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### 'Bacteroides cutis,' a new bacterial species isolated from human skin

#### S. Belkacemi, N. Cassir, J. Delerce, F. Cadoret and B. La Scola

Unité de Recherche sur les Maladies Infectieuses et Tropicales Emergentes, CNRS (UMR 7278), IRD (198), INSERM (U1095), AMU (UM63), Institut Hospitalo-Universitaire Méditerranée-Infection, Faculté de médecine, Aix-Marseille Université, Marseille, France

#### Abstract

We report the main characteristics of 'Bacteroides cutis' sp. nov., strain Marseille-P4118<sup>T</sup> (= CSUR P4118), a new species within the genus *Bacteroides*. This strain was isolated from a skin sample of a 75-year-old man from Marseille. © 2017 Published by Elsevier Ltd.

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**Corresponding author**: N. Cassir, URMITE, CNRS (UMR 7278), IRD (198), INSERM (U1095), AMU (UM63), Faculté de Médecine, Aix-Marseille Université, 27 Boulevard Jean Moulin, 13385 Marseille Cedex 5, France

E-mail: cassirnadim@gmail.com

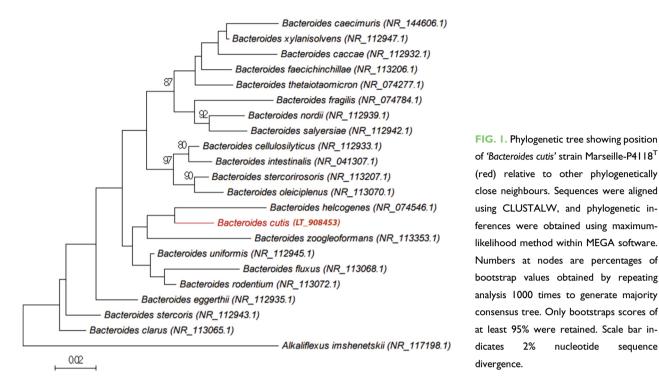
In April 2017, as part of the culturomics study to assess the microbial diversity of the human skin [1,2], a new bacterial species was isolated from a skin sample of a 75-year-old man hospitalized in the Nord hospital, Marseille, France. Although the patient was dermatologically asymptomatic, we isolated from his neck a bacterial strain that could not be identified by our systematic matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS) screening on a MicroFlex spectrometer (Bruker Daltonics, Bremen, Germany) [3]. The study was approved by the Institut Fédératif de Recherche 48 (agreement number 09-022, Marseille, France), and the patient's consent was obtained.

Strain Marseille-P4118<sup>T</sup> was first isolated on 5% sheep's blood–enriched Columbia agar (bioMérieux, Marcy l'Etoile, France) at 37°C in anaerobic atmosphere (AnaeroGen Compact; Oxoid, Thermo Scientific, Dardilly, France) after a 3-day preincubation in an anaerobic bottle of blood culture (Becton Dickinson, Le Pont-de-Claix, France) containing 5% sheep's blood supplemented with 5% rumen fluid previously filter sterilized through a 0.2  $\mu$ m pore filter (Thermo Fisher Scientific, Villebon sur Yvette, France). Colonies were

colourless, with a diameter ranging from 0.5 to 1.5 mm. Bacterial cells were Gram-negative, motile and rod-shaped bacilli with 0.6  $\mu m$  wide by 2.8  $\mu m$  long.

The strain Marseille-P4118<sup>T</sup> grew after 48 hours of incubation under anaerobic conditions at a temperature ranging from 28°C to 40°C, but optimally at 37°C. Growth was impossible in microaerophilic (campyGEN; Oxoid) conditions. The bacterial cells tolerated a pH of 6.5 to 7, and a NaCl concentration less than 50 mg/L. After 20 minutes of thermal shock at 80°C, this bacterium did not grow at 37°C on Columbia agar enriched with 5% sheep's blood, confirming the spore search result by microscopy, which was negative. The catalase and oxidase tests for the strain Marseille-P4118<sup>T</sup> were negative.

After three failed identifications by MALDI-TOF MS screening on a MicroFlex spectrometer (Bruker Daltonics) [3], the 16S rRNA gene was sequenced using universal primers FDI and RP2 (Eurogentec, Angers, France) as previously described [4] and a 3130-XL sequencer (Applied Biosciences, Saint Aubin, France). Strain Marseille-P4118<sup>T</sup> exhibited a 97.4% sequence identity with the type strain *Bacteroides uniformis*, the phylogenetically closest species with standing in nomenclature (Fig. 1). Therefore, strain Marseille-P4118 was classified as a member of the genus *Bacteroides* in the order of *Bacteriodales* within the *Bacteriodetes* phylum. Because the sequence identity with the phylogenetically closest validated species was <98.7%, which is the threshold recommended to define a species according to the nomenclature [5,6], we suggest the creation of a new species named 'Bacteroides cutis' sp. nov. strain Marseille-P4118<sup>T</sup>



(ku.tis, L. fem., from cutis, 'skin,' the site from which strain Marseille-P4118<sup>T</sup> was first isolated).

#### **MALDI-TOF MS** spectrum

The MALDI-TOF MS spectrum of 'Bacteroides cutis' Marseille-P4118<sup>T</sup> is available online (http://www.mediterraneeinfection. com/).

#### Nucleotide sequence accession number

The 16S rRNA gene sequence was deposited in GenBank under accession number LT908453.

#### Deposit in a culture collection

Strain Marseille-P4118<sup>T</sup> was deposited in the Collection de Souches de l'Unité des Rickettsies (CSUR) under number P4118.

#### Acknowledgement

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**Conflict of interest** 

None declared.

#### References

[1] Lagier JC, Khelaifia S, Alou MT, Ndongo S, Dione N, Hugon P, et al. Culture of previously uncultured members of the human gut microbiota by culturomics. Nat Microbiol 2016;1:16203.

2%

nucleotide

sequence

- [2] Cassir N, Papazian L, Fournier PE, Raoult D, La Scola B. Insights into bacterial colonization of intensive care patients' skin: the effect of chlorhexidine daily bathing. Eur J Clin Microbiol Infect Dis 2015;34: 999-1004.
- [3] 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 ionization-time of flight mass spectrometry. J Clin Microbiol 2013;51:2182-94.
- [4] 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.
- [5] Stackebrandt E, Ebers J. Taxonomic parameters revisited: tarnished gold standards. Microbiol Today 2006;33:152.
- [6] Ramasamy D, Mishra AK, Lagier JC, Padhmanabhan R, Rossi M, Sentausa E, et al. A polyphasic strategy incorporating genomic data for the taxonomic description of novel bacterial species. Int J Syst Evol Microbiol 2014;64:384-91.

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