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Genome sequence and description of Alterileibacterium massiliense gen. nov., sp. nov., a

new bacterium isolated from human ileum of a patient with Crohn disease

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Running title: Alterileibacterium massiliense gen. nov., sp. nov.

Keywords: Culturomics, taxono-genomics, Alterileibacterium massiliense

1 Abstract:

- 2 *Alterileibacterium massiliense* gen. nov. sp. nov. strain Marseille-P3115^T (= CSURP-3115;
- 3 DSM 103486), formerly proposed as *Ileibacterium massiliense*, is a new genus isolated from
- 4 human ileum of a patient with Crohn disease.

6 Introduction

Alterileibacterium massiliense was isolated using culturomics approach that is based
on the use of a large culture conditions panel in order to describe the microbial composition of
a sample by high-throughput culture [1-4]. A taxono-genomics approach including matrixassisted laser desorption-ionization time-of-flight mass spectrometry (MALDI-TOF MS),
phylogenetic analysis, main phenotypic description and genome sequencing, was used to
describe it [5-6].

13 Isolation and growth conditions

14 In April 2016, an unidentified bacterial strain was isolated from the human ileum of a patient with Crohn disease and provisionally named *Ileibacterium massiliense* [7]. Tentative 15 16 identification was done by MALDI-TOF MS on a Microflex LT spectrometer (Bruker 17 Daltonics, Bremen, Germany) as previously described [8]. The obtained spectra (Figure 1) 18 was imported into MALDI Biotyper 3.0 software (Bruker Daltonics) and analyzed against the 19 main spectra of the bacteria included in two databases (Bruker Daltonics and constantly 20 updated MEPHI databases (http://www.mediterranee-21 infection.com/article.php?larub=280&titre=urms-database). This strain was cultured routinely 22 on columbia sheep blood agar (Biomerieux, Marcy l'Etoile, France) at 37°C under anaerobic 23 conditions.

24 Strain identification

To identify this bacterium, the 16S rRNA gene was amplified using the primer pair
fD1 and rP2 (Eurogentec, Angers, France) and sequenced using the Big Dye® Terminator
v1.1 Cycle Sequencing Kit and 3500xLGenetic Analyzer capillary sequencer (Thermofisher,
Saint-Aubin, France) as previously described [9]. The 16S rRNA nucleotide sequence was
assembled and corrected using CodonCode Aligner software (http://www.codoncode.com).

30	The 16s rDNA gene sequence strain Marseille-P3115 exhibited a 90.7 % sequence similarity
31	with <i>Mogibacterium neglectum</i> ATCC700924 ^T (GenBank accession no. AB037875) the
32	phylogenetically closest species with standing in nomenclature (Figure 3). We consequently
33	classify this strain as a member of a new species within the new genus Alterileibacterium,
34	family Clostridiales XIII. Incertae Sedis, phylum Firmicutes.
35	

36 **Phenotypic characteristics (Table 1)**

Microcolonies are white and circular with a mean diameter of 0.08 mm. Bacterial cells of this gram-positive bacterium are easily discolored and appear rather as Gram-negative bacilli that differs with their neighbor's genus *Eubacterium* and *Mogibacterium*. They present a mean diameter of 0.3 μ m and a mean length of 1 μ m. (Figure 2) Strain Marseille-P3115 showed catalase-negative and oxidase-negative activities.

42 Genomics

Genomic DNA of the bacterium was sequenced using the MiSeq Technology (Illumina Inc, San Diego, CA, USA) with the mate pair strategy as previously described [10]. Total information of 5.1 Gb was obtained from a 544K/mm² cluster density with a cluster passing quality control filters of 96.8 % (10.139.000 passing filter paired reads). Within this run, the index representation for *A. massiliense* was determined to 4.43%. The 449.618 paired reads were trimmed using the Trimmomatic software [11], GapCloser [12] was used to reduce gaps, then assembled with the Spades software [13] in 2 scaffolds.

50 The genome of strain Marseille-P3115 is 1.450.823 bp long with a 35.9 mol% G+C 51 content. The degree of genomic similarity of *A. massiliense* strain Marseille-P3115 with 52 closely related species was estimated using the OrthoANI software [14]. Values among 53 closely related species (Figure 4) ranged from 63.39% between *Eubacterium brachy* ATCC

54	33089 ^T (GenBank: AXUD01000001.1) and <i>Eubacterium pyruvativorans</i> ATCC BAA-574 ^T
55	(GenBank FNBF01000001.1) to 83.43% between <i>Mogibacterium diversum</i> ATCC700923 ^T
56	(RefSeq NZ_CP027228.1) and <i>Mogibacterium pumilium</i> ATCC700696 ^T (RefSeq
57	NZ_CP016199.1). When the isolate was compared to these closely species, values ranged
58	from 64.02% with Eubacterium pyruvativorans ATCCBAA574 ^T (GenBank
59	FNBF01000001.1) to 68.04% with <i>Mogibacterium timidum</i> ATCC33093 ^T (GenBank
60	JALU01000001.1).

61

62 **Conclusion**

Strain Marseille-P3115^T exhibiting a 16S rRNA sequence divergence > 5 % with its
phylogenetically closest species with standing in nomenclature, is consequently proposed as
the type strain of the new genus *Alterileibacterium massiliense* gen. nov., sp. nov.
(Alter.il.ei.bac.te[']ri.um, Gr. adj. Alter 'other', Gr. n. ilei, 'ileum'; Gr. n. bakterion,

67 'bacterium'; N.L. neut. n. Ileibacterium, 'bacterium isolated from the human ileum sample

⁶⁸ 'mas.si.li. en'se, L. neut. adj., massiliense for Massilia, the Latin name of Marseille, where the

69 strain was first isolated). This strain was provisionally named *Ileibacterium massiliense* [7].

70 However, in the same time the genus name *Ileibacterium* gen. nov. was used to name an

71 isolate of the family *Erysipelotrichaceae* [15]. This is the reason we modified the name of our

72 isolate from *Ileibacterium massiliense* gen. nov. sp. nov. to *Alterileibacterium massiliense*

73 gen. nov. sp. nov.

Nucleotide sequence accession number. The 16S rRNA gene and genome sequences were
 deposited in GenBank under accession number LT598557 and FNWE00000000, respectively.
 Deposit in culture collections. Strain Marseille-P3115^T was deposited in two different strain
 collections under number CSURP3115 and DSM103486.

7	8
	U

79	Conflict of interest:
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80 None to declare.

81 Acknowledgements

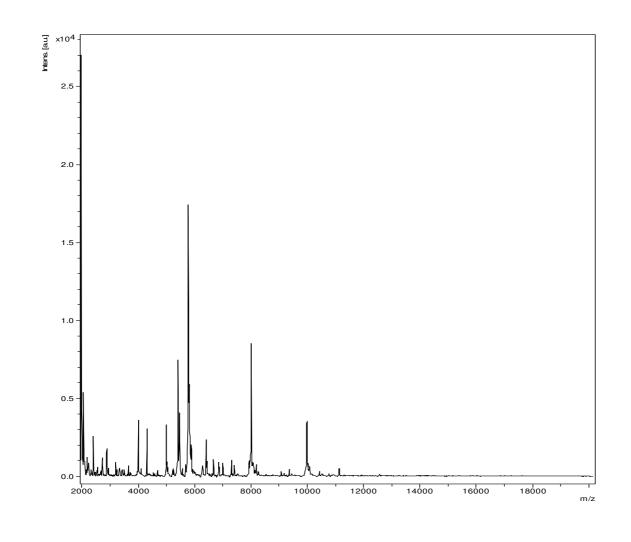
- 82 The authors are indebted to Catherine Robert for sequencing the genome, Aurelia Caputo for
- 83 submitting the genome sequence to GenBank and the platform of electron microscopy of IHU
- 84 for the electron micrographs.
- 85

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- 90 FEDER PRIMI

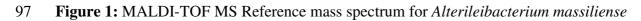
- **Table 1**: Description of *Alterileibacterium massiliense* according to the digitalized protologue
- 92 under the number TA00884 on the www.imedea.uib.es/dprotologue website.

	T + 0000 4
Taxonumber	TA00884
Date of the entry	2019-01-29
Version	Draft
Species name	Alterileibacterium massiliense
Genus name	Alterileibacterium
Specific epithet	massiliense
Species status	gen. nov; sp. nov.
Species etymology	Alterileibacterium massiliense gen. nov., sp. nov. (Alter.il.ei.bac.te'ri.um, Gr. adj. Alter 'other', Gr. n. ilei, 'ileum'; Gr. n. bakterion, 'bacterium'; N.L. neut. n. Ileibacterium, 'bacterium isolated from the human ileum sample';'mas.si.li. en'se, L. neut. adj., massiliense for Massilia, the Latin name of Marseille, where the strain was first isolated.)
Submitter	BOXBERGER Manon
E-mail of the submitter	manon.boxberger@hotmail.fr
Designation of the type strain	Marseille-P3115
Strain collection numbers	CSURP3115=DSM103486
16S rRNA gene accession number	LT598557
Genome accession number [EMBL]	FNWE0000000
Genome status	Draft
Genome size	1,450,823 bp
GC mol %	35.9
Data on the origin of the sample from whi	ich the strain had been isolated
Country of origin	France
Region of origin	Marseille
Date of isolation	2016-01-01
Source of isolation	Human ileum
Sampling date	2019-01-01
Growth medium, incubation conditions	Columbia agar supplemented with 5% sheep blood, 37°C for 48h
[Temperature, pH, and further information] used for standard cultivation	of incubation
Gram stain	Negative
Cell shape	Bacilli
Cell size (mean length; mean diameter)	1 ; 0.3 (μm)
Colony morphology	White, circular
Motility	Non motile
Sporulation	No sporulation
Temperature range	37°C
Temperature optimum	37°C
Lowest pH for growth	7
Highest pH for growth	7,5
Relationship to O ₂	Strictly anaerobe
O ₂ conditions for strain testing	Aerobiosis, Anaerobiosis, Microaerophilic
Oxidase	Negative
Catalase	Negative



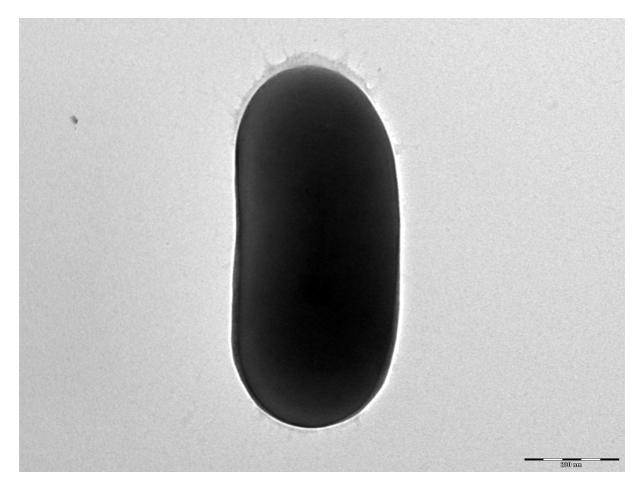






98 Marseille-P3115. Spectra from 12 individual colonies were compared and a reference

99 spectrum was generated.





109 Figure 2. Micrograph electron microscopy of strain *Alterileibacterium massiliense* gen. nov.,

sp. nov. A colony was collected from agar and fixed with 2.5 % glutaraldehyde in 0.1M

111 cacodylate buffer for at least 1h at 4°C. A drop of cell suspension was deposited for

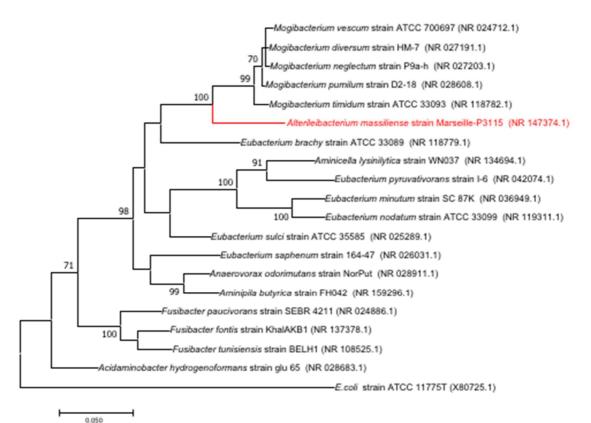
approximately 5 minutes on glow-discharged formvar carbon film with 400 mesh nickel grids

113 (FCF400-Ni, EMS). The grids were dried on blotting paper and the cells were negatively

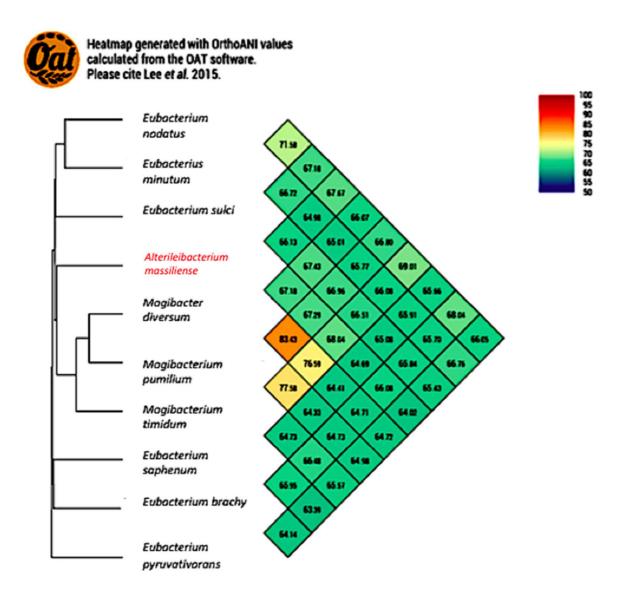
114 stained for 10 seconds with 1% ammonium molybdate solution in filtered water at RT.

115 Electron micrographs were acquired with a Morgagni 268D (Philips) transmission electron

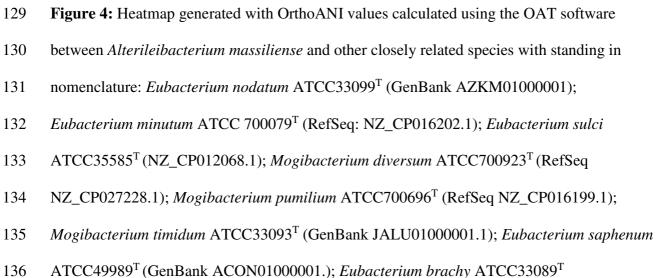
116 microscope operated at 80 keV. Scale is shown of figures.



120 Figure 3: Phylogenetic tree showing the position of Alterileibacterium massiliense strain 121 Marseille-P3115 relative to other phylogenetically-close neighbors. The respective GenBank 122 accession numbers for 16S rRNA genes are indicated in parenthesis. Sequences were aligned 123 using Muscle v7.0.26 with default parameters and phylogenetic inferences were obtained 124 using the maximum likelihood method within MEGA 7 software. Numbers at the nodes are 125 percentages of bootstrap values obtained by repeating the analysis 1000 times to generate a 126 majority consensus tree. Only bootstrap values > 70 % were retained. The scale bar indicates a 5 % nucleotide sequence divergence. 127







- 137 (AXUD01000001.1); *Eubacterium pyruvativorans* ATCCBAA574^T (GenBank
- 138 FNBF01000001.1)

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