TAXONOGENOMICS: GENOME OF A NEW ORGANISM

Detailed description of *Senegalia massiliensis* strain SIT17^T^, a bacterium isolated from the human gut

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**Abstract**

Strain SIT17^T^ was isolated from the stool of a healthy 13-month-old Senegalese boy. It is a Gram-positive, anaerobic, rod-shaped, non-spore-forming and mobile bacterium. It exhibited 92.74% 16S rRNA gene sequence similarity with the *Brassicibacter thermophilus* strain Cel2f, the phylogenetically most closely related species. Its genome is about 2.87 Mb long with 27.39 mol% G + C content. We provide more details of *Senegalia massiliensis* strain SIT17^T^ (= CSURP2130 = DSM 103071), the creation of which was previously announced.

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**Introduction**

Recently, the culturomics concept developed in our laboratory has allowed us to change the paradigm of the human gut microbiota [1]. Indeed, by this method, >50% of the microorganisms present in the human gut microbiota are known [2]. To improve culture and bacterial identification, culturomics is associated with a new process named taxonogenomics to provide exhaustive information and to better characterize bacterial species [3-5]. Combining phenotypic characteristics and genomic analysis and comparison, this polyphasic approach exceeds the limits of conventional methods long used for the description of new species [6-8].

Here, we present the classification and features of *Senegalia massiliensis* strain SIT17^T^, including a description of the complete genome sequencing and annotation.

**Isolation and growth conditions**

Strain SIT17^T^ was first isolated in 2015 from the stool of a healthy 13-month-old Senegalese boy [9]. The sample was collected in Senegal and was then frozen at –80°C. Subsequently, it was transported in dry ice to Marseille, where the bacterial culture was started. The initial growth of bacterial cells was obtained on Columbia agar with 5% sheep’s blood after 2 days of anaerobic incubation at 37°C. The identification of strain SIT17^T^ using matrix assisted laser desorption/ionization time-of-flight mass spectrometry was unsuccessful. The process was performed on a Microflex LT spectrometer (Bruker, Daltonics, Bremen, Germany) as previously described [10-11]. The spectra obtained were imported and analysed using the BIOTYPER 3.0 software against the Bruker database, which is permanently improved with the local MEPHI database (Fig. 1).

**Strain identification and phylogenetic analysis**

In order to identify the strain SIT17^T^, the 16S rRNA gene was amplified using the fD1 and rP2 primer pair (Eurogentec, Angers, France) and sequenced using the Big Dye® Terminator v1.1 Cycle Sequencing Kit and 3500xL Genetic Analyzer capillary sequencer (Thermo Fisher, Saint-Aubin, France), as previously reported [12]. The 16S rRNA nucleotide sequences were assembled and corrected using CodonCode Aligner software...
FIG. 1. Reference mass spectrum from *Senegalia massiliensis* strain SIT17T.

FIG. 2. Phylogenetic tree highlighting the position of *Senegalia massiliensis* strain SIT17T relative to other close species. Sequences were aligned using CLUSTALW, with default parameters, and phylogenetic inferences were obtained using the maximum likelihood method within the MEGA7 software. Numbers at the nodes are percentages of bootstrap values obtained by repeating the analysis 500 times to generate a majority consensus tree. The scale bar indicates a 2% nucleotide sequence divergence.
The PCR-amplified genes coding for 16S rRNA of *Senegalia massiliensis* yielded 92.74% similarity level with *Brassicibacter thermophilus* strain Cel2f (GenBank accession no: NR137216) [13], the phylogenetically closest species with standing in nomenclature (Fig. 2). This value was lower than 95%, which is the recommended threshold for delineating a new bacterial genus based on 16S rRNA gene sequence without DNA–DNA hybridization [14-15]. Classification and general features are summarized in Table 1.

### Phenotypic and biochemical characteristics

Colonies of the strain SIT17\(^T\) were grey and translucent with a size of 0.5–1 mm on Columbia agar with 5% sheep’s blood. Growth was observed from 28°C to 45°C, with optimal growth at 37°C, and colonies were obtained after 48 hours of culture. Bacterial cells were Gram-positive, rod-shaped and motile, but non-spor-forming (Fig. 3a). Observed under electronic microscopy, the cells presented a mean diameter of 0.4 μm and a mean length of 3.2 μm (Fig. 3b). The bacterium was catalase positive but had no oxidase activity. *Senegalia massiliensis* is able to grow in an environment with a pH ranging from 6 to 8.5, with an optimal value of 7. Strain SIT17\(^T\) is an anaerobic bacterium that can grow in a microaerophilic atmosphere. On the other hand, no growth was observed under aerobic conditions. The biochemical and phenotypic features of strain SIT17\(^T\) were compared with those of other close representative strains in the *Clostridiaceae* family (Table 2).

Using API ZYM strips (bioMérieux, Marcy l’Étoile, France), positive reactions were observed for esterase, esterase lipase, alkaline phosphatase, α-chymotrypsin, acid phosphatase, naphthol-AS-BI-phosphohydrolase and β-galactosidase. However, we noted that the enzymatic activities for lipase, leucine arylamidase, cystine arylamidase, trypsin, α-galactosidase, β-glucuronidase, β-glucosidase, N-acetyl-β-glucosaminidase, α-mannosidase and α-fucosidase, were negative. Using API 50 CH, positive reactions were observed for glycerol, D-ribose, L-xyllose, D-galactose, D-glucose, D-fructose, D-mannose, L-rhamnose, inositol, D-mannitol, D-sorbitol, methyl D-D-glucopyranoside, N-acetylglucosamine, amygdalin, arbutin, salicin, D-cellulbiose, D-maltose, D-lactose, D-sucrose, D-trehalose, D-melezitose, D-raffinose, D-turanose, D-xyllose, D-fucose, D-fucitol, D-arabitol, potassium gluconate and starch. However, there was no metabolism for the following carbohydrates: erythritol, L-arabinose, D-adonitol, methyl β-D-xylpyranoside, methyl α-D-mannopyranoside, D-arabinose, inulin

### Table 1. Classification and general features of *Senegalia massiliensis* strain SIT17\(^T\)

| Property                        | Terms                                                                 |
|---------------------------------|----------------------------------------------------------------------|
| Current classification          | Domain: *Bacteria*  
Phylum: *Firmicutes*  
Class: *Clostridia*  
Order: *Clostridiales*  
Family: *Clostridiaceae*  
Genus: *Senegalia*  
Species: *Senegalia massiliensis*  
Type: strain SIT17\(^T\) |
| Gram stain                      | Positive                                                               |
| Cell shape                      | Rod                                                                 |
| Motility                        | Motile                                                                |
| Sporulation                     | Not sporulating                                                       |
| Temperature range               | 28°C–45°C                                                             |
| Optimum temperature             | 37°C                                                                  |
| pH range (optimum)              | 7                                                                     |
| Oxygen requirement              | Anaerobic                                                             |
| Carbon source                   | Unknown                                                               |
| Habitat                         | Human gut                                                             |
| Basic relationship              | Free-living                                                           |
| Pathogenicity                   | Unknown                                                               |

![FIG. 3. The morphology of bacterial cells of strain SIT17\(^T\). (a) Gram staining of *Senegalia massiliensis* strain SIT17\(^T\). (b) Transmission electron microscopy of *Senegalia massiliensis* strain SIT17\(^T\) using Tecnai G20 electron microscope (FEI Company). The scale bar represents 500 nm.](#)
TABLE 2. Differential characteristics of Senegalia massiliensis strain SIT17T (data from this study) compared to other close bacteria

| Properties                   | Senegalia massiliensis | Clostridiisalibacter paucivorans | Alkaliphilus oremlandii | Alkaliphilus transvaalenis | Proteiniborus ethanoligenes | Sporosallibacter fouaurense |
|------------------------------|------------------------|----------------------------------|-------------------------|---------------------------|-------------------------------|-----------------------------|
| Cell diameter (μm)           | 0.3–0.5                | 0.5                              | 0.5                     | 0.4–0.7                   | 0.5–0.6                       | 0.5                         |
| Oxygen requirement           | Anaerobic              | Anaerobic                        | Anaerobic               | Anaerobic                 | Anaerobic                     | Anaerobic                   |
| Shape                        | bacilli                | bacilli                          | bacilli                 | bacilli                   | bacilli                        | bacilli                     |
| Gram stain                   | +                      | +                                | +                       | +                         | +                             | +                           |
| Motility                     | +                      | +                                | +                       | +                         | +                             | +                           |
| Motility                     | +                      | +                                | +                       | +                         | +                             | +                           |
| Motility                     | +                      | +                                | +                       | +                         | +                             | +                           |
| Production of:               |                        |                                  |                         |                           |                               |                             |
| Alkalinephosphatase          | +                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| Catalase                     | +                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| Oxidase                      | —                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| Nitrate reductase            | —                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| Urease                       | —                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| β-galactosidase              | +                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| N-acetylglucosamine          | —                      | NA                               | NA                      | NA                        | NA                            | NA                          |
| Acid from:                   |                        |                                  |                         |                           |                               |                             |
| L-arabinose                  | —                      | +                                | NA                      | NA                        |                               |                             |
| Ribose                      | +                      | NA                               | NA                      | NA                        |                               |                             |
| Mannose                     | +                      | —                                | NA                      | NA                        |                               |                             |
| Manitol                      | +                      | NA                               | NA                      | NA                        |                               |                             |
| D-glucose                   | —                      | NA                               | NA                      | NA                        |                               |                             |
| D-fructose                  | +                      | —                                | NA                      | NA                        |                               |                             |
| L-malate                     | +                      | NA                               | NA                      | NA                        |                               |                             |
| L-lactate                    | +                      | NA                               | NA                      | NA                        |                               |                             |
| G + C content (%)           | 27.4                   | 33.0                             | 36.1                    | 36.4                      | 38.0                          | 37.7                        |
| Habitat                     | Human colon            | Wastewater                       | Environment             | Environment               | Environment                   | Soil                         |

**, negative reaction; +, positive reaction; NA, not available data.

and glycogen. Cellular fatty acid methyl esters analysis of the strain SIT17T was carried out by operating gas chromatography/mass spectrometry as previously described [16-17]. The result showed that hexadecanoic acid (32.6%), 9-octadecenoic acid (21.6%) and 13-methyl-tetradecanoic acid (11.9%), are the most abundant fatty acids. Other saturated and unsaturated fatty acids are also found (Table 3).

Genomic properties and comparison
The genome of strain SIT17T is 2,866,883 bp long with 27.39 mol% G + C content and it contains 2933 coding genes (Fig. 4). It is composed of 22 contigs. By comparing it with related species, its genome (2.87 Mbp) is smaller than those of Alkaliphilus oremlandii strain OhiLAsm, Proteiniborus ethanoligenes strain DSM 21650, Clostridiisalibacter paucivorans strain DSM 22131, Alkaliphilus transvaalenis strain ATCC 700919, Paramaledivibacter caminithermalis strain DSM 15212, Alkaliphilus peptidifermentans strain DSM 18978 and Alkaliphilus metalredi gens strain L21-TH-D2 (3.12, 3.16, 3.24, 4.02, 4.05, 4.45 and 4.93 Mbp, respectively), but larger than the genome of Caldisalibacter kiritimatiensis (2.79 Mbp). The G + C content of strain SIT17T (27.39 mol%) is smaller than those of A. oremlandii, Proteiniborus ethanoligenes, Clostridiisalibacter paucivorans, A. transvaalenis, Paramaledivibacter caminithermalis, A. peptidifermentans, A. metalredi gens and Caldisalibacter kiritimatiensis (36.3, 32.6, 31.4, 34.0, 30.5, 34.1, 36.8 and 30.1 mol%, respectively). The gene content of strain SIT17T (293 genes) is larger than those of A. oremlandii (n = 2898), Caldisalibacter kiritimatiensis (n = 2557) and Proteiniborus ethanoligenes (n = 2846), but smaller than Clostridiisalibacter paucivorans (n = 3014), A. peptidifermentans (n = 4072), A. metalredi gens (n = 4641), A. transvaalenis (n = 3640) and Paramaledivibacter caminithermalis (n = 3543). Distribution of functional classes of predicted genes according to the clusters of orthologous groups is shown in Table 3.

TABLE 3. Cellular fatty acid profiles (%) of Senegalia massiliensis strain SIT17T compared with other species

| Fatty acids     | Names                      | 1 | 2 | 3 | 4 | 5 |
|-----------------|----------------------------|---|---|---|---|---|
| 12:0            | Dodecanoic acid            | 1.2 | 1.2 | 1.2 | 1.2 | 1.2 |
| 13:0            | Tridecanoic acid           | TR | 2.3 | 2.3 | 2.3 | 2.3 |
| 14:0            | Tetradecanoic acid         | 9.2 | 14.3 | 15.8 | 21.6 | 21.6 |
| 15:0 iso        | 12-methyl-tetradecanoic acid | 1.0 | 1.3 | 2.8 | 3.9 | 3.9 |
| 15:0 anteiso    | 13-methyl-tetradecanoic acid | 11.9 | 6.6 | 51.6 | 43.0 | 41 |
| 16:0            | Hexadecanoic acid          | 32.6 | 7.6 | 3.9 | 25.4 | 1.2 |
| 16:1n5         | 11-Hexadecenoic acid       | TR | 1.9 | 6.1 | 6.1 | 6.1 |
| 17:0            | Heptadecanoic acid         | TR | 0.6 | 0.6 | 0.6 | 0.6 |
| 17:1 anteiso    | 10-Heptadecenoic acid      | TR | 9.3 | 12.2 | 9.49 | 9.49 |
| 18:0            | Octadecanoic acid          | 4.7 | 7.2 | 12.0 | 1.3 | 1.3 |
| 18:1n7         | 11-Octadecenoic acid       | 1.7 | 2.0 | 2.0 | 2.0 | 2.0 |
| 18:1n9         | 9-Octadecenoic acid        | 21.6 | 1.1 | 11.2 | 11.2 | 11.2 |

1. Senegalia massiliensis strain SIT17T; 2. Clostridiisalibacter paucivorans strain 37H5600 [19]; 3. Alkaliphilus oremlandii strain SAGM1T [20]; 4. Proteiniborus ethanoligenes strain GW1 [21]; 5. Sporosallibacter fouaurense strain SOL397T [11].

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FIG. 4. Graphical circular map of the chromosome. From outside to the centre: Contigs (red), cluster of orthologous groups (COG) category of genes on the forward strand (three circles), genes on forward strand (blue circle), genes on the reverse strand (red circle), COG category on the reverse strand (three circles), G + C content.
groups is reported in Table 4. Results from pairwise genome comparison obtained from analysis of the digital DNA–DNA hybridization using GGDC software [18] are shown in Table 5. ORTHOANI values among the closely related species ranged from 64.37%, between A. metalliredigens and Clostridii salibacter paucivorans, to 70.05% between Caldisalini bacter kiritimatiensis and S. massiliensis. When S. massiliensis was compared with these closely related species, values ranged from 65.93% with A. metalliredigens to 70.05% with Caldisalini bacter kiritimatiensis (Fig. 5).

**TABLE 4. Distribution of functional classes of predicted genes according to the clusters of orthologous groups of proteins of Senegalia massiliensis strain SIT17**

| Code | Value | Description |
|------|-------|-------------|
| [J]  | 244   | Translation, ribosomal structure and biogenesis |
| [A]  | 0     | RNA processing and modification |
| [C]  | 225   | Transcription |
| [D]  | 114   | Replication, recombination and repair |
| [E]  | 1     | Chromatin structure and dynamics |
| [F]  | 52    | Cell-cycle control, cell division, chromosome partitioning |
| [G]  | 0     | Nuclear structure |
| [H]  | 80    | Defence mechanisms |
| [I]  | 181   | Signal transduction mechanisms |
| [J]  | 148   | Cell wall/membrane/envelope biogenesis |
| [K]  | 71    | Cell motility |
| [L]  | 0     | Cytoskeleton |
| [M]  | 11    | Extracellular structures |
| [N]  | 28    | Intracellular trafficking, secretion and vesicular transport |
| [O]  | 116   | Post-translational modification, protein turnover, chaperones |
| [P]  | 26    | Nucleotide transport and metabolism |
| [Q]  | 173   | Energy production and conversion |
| [R]  | 144   | Carbohydrate transport and metabolism |
| [S]  | 202   | Amino acid transport and metabolism |
| [T]  | 92    | Nucleotide transport and metabolism |
| [U]  | 124   | Coenzyme transport and metabolism |
| [V]  | 89    | Lipid transport and metabolism |
| [W]  | 145   | Inorganic ion transport and metabolism |
| [X]  | 34    | Secondary metabolites biosynthesis, transport and catabolism |
| [Y]  | 563   | General function prediction only |
| [Z]  | 255   | Function unknown |
| [Q]  | 203   | Hypothetical protein |

**TABLE 5. Pairwise comparison of Senegalia massiliensis strain SIT17** with other species using GGDC formulaa

| Species                                | Senegalia massiliensis | Alkaliphilus metalliredigens | Alkaliphilus oremlandii | Alkaliphilus transvaalenensis | Alkaliphilus peptidifermentans | Proteiniborus ethanoligenes | Clostridii salibacter paucivorans | Parmale divibacter caminithermales | Caldisalini bacter kiritimatiensis |
|----------------------------------------|------------------------|------------------------------|------------------------|------------------------------|--------------------------------|-------------------------------|----------------------------------|-----------------------------------|----------------------------------|
| Caldisalini bacter kiritimatiensis     | 19.7 ± 2.4             | 37.2 ± 5                     | 34 ± 4.9               | 17.5 ± 4.4                   | 18 ± 4.5                       | 18.9 ± 4.6                      | 16.9 ± 4.4                       | 23.6 ± 4.7                         | 100%                             |
| Parmale divibacter caminithermales     | 15.2 ± 4.3             | 28.4 ± 4.9                   | 26.4 ± 4.8             | 16 ± 4.3                     | 19.5 ± 4.6                     | 19 ± 4.5                       | 23 ± 4.7                         | 100%                              |
| Clostridii salibacter paucivorans      | 17.5 ± 4.5             | 25 ± 4.8                     | 16.7 ± 4.4             | 31 ± 4.9                     | 17.8 ± 5.4                     | 18.6 ± 5.5                     | 100%                             |
| Proteiniborus ethanoligenes            | 17.3 ± 4.1             | 19.9 ± 4.6                   | 19.8 ± 4.6             | 18.7 ± 4.6                   | 16.2 ± 4.3                     | 100%                           |
| Alkaliphilus peptidifermentans         | 17.2 ± 4.4             | 25.5 ± 5.2                   | 24.8 ± 4.8             | 19.9 ± 4.7                   | 100%                           |
| Alkaliphilus transvaalenensis          | 27.4 ± 4.8             | 23.5 ± 4.8                   | 22.2 ± 4.7             | 100%                         |
| Alkaliphilus oremlandii                | 29.9 ± 4.9             | 26.8 ± 4.9                   | 100%                   |
| Alkaliphilus arminandi                 | 33.9 ± 4.9             | 100%                         |

*aGGDC formula 2: (Identities/high-scoring segment pairs (HSPs)). The confidence intervals indicate the inherent uncertainty in estimating DNA–DNA hybridization values from intergenomic distances based on models derived from empirical test data sets (which are always limited in size). These results are in accordance with the 16S rRNA and phylogenomic analyses as well as the GGDC results.*
Conclusion

On the basis of phenotypic, phylogenetic and genomic analyses, we formally propose the creation of *Senegalia massiliensis* gen. nov. sp. nov., that contains the strain SIT17\(^T\). Hence, the combination of culturomics and taxonogenomics has contributed to a better knowledge of the associated human microorganisms and to better understanding of physiological functioning in health and disease.

**Description of Senegalia gen. nov**

*Senegalia* (Se.ne.ga.lia. L. gen. n. Senegalia, the Latin name of Senegal, where the stool specimen was collected). Cells are Gram-positive, non spore-forming, motile and anaerobic bacilli. The type species is *Senegalia massiliensis* sp. nov.

**Description of Senegalia massiliensis sp. nov**

*Senegalia massiliensis* gen. nov., sp. nov. (mas.si.li.en’sis. L. fem. adj., from massiliensis, of Massilia, from the Latin name of Marseille where the strain was first isolated). It is classified as a member of the family *Clostridiaceae* within the phylum *Firmicutes*. The strain SIT17\(^T\) designed the type strain of *Senegalia massiliensis* gen. nov., sp. nov., and was deposited in CSUR (CSURP2130) and DSMZ (DSM 103071) collections. It is a Gram-positive bacillus, motile, catalase-positive, oxidase-negative and non-spore-forming. Strain SIT17\(^T\) was first isolated from the stool of a healthy 13-month-old Senegalese boy. Its genome is 2 866 883 bp long with 27.39 mol% G + C content and possesses 2933 coding genes. The genome and 16S rRNA sequences of the strain SIT17\(^T\) are both deposited in GenBank under accession numbers UZAQ00000000 and LN881608, respectively.

**Conflict of interest**

The authors declare no conflict of interest.

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Ethics and consent

The child's parents provided signed informed consent and the study was approved by the ethics committee of the Institut Fédératif de Recherche IFR48 under number 09-022.

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