

Mycobacterium bovis Pulmonary Tuberculosis, Algeria

Fatah Tazerart, Jamal Saad, Abdellatif Niar, Naima Sahraoui, Michel

Drancourt

▶ To cite this version:

Fatah Tazerart, Jamal Saad, Abdellatif Niar, Naima Sahraoui, Michel Drancourt. Mycobacterium bovis Pulmonary Tuberculosis, Algeria. Emerging Infectious Diseases, 2021, 27 (3), pp.972-974. 10.3201/eid2703.191823 . hal-03666147

HAL Id: hal-03666147 https://amu.hal.science/hal-03666147v1

Submitted on 12 Oct 2023

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

Despite temporary suspension of all flights to or from Brazil from or through the United Kingdom as of December 25, 2020 (http://www.gov.uk/foreigntravel-advice/brazil), it is likely that the number of SARS-CoV-2 lineage B.1.1.7 infections in Brazil is higher than that reported. Increasing genomic surveillance of B.1.1.7 and other variants of concern that carry mutations of potential biological significance (e.g., E484K in the spike protein; C.M. Voloch, unpub data, https://www.medrxiv.org/content/10.1 101/2020.12.23.20248598v1) is imperative for monitoring vaccination effectiveness and contextualizing the epidemiology and evolution of SARS-CoV-2 in Latin America.

Acknowledgments

We thank all researchers who are working around the clock to generate and share genome data worldwide on GISAID (http://www.gisaid.org). GISAID acknowledgment tables are available at https://github.com/CADDE-CENTRE/ VOC-Lineage-Brazil.

This project was supported by a Medical Research Council-São Paulo Research Foundation (FAPESP) CADDE partnership award (MR/S0195/1 and FAPESP 18/14389-0) (http://caddecentre.org/). N.R.F. is supported by a Wellcome Trust and Royal Society Sir Henry Dale Fellowship (204311/Z/16/Z).

About the Author

Mrs. Claro is a PhD student at the Department of Infectious Disease, School of Medicine & Institute of Tropical Medicine, University of São Paulo, São, Paulo, Brazil. She is a part of the Centre for Arbovirus Discovery, Diagnostics, Genomics and Epidemiology team and has primary research interests in real-time epidemiologic surveillance of viruses of public importance for Brazil.

References

- Vogels CBF, Watkins AE, Harden CA, Brackney DE, Shafer J, Wang J, et al. SalivaDirect: a simplified and flexible platform to enhance SARS-CoV-2 testing capacity. Med. 2020 Dec 26 [Epub ahead of print]. https://doi.org/10.1016/j. medj.2020.12.010
- Rambaut A, Holmes EC, O'Toole Á, Hill V, McCrone JT, Ruis C, et al. A dynamic nomenclature proposal for SARS-CoV-2 lineages to assist genomic epidemiology. Nat Microbiol. 2020;5:1403–7. https://doi.org/10.1038/ s41564-020-0770-5
- Minh BQ, Schmidt HA, Chernomor O, Schrempf D, Woodhams MD, von Haeseler A, et al. IQ-TREE 2: new models and efficient methods for phylogenetic inference in the genomic era. Mol Biol Evol. 2020;37:1530–4. https://doi.org/10.1093/molbev/msaa015
- 4. Candido DS, Claro IM, de Jesus JG, Souza WM, Moreira FRR, Dellicour S, et al. Evolution and epidemic spread of SARS-

CoV-2 in Brazil. Science. 2020;369:1255–60. https://doi.org/10.1126/science.abd2161

- Katoh K, Standley DM. MAFFT multiple sequence alignment software version 7: improvements in performance and usability. Mol Biol Evol. 2013;30:772–80. https://doi.org/10.1093/molbev/mst010
- Shu Y, McCauley J. GISAID: Global Initiative on Sharing All Influenza Data – from vision to reality. Euro Surveill. 2017;30;22:30494.

Address for correspondence: Nuno R. Faria, St Mary's Hospital, Praed St, Paddington, London W2 1NY, UK; email: nfaria@ ic.ac.uk; and José Eduardo Levi, Avenida Juruá 548, Alphaville, Barueri, SP 06455-010, Brazil; email: jose.levi.ext@dasa.com.br

Mycobacterium bovis Pulmonary Tuberculosis, Algeria

Fatah Tazerart, Jamal Saad, Abdellatif Niar, Naima Sahraoui,¹ Michel Drancourt¹

Author affiliations: Université Ibn Khaldoun de Tiaret, Tiaret, Algeria (F. Tazerart); Université de Blida, Blida, Algeria (F. Tazerart, N. Sahraoui); Institut Hospitalo-Universitaire Méditerranée Infection, Marseille, France (F. Tazerart, J. Saad, M. Drancourt); Aix-Marseille-University, Marseille (J. Saad, M. Drancourt); Laboratoire de Reproduction des Animaux de la Ferme, Université Ibn Khaldoun de Tiaret, Tiaret (A. Niar)

DOI: https://doi.org/10.3201/eid2703.191823

We analyzed 98 *Mycobacterium tuberculosis* complex isolates collected in 2 regions of Algeria in 2015–2018 from 93 cases of pulmonary tuberculosis. We identified 93/98 isolates as *M. tuberculosis* lineage 4 and 1 isolate as *M. tuberculosis* lineage 2 (Beijing). We confirmed 4 isolates as *M. bovis* by whole-genome sequencing.

In Algeria, interpreting tuberculosis (TB) incidence, estimated at 53–88 cases/100,000 population in 2017 (1), is limited by the fact that the diagnosis relies on microscopic examination of clinical samples. Iso-

¹These authors equally contributed to this work.

lates are presumptively identified as *Mycobacterium tuberculosis* complex based on colony phenotype.

We analyzed 98 sputum isolates identified as M. tuberculosis complex by 5 Tuberculosis and Respiratory Disease Control Service facilities in 2015-2018 (Appendix Table 1, Figure, https://wwwnc.cdc. gov/EID/article/27/3/19-1823-App1.pdf). Exact tandem repeat D analysis (2) confirmed these 98 isolates as M. tuberculosis complex. Large-sequence polymorphism analysis using PCR sequencing of genomic regions RD105, RD239, and RD750 and of the polyketide synthase gene pks15/1 (3) yielded 88 (89.8%) M. tuberculosis sensu stricto Euro-American lineage 4 isolates and 1 East Asian lineage 2 (Beijing) isolate. Whole-genome sequencing (WGS) of 5 RD deletion-free unidentified isolates indicated that these 5 isolates, P9982(ERR3588223), P9983(ERR3588225), P9985(ERR3588243), P9984(ERR3588246), and P9986(ERR3588247), were M. tuberculosis sensu stricto Euro-American lineage 4. We conducted WGS analysis using TB-profiler for M. tuberculosis online tool (https://tbdr.lshtm.ac.uk/upload) for lineage and sublineage determination. Altogether, M. tuberculosis lineage 4 was the predominant lineage in the 5 Algerian departments and the sole lineage documented in Bgayet, Tizi-Ouzou, and Medea (Appendix Table 2); it was found to be the cause of pulmonary TB in 79/93 (85%) cases, pleural TB in 11 (12%) cases, and lymph node TB in 3 (3%) cases. These observations updated those issued from a previous study conducted in 14 departments including 114 (88%) cases of pulmonary localization and 15 cases (12%) of extrapulmonary localization (4). In a later study, spoligotyping revealed that most isolates belonged to *M. tuberculosis* Euro-American lineage 4; the Haarlem clade accounted for 29.5% of studied isolates; the Latin American-Mediterranean clade, 25.6%; and the T clade, 24.8% (4). In our study, 1 *M. tuberculosis* Beijing strain was isolated from a bronchial fluid sample collected in Blida from the location at which 15 *M. tuberculosis* Beijing isolates had been identified ≈10 years earlier from 14 workers from Algeria and 1 from China (5). Our observation suggests that 10-year circulation of *M. tuberculosis* Beijing strain in the community in Blida area most probably followed immigration of workers from China employed in the construction sector.

WGS analysis of 4 additional isolates exhibiting a 6-bp deletion in the *pks*15/1 gene identified them as *M. bovis*. Using a Roary pangenome pipeline (https:// sanger-pathogens.github.io/Roary), we found that *M. bovis* CSURP9981 grouped with *M. bovis* CSURP9979 and that *M. bovis* CSURP9980 grouped with *M. bovis* CSURP9978 (Figure). Further analysis based on the 3,732,808-bp core genome detected 3,761-bp (0.1%) of single-nucleotide polymorphisms (SNPs) between the 4 isolate genomes. Whole-genome sequences of *M. bovis* strains in the study have been deposited in GenBank (sequence P9978, accession no. ERR3587501; P9979, no. ERR3587591; P9980, no. ERR3587597; and P9981, no. ERR3588222).

All 4 patients had pulmonary TB and had no detectable lymph node swelling and no scrofula (6). Two case-patients in Blida were a 27-year-old unemployed man and a 60-year-old taxi driver who both declared that they did not consume raw milk and had no contacts with cattle; a neighbor of the 60-year-old patient was a butcher with whom he

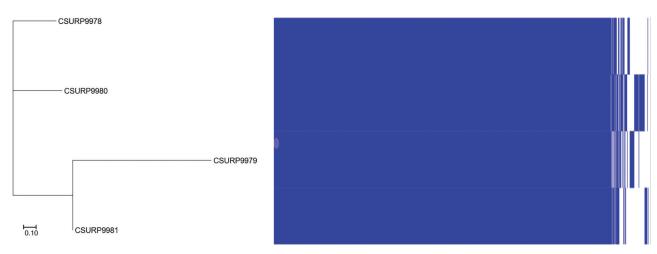


Figure. Pangenome-based tree of 4 human *Mycobacterium bovis* isolates, Algeria. The tree was generated by Roary (https://sanger-pathogens.github.io/Roary) from binary gene presence or absence in the accessory genome. Scale bar indicates 10% sequence divergence.

spent a lot of time. Two case-patients in Ain Defla were 18-year-old and 43-year-old housewives living in 2 different rural areas. The interviews of these patients did not reveal contacts with cattle. Identification of human cases of *M. bovis* was unexpected because in 50 years, only 7 cases of *M. bovis* human infection have been reported in Algeria: 3 cases of pulmonary TB and 2 cases of cervical lymphatic TB detected in a total of 1,183 (0.4%) phenotypically identified *M. bovis* isolates (7), and 2 additional cases reported in 2009 (8).

M. bovis TB is clinically, pathologically, and radiologically indistinguishable from M. tuberculosis; diagnosis requires accurate identification of the causative mycobacterium, most efficiently by using WGS. Our report illustrates pitfalls in precisely tracing the natural history of *M. bovis* TB in patients, including sources, routes of transmission, and primary route of entry, which may determine the pathology of the infection. Zoonotic M. bovis TB was most often transmitted to humans by the consumption of *M. bovis*-contaminated dairy products that caused lymphatic TB, eventually becoming pulmonary TB (9). We previously reported a hidden circumstance for contacts with M. bovis-infected animals, tracing 1 M. bovis pulmonary TB case in a patient in Tunisia to contacts with an infected sheep during religious festivities in 2018 (10). In the case we report here, foodborne transmission cannot be ruled out, but it is possible that this may be a rare case of aerosol transmission.

Algeria is a bovine TB-enzootic country. We recommend comparing the genome sequences from the 4 patients reported here with those of future bovine isolates in the same departments to trace zoonotic *M. bovis* TB in Algeria and contribute to the understanding of its natural history.

Acknowledgments

We thank Graba L., Lamri Zahir, Cherair Ali, and Kerrouche Moussa and those who are responsible for the Tuberculosis and Respiratory Disease Control Service facilities of Tizi Ouzou for their valuable assistance. We also thank Mohamed Rahal, Asma Aiza, and especially Sofiane Tahrikt.

About the Author

Dr. Tazerart is a veterinarian with a magister degree in animal epidemiology. He is an instructor at Blida University and is a PhD student at Tiaret University, Algeria. His research interests include bovine and human tuberculosis in Algeria, with a focus on characterization of animal and human *M. bovis* strains.

References

- Bouziane F, Allem R, Sebaihia M, Kumanski S, Mougari F, Sougakoff W, et al.; CNR-MyRMA. First genetic characterisation of multidrug-resistant *Mycobacterium tuberculosis* isolates from Algeria. J Glob Antimicrob Resist. 2019;19:301–7. https://doi.org/10.1016/j.jgar.2019.05.010
- Djelouadji Z, Raoult D, Daffé M, Drancourt M. A single-step sequencing method for the identification of *Mycobacterium tuberculosis* complex species. PLoS Negl Trop Dis. 2008;2:e253. https://doi.org/10.1371/journal.pntd.0000253
- Gagneux S, Small PM. Global phylogeography of Mycobacterium tuberculosis and implications for tuberculosis product development. Lancet Infect Dis. 2007;7:328–37. https://doi.org/10.1016/S1473-3099(07)70108-1
- Ifticene M, Kaïdi S, Khechiba MM, Yala D, Boulahbal F. Genetic diversity of *Mycobacterium tuberculosis* strains isolated in Algeria: results of spoligotyping. Int J Mycobacteriol. 2015;4:290–5. https://doi.org/10.1016/j.ijmyco.2015.06.004
- Ifticene M, Gacem FZ, Yala D, Boulahbal F. Mycobacterium tuberculosis genotype Beijing: about 15 strains and their part in MDR-TB outbreaks in Algeria. Int J Mycobacteriol. 2012;1:196–200. https://doi.org/10.1016/j.ijmyco.2012.10.006
- Forget N, Challoner K. Scrofula: emergency department presentation and characteristics. Int J Emerg Med. 2009;2:205–9. https://doi.org/10.1007/s12245-009-0117-8
- de Beco O, Boulahbal F, Grosset J. Incidence of the bovine bacillus in human tuberculosis at Algiers in 1969 [in French]. Arch Inst Pasteur Alger. 1970;48:93–101.
- Gharnaout M, Bencharif N, Abdelaziz R, Douagui H. Tuberculosis from *Mycobacterium bovis*: about two cases [in French]. Rev. Mal. Respir. 2009;26:141. https://www.emconsulte.com/rmr/article/197004
- Michel AL, Müller B, van Helden PD. Mycobacterium bovis at the animal-human interface: a problem, or not? Vet Microbiol. 2010;140:371–81. https://doi.org/10.1016/j.vetmic.2009.08.029
- Saad J, Baron S, Lagier JC, Drancourt M, Gautret P. *Mycobacterium bovis* pulmonary tuberculosis after ritual sheep sacrifice in Tunisia. Emerg Infect Dis. 2020;26:1605–7. https://doi.org/10.3201/eid2607.191597

Address for correspondence: Michel Drancourt, Aix Marseille Université, Institut Hospitalo-Universitaire Méditerranée Infection, 19-21 Bd Jean Moulin, 13005 Marseille, France; email: michel.drancourt@univ-amu.fr