterium abscessus* subsp. *abscessus*, *M. massiliense* and *M. bolletii* are placed in the same subspecies designated *Mycobacterium abscessus* subsp. *bolletii* (4). This tenuous merging of *M. massiliense* and *M. bolletii* is still being debated as recent publications support the previous three-subspecies classification (5). Here, we present more evidence for the retention of the former three-subspecies taxonomic division, which correlates better with the expected treatment outcomes in infected patients (6).

(This research was conducted by J. L. Tan in partial fulfillment of

the requirements for a Ph.D. from University of Malaya, Kuala Lumpur, Malaysia.)

For our genomic and amino acid analyses, we used 12 genomes from strains isolated in the Diagnostic Microbiology Laboratory

Received 26 February 2015 Returned for modification 3 May 2015

Accepted 9 June 2015

Accepted manuscript posted online 8 July 2015

Citation Tan JL, Ngeow YF, Choo SW. 2015. Support from phylogenomic networks and subspecies signatures for separation of *Mycobacterium massiliense* from *Mycobacterium bolletii*. *J Clin Microbiol* 53:3042–3046. doi:10.1128/JCM.00541-15.

Editor: G. A. Land

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Supplemental material for this article may be found at <http://dx.doi.org/10.1128/JCM.00541-15>.

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doi:10.1128/JCM.00541-15

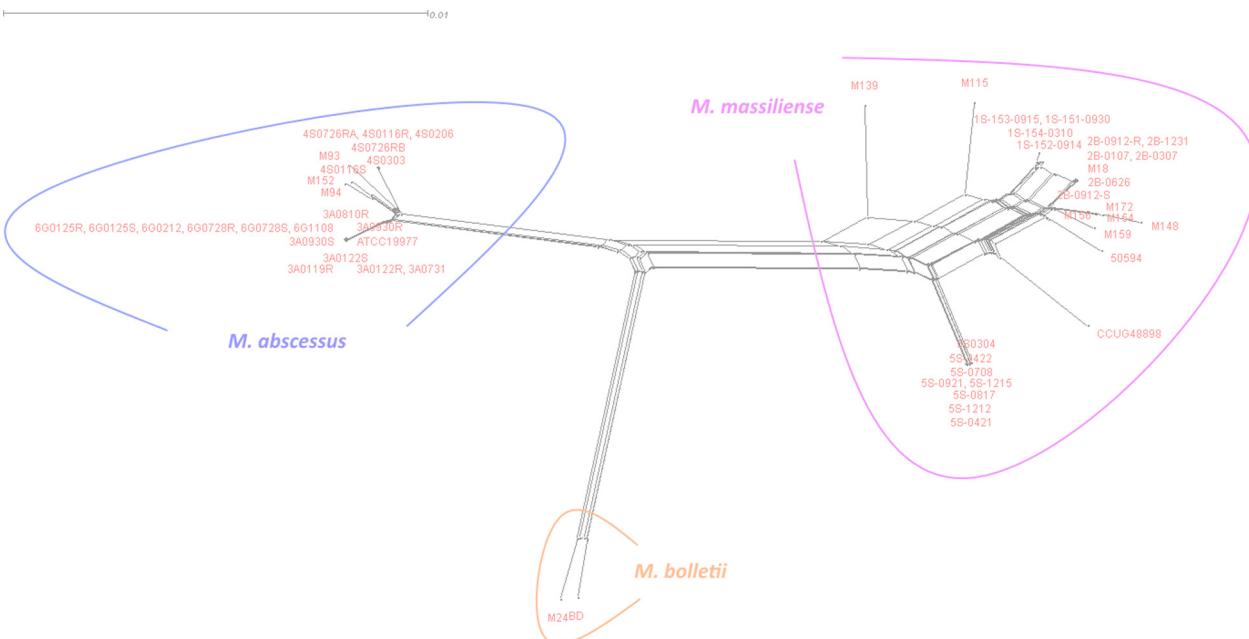


FIG 1 Phylogenomic split network tree obtained from the concatenation of single-copy core genes from *M. abscessus* subspecies. *M. massiliense* (right), *M. bolletii* (center), and *M. abscessus* (left) can be seen clearly as distinct groups.

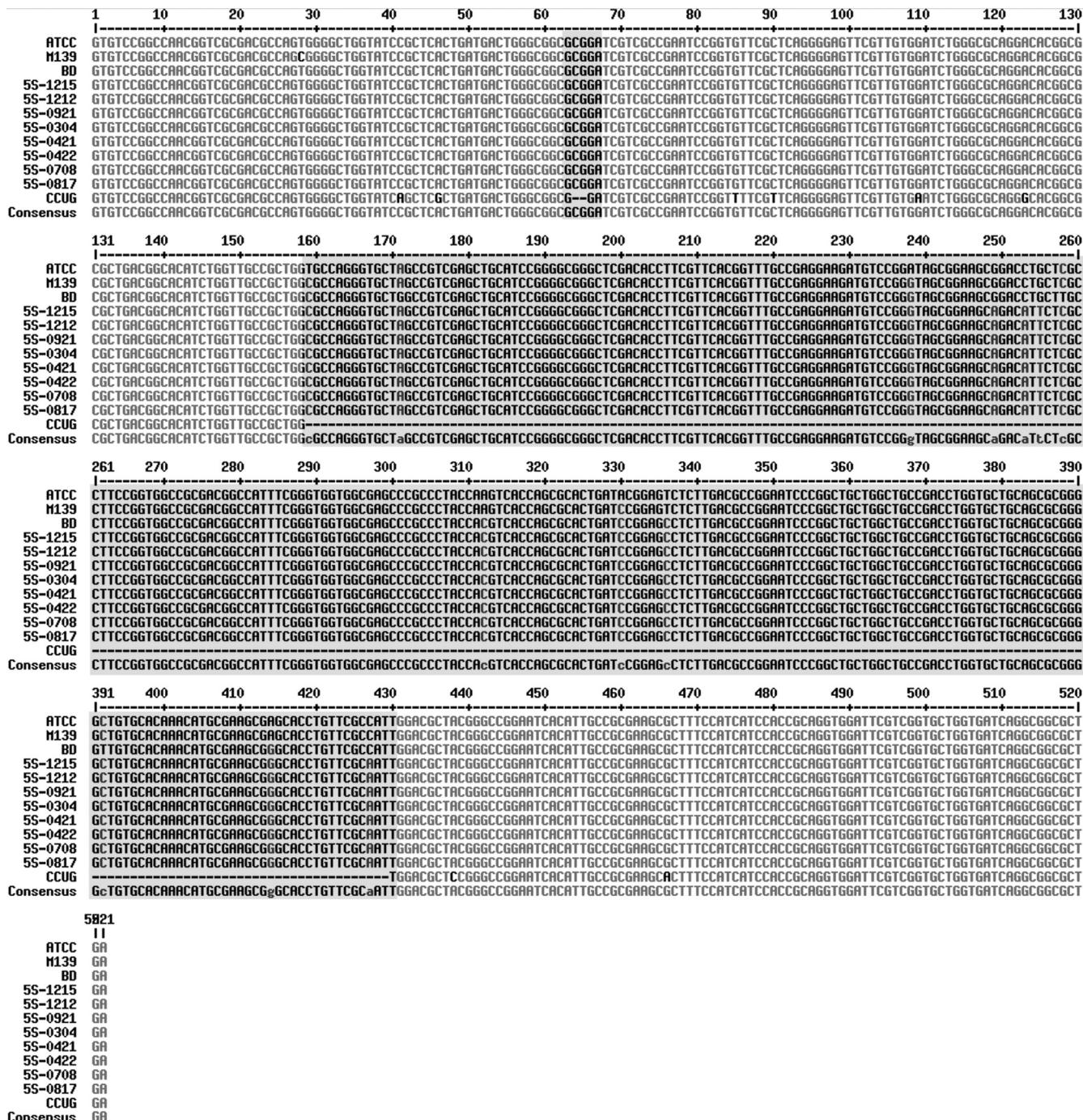


FIG 2 Multiple sequence alignments of *erm41* showing features of *M. massiliense* M139 and 55 strains compared to those of the type strains of *M. abscessus* ATCC 19977^T, *M. massiliense* CCUG 48898^T, and *M. bolletii* BD^T. The *M. massiliense* signatures are (i) deletions at positions 64 and 65 and (ii) a 274-bp deletion after position 159.

of the University of Malaya Medical Centre (UMMC), Kuala Lumpur, Malaysia, and 41 downloaded from the NCBI Genome database on July 2014 (see Table S1 in the supplemental material). Eleven of the UMMC strains have been previously reported to be *M. abscessus* (M93, M94, and M152), *M. bolletii* (M24), and *M. massiliense* (M18, M115, M152, M172, M159, M156, and M148). One strain, M139, was shown to have an ambiguous taxonomic position in a number of studies (7, 8).

The protein sequences for all strains were retrieved using the

self-training structural annotation algorithm of GeneMarkS (9). To define orthologous sequences, we used the CD-HIT program (10) with the following criteria: word length of 2, local sequence identity threshold of 0.4, alignment coverage for both sequences of 0.4, and greedy algorithm off. We also used the BLASTClust program with the following parameters: reference and query sequences must cover at least 40% of the aligned sequence and reference and query sequences must have a minimum identity of 40% (11). To reduce false-positive results due to algorithmic er-

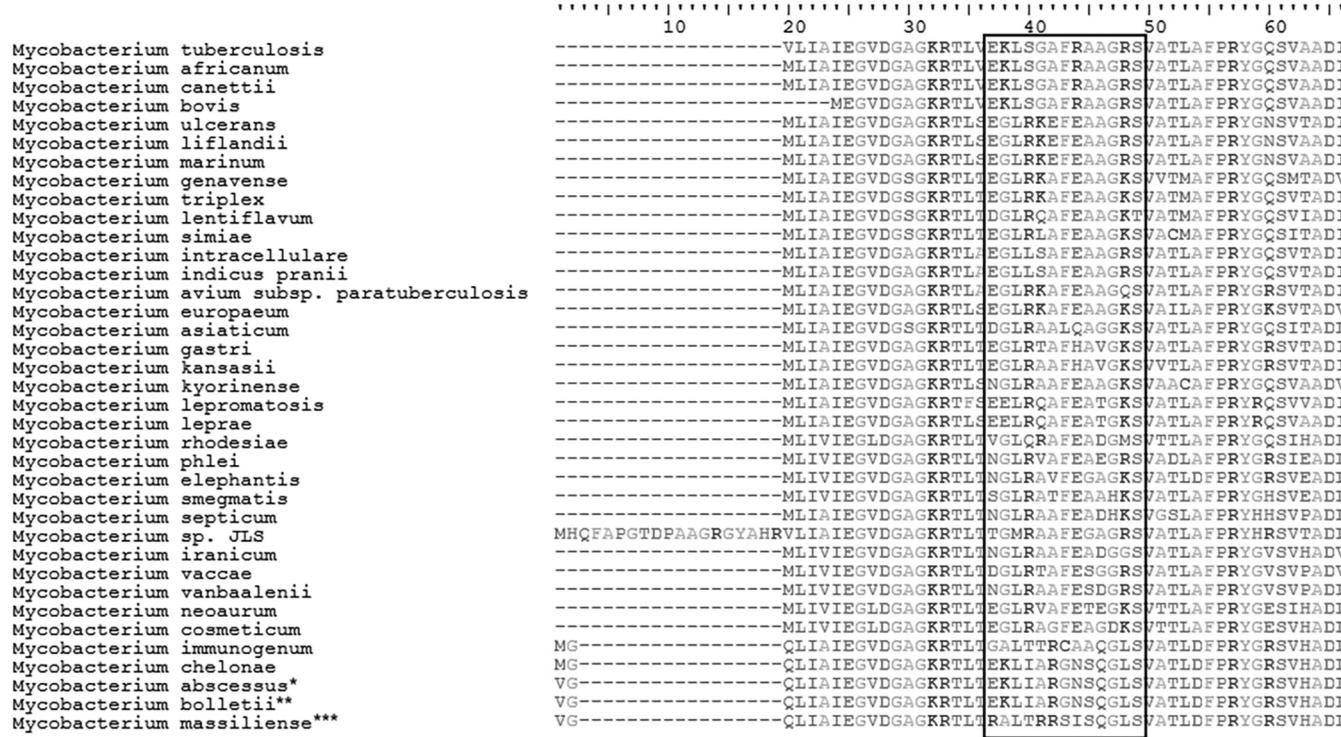


FIG 3 Consistent protein signatures in *M. massiliense* identified in multiple alignments of thymidylate kinase from *M. abscessus* and other selected mycobacteria: *23 strains of *M. abscessus* subsp. *abscessus*; **2 strains of *M. bolletii*; ***28 strains of *M. massiliense*.

rors, only the consensus sequences from both programs were extracted and used as the final list of orthologs. Nonduplicated conserved protein orthologs were aligned in MAFFT (12).

The protein sequence alignments were used as the reference for codon alignments in PAL2NAL (13). The aligned nucleotide sequences were concatenated into supersequences for phylogenomic analysis using the Neighbor-Net algorithm implemented in SplitsTree4 (14). This algorithm was considered the best for the resolution of complex taxonomy (15). To assess the subspecies classification derived from our network tree, we looked for subspecies-specific polymorphisms previously described for the erythromycin ribosome methyltransferase (*erm41*) and 16S to 23S internal transcribed spacer (ITS) genes.

Our network-based phylogenomic tree showed reticulated branches leading to three clearly distinctive monophyletic groups representing the three subspecies of *M. abscessus* (Fig. 1). The M139 and the 5S strains (5S-0421, 5S-0422, 5S-0708, 5S-0817, 5S-0921, 5S-1212, 5S-1215, and 5S-0304) clustered with the other *M. massiliense* strains. None of the branches in any of the three major clusters bifurcated to the other two clusters. The presence of 3-dimensional-like splits within the branches indicated incompatible phylogenetic signals that are likely to be the result of recombination following the horizontal transfer of genetic material among strains. Indeed, the recombination among our *M. abscessus* strains is statistically supported by the pairwise homoplasy index (PHI) ($P = 0$) (16). The incompatible signals occurred at random points in the tree, suggesting that recombination has occurred in ancestral states and within the respective subspecies. We also noticed unusual conflicting signals within the *M. massiliense* cluster, appearing as a major reticulation connecting the *M. massiliense* strains and suggesting a higher degree of genetic re-

combination in *M. massiliense* compared to that in the other two subspecies. To test the validity of this network phylogenomics approach, we used it on three members of the *M. avium* complex and found a clear separation of *Mycobacterium avium* subsp. *paratuberculosis*, *Mycobacterium avium* subsp. *hominissuis*, and *Mycobacterium avium* subsp. *avium* into three distinctive monophyletic groups, as observed with the *M. abscessus* complex (see Fig. S1 in the supplemental material).

M. massiliense is known to be different from the other two subspecies in having a truncated *erm41* with nucleotide deletions at the 64th to 65th and 159th to 432nd positions, as well as mutations in the ITS (a A to G substitution at the 60th position and a C insertion at the 102nd position) (2). M139 and the eight 5S strains previously classified as *M. massiliense* and appearing as *M. massiliense* in our phylogenomic tree did not show the *erm41* features associated with *M. massiliense* (Fig. 2). M139 additionally lacked the ITS mutations characteristic of *M. massiliense* and did not show inducible resistance to macrolides (17). Overall, however, there was good concordance (83%) between the subspecies classifications by *erm41* signatures and by the network tree.

In the multiple sequence alignment of the orthologous proteins from our 53 strains, we noted 46 proteins with at least one amino acid that can be used to differentiate the three subspecies (see Table S2 in the supplemental material) and another two proteins (thymidylate kinase [tk] and 30S ribosomal protein S3 [S3]) that can differentiate *M. massiliense* from the other two subspecies (Fig. 3 and 4). We used BLAST to search the amino acid sequences of tk and S3 against all *Mycobacterium* genomes in the NCBI database and found them in 37 and 44 species, respectively. After realigning against these mycobacterial species, we confirmed the amino acid signatures of tk (RALTRRSISQGLS at position 20 to

Mycobacterium tuberculosis
 Mycobacterium africanum
 Mycobacterium canetti
 Mycobacterium asiaticum
 Mycobacterium kansasii
 Mycobacterium gastri
 Mycobacterium liflandii
 Mycobacterium marinum
 Mycobacterium triplex
 Mycobacterium genavense
 Mycobacterium yongonense
 Mycobacterium intracellulare
 Mycobacterium indicus pranii
 Mycobacterium lentiflavum
 Mycobacterium simiae
 Mycobacterium nebraskense
 Mycobacterium europaeum
 Mycobacterium gilvum
 Mycobacterium iranicum
 Mycobacterium rufum
 Mycobacterium obuense
 Mycobacterium vanbaalenii
 Mycobacterium kyorinense
 Mycobacterium xenopi
 Mycobacterium chubuense
 Mycobacterium vaccae
 Mycobacterium sp. JLS
 Mycobacterium smegmatis
 Mycobacterium setense
 Mycobacterium septicum
 Mycobacterium rhodesiae
 Mycobacterium tusciae
 Mycobacterium phlei
 Mycobacterium mageritense
 Mycobacterium aromaticivorans
 Mycobacterium avium
 Mycobacterium elefantis
 Mycobacterium immunogenum
 Mycobacterium leprae
 Mycobacterium hassiacum
 Mycobacterium cheloneae
 Mycobacterium bolletii**
 Mycobacterium abscessus*
 Mycobacterium massiliense***

| | 240 | 250 | 260 | 270 | 280 | 290 | 300 |
|----------------------------------|-------------|--------|---------------------------|--------------|-------------|-------|---------|
| RPRSGASGTTAT-GTDAGRAAGG | EE- | - | - | AAPDAAA | PV | - | EAQSTES |
| RPRSGASGTTAT-GTDAGRAAGG | EE- | - | - | AAPDAAA | PV | - | EAQSTES |
| RPRSGASGTTAT-GTDAGRAAGG | EE- | - | - | AAPDAAA | PV | - | EAQSTES |
| RPRSGASGTTAT-GTDAGRAAGG | EEGSAPA | - | - | AAEAAA | PAV | - | EAQSTES |
| RPRSGASGTTAT-GTDAGRAAGG | EEG-TA | - | - | AVGNAAA | PAV | - | EAQSTE- |
| RPRSGASGTTAT-GTDAGRAAGS | EEG-TA | - | - | AAGNEAA | PAV | - | EAQSTE- |
| RPRSGASGTTAT-GTEARRAVGS | EE-PA | - | - | AAESATT | P | - | EAQSTES |
| RPRSGASGTTAT-GTEAGRAGVS | EE-PA | - | - | AAESATT | P | - | EAQSTES |
| RPRSGASGTTAT-STEAGRAADAG | - | - | - | EPPADSAP | P | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAADAG | - | - | - | EPAADSAP | P | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAASA | EEG- | - | - | AASAAA | PA | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAASV | EEG- | - | - | AAAAA | PAA | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAASA | EEG- | - | - | AAAAA | PAA | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAAGA | EEN-TA | - | - | NAAAESA | PAP | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAAGAEG | TAC | TET-TA | - | TAAAEGA | PA | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAAGA | EEA-AAA | - | - | TAAAAT | TA | - | EPQSTES |
| RPRSGASGTTAT-STEAGRAAGA | EES-TA | - | - | TAAAETP | FAA | - | EPQSTES |
| RPRSGASGTTAT-STDAGRAAT | EEA-PA | - | - | TDAAAATA | PAAG | - | QPETTES |
| RPRSGASGTTAT-STDAGRAAS | EEA-PA | - | - | PEAAA | AA | - | AAESTES |
| RPRSGASGTTAT-STDAGRAATEGS | VEA-PAV | - | - | AEASVG | TEAA | - | AGESTES |
| RPRSGASGTTAT-STDAGRAATESP | AEA-PAA | - | - | VEATAGA | P | - | APESTES |
| RPRSGASGTTAT-STDAGRAASEGT | VEA-PA | - | - | TEAAAATAPSAG | - | - | QPETTES |
| RPRSGAAGTTAT-STDAGRAASGG | EEA- | - | - | TAAAATP | FAAEEQAET | QSTES | |
| RPRSGAAGTTGA-TTEAGRAAGA | EEA- | - | - | AAAPASA | EFAL | - | ETQSTES |
| RPRSGASGTTAT-STDAGRAASESP | AEA-PAI | - | - | AEATQG | TEAAAGTSAE | - | TTENSGS |
| RPRSGASGTTAT-STDAGRA | AEA-PAV | - | - | AEATQG | TEAAAAGAAAE | - | TTENSGS |
| RPRSGASGTTAT-STDAGRAASGT | QEA-PAA | - | - | AEAAA | GTEAAAGAAET | - | TTQNPGS |
| RPRSGASGTTAT-STEAGRAART | SDA-PAA | - | - | GTAAAAA | EAP | - | AESTES |
| RPRSGASGTTAT-STEAGRAATG | DDA-SSA | - | - | TEAAAAS | AP | - | AAESTES |
| RPRSGASGTTAT-STEAGRAATG | DDA-PAA | - | - | TEAAAAS | AP | - | AAESTES |
| RPRSGASGTTAT-STEAGRAAT | EEA-PAAPAVA | EAT | TTGTEAAAGDAA | - | - | - | APESTES |
| RPRSGASGTTAT-STEAGRAAT | EEA-PAAPAVA | EAT | TTGTEAAAGQASPTQTTEAPSSEES | - | - | - | |
| RPRSGSSGTTAT-STEAGRAATGT | EDA-PAA | - | - | AEATGATEAAA | QA | - | ATENTES |
| RPRSGSSGTTAT-STEAGRAA | EETAESA | - | - | VPATA | EA | - | SAENTES |
| RPRSGASGTTAT-STEAGRAAS | EETAESA | - | - | VPVTAA | EAPAVEP | - | AAENTES |
| RPRSGASGTTAT-STDAGRAAES | TEN-TAV | - | - | QENAATAE | Q | - | ATQSTES |
| RPRSGAAGNTAT-GTEAGRAASGVDTGRAAAC | AEAA-PAV | - | - | AEAGGTE | AAAGAAET | - | TTENTES |
| RPRSGASGTTAT-STEAGRAAV | ENT- | - | - | ATPDASA | - | - | AETSTES |
| RPRSGAAGTGT-VTDAGRAVGG | EES-AATN | - | - | IGHSSD | SVVT | H | EPQIAES |
| RPRSGASGTTATGSTDAGRAATSG | ETA-TAV | - | - | AEEAAATE | AAAGQATE | - | TTTSGES |
| RPRSGASGTTAT-STEAGRAAA | ETP- | - | - | ASDGASAP | - | - | SAETTES |
| RPRSGSSGTTAT-STEAGRAAA | ETP- | - | - | ASDGASAP | - | - | SAETTES |
| RPRSGSSGTTAT-STEAGRAAA | ETP- | - | - | ASDGASAP | - | - | SAETTES |
| RPRSGSSGTTAT-STEAGRAAA | ETG- | - | - | GNTSAE | AP | - | AETSTES |

FIG 4 Consistent protein signatures in *M. massiliense* identified in multiple alignments of 30S ribosomal protein S3 from *M. abscessus* and other selected mycobacteria: *23 strains of *M. abscessus* subsp. *abscessus*; **2 strains of *M. bolletii*; ***28 strains of *M. massiliense*.

30) and S3 (ETGGNTSAEAPAETSTES at position 260 to 277) to be specific for *M. massiliense* (Fig. 3 and 4). The presence of these signatures in M139 and the 5S strains supported their classification as *M. massiliense*, in agreement with the classification by the phylogenomic network. They will need to be experimentally verified as suitable biomarkers for the identification of *M. massiliense* in clinical material.

It is well known that *M. abscessus* subspecies exhibit different clinical and epidemiological features (18, 19). *M. massiliense* is more susceptible to antibiotics but is also more often associated with clinical infections. *M. bolletii*, on the other hand, is rarely isolated from clinical material but is more highly antibiotic resistant. While the reasons behind these differences are still unclear, there is sufficient justification for subspecies identification in patient care. Our analyses support the division of *M. abscessus* into three subspecies and the reinstatement of *M. massiliense* as a taxon independent of *M. bolletii*. The specific identification of these two subspecies which show different antibiotic susceptibilities will enable the clinician to prescribe appropriate antibiotics for the effective treatment of infections.

ACKNOWLEDGMENTS

This study was supported by research grants UM.C/625/1/HIR/MOHE/CHAN/14/4 and UM.C/HIR/MOHE/08 from the University of Malaya.

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