'Congobacterium massiliense' gen. nov. sp. nov., a new bacterium isolated from the gut of a pygmy (Baka) woman

M. Bilen¹, F. Cadoret¹, Z. Daoud², P. E. Fournier¹ and D. Raoult¹

1) Aix-Marseille Université, URMITE, UM63, CNRS7278, IRD198, Inserm 1095, Institut Hospitalo-Universitaire Méditerranée-Infection, Faculté de médecine, Marseille, France and 2) Clinical Microbiology Department, Faculty of Medicine and Medical Sciences, University of Balamand, Amioun, Lebanon

Abstract

We report the main characteristics of a new genus, 'Congobacterium,' and a new species, 'Congobacterium massiliense,' strain Marseille-P3295 (CSUR P3295), a new member in the order Coribacteriacea, which was isolated from a stool sample of a healthy 50-year-old pygmy (Baka) woman. © 2016 The Authors. Published by Elsevier Ltd on behalf of European Society of Clinical Microbiology and Infectious Diseases.

Keywords: Culturomics, '*Congobacterium massiliense*', emerging bacteria, gut microbiota, human microbiota Original Submission: 20 October 2016; Revised Submission: 3 November 2016; Accepted: 9 November 2016 Article published online: 15 November 2016

Corresponding author: D. Raoult, Aix-Marseille Université, URMITE, UM63, CNRS7278, IRD198, Inserm 1095, Institut Hospitalo-Universitaire Méditerranée-Infection, Faculté de médecine, 27 Boulevard Jean Moulin, 13385, Marseille cedex 05, France **E-mail: didier.raoult@gmail.com**

In 2015, stool samples were collected from Congolese persons as part of the project aiming to describe the human gut microbiome by culturomics [1]. An approval from the ethics committee under the number 09-022 was obtained from the Institut Fédératif de Recherches IFR48 (Marseille, France).

Samples were inoculated in blood culture media after being diluted with 1 mL of phosphate-buffered saline. Then 5 mL of sheep's blood was added to the culture bottle along with 5 mL of filtered rumen and incubated at 37°C under anaerobic conditions. On day 10, strain Marseille-P3295 was isolated on 5% sheep's blood-enriched Columbia agar (bioMérieux, Marcy l'Etoile, France). Colonies were smooth with a mean diameter of 0.4 to 0.8 mm. Strain Marseille-P3295 cells were Gram-positive bacilli, catalase and oxidase negative with an average length of 1.58 µm. Colonies could not be identified by our systematic matrix-assisted desorption ionization-time of flight mass spectrometry (MALDI-TOF MS) screening on a Microflex spectrometer (Bruker Daltonics, Bremen, Germany) [2]. Thus, 16S rRNA gene sequencing was done using fDI-rP2 primers (Eurogentec, Seraing, Belgium) on a 3130-XL sequencer (Applied Biosciences, Saint Aubin, France) as previously described [3].

'Congobacterium massiliense' strain Marseille-P3295 exhibited a 93.8% 16S rRNA gene sequence similarity with Mogibacterium neglectum type strain P9a-h (Z36274), the phylogenetically closest species with standing in nomenclature (Fig. 1). Thus, strain Marseille-P3295 can be classified as a new genus because it exhibits a 16S rRNA gene sequence divergence of more than 5% with its phylogenetically closest species with a validly published name with standing in nomenclature [4]. We propose the creation of the new genus, 'Congobacterium' (con.go.bac.ter'ium, N.L. masc. gen. n. Congobacterium, for Congo, the originary country of samples out of which strain Marseille-P3295 was isolated). Marseille-P3295 is the type strain of the new species 'Congobacterium massiliense' gen. nov., sp. nov. (mas.il.i.en'se, L. gen. masc. n. massiliense, pertaining to Massilia, the antic name of the city of Marseille where this bacteria was discovered.

MALDI-TOF MS spectrum accession number

The MALDI-TOF MS spectrum of 'C. massiliense' is available online (http://www.mediterranee-infection.com/article.php? laref=256&titre=urms-database).

Nucleotide sequence accession number

The 16S rRNA gene sequence was deposited in GenBank under accession number LT623899.

New Microbe and New Infect 2017; 15: 63-64



FIG. 1. Phylogenetic tree showing position '*Congobacterium massiliense*' strain Marseille-P3295 among phylogenetically closest species. Multiple sequence alignment was performed using CLUSTALW, and phylogenetic inferences were done using maximum-likelihood method by MEGA software. Bootstraps values are shown on nodes after 500 repeats, and values less than 90% were eliminated. Scale bar indicates 2% nucleotide sequence divergence.

Deposit in a culture collection

Strain Marseille-P3295 was deposited in the Collection de Souches de l'Unité des Rickettsies (CSUR, WDCM 875) under number P3295.

Acknowledgement

This study was funded by the Fondation Méditerranée Infection.

Conflict of Interest

None declared.

References

- [I] Lagier JC, Hugon P, Khelaifia S, Fournier PE, La Scola B, Raoult D. The rebirth of culture in microbiology through the example of culturomics to study human gut microbiota. Clin Microbiol Rev 2015;28:237-64.
- [2] Seng P, Abat C, Rolain JM, Colson P, Lagier JC, Gouriet F, et al. Identification of rare pathogenic bacteria in a clinical microbiology laboratory: impact of matrix-assisted laser desorption ionizationtime of flight mass spectrometry. J Clin Microbiol 2013;51: 2182–94.
- [3] Drancourt M, Bollet C, Carlioz A, Martelin R, Gayral JP, Raoult D. 16S ribosomal DNA sequence analysis of a large collection of environmental and clinical unidentifiable bacterial isolates. J Clin Microbiol 2000;38: 3623–30.
- [4] Kim M, Oh HS, Park SC, Chun J. Towards a taxonomic coherence between average nucleotide identity and 16S rRNA gene sequence similarity for species demarcation of prokaryotes. Int J Syst Evol Microbiol 2014;64:346–51.