Lactobacillus garii sp. nov., isolated from a fermented cassava product

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Abstract

A novel Gram-positive, catalase negative, rod-shaped strain, FI11369[T], was isolated from *gari*, a traditional West African fermented food derived from cassava. Based on 16S rRNA gene sequence similarity, the closest type strains were *Lactobacillus xiangfangensis* LMG 26013[T] (99.4% similarity), *Lactobacillus plajomi* NBRC 107333[T] (99.1%), *Lactobacillus paraplantarum* DSM 10667[T] (99.1%), *Lactobacillus pentosus* DSM 20314[T] (99.0%), *Lactobacillus plantarum* subsp. *plantarum* ATCC 14917[T] (99.0%), *Lactobacillus modestisalitolerans* NBRC 107235[T] (98.9%), *Lactobacillus plantarum* subsp. *argentoratensis* DSM 16365[T] (98.9%) and *Lactobacillus daowaiensis* NCIMB 15183[T] (98.8%). The genome of strain FI11369[T] was sequenced and the average nucleotide identity (ANI) was compared with its closest relatives. ANI analysis showed that the closest relative, *L. xiangfangensis* DSM 27103[T], had only a 82.4% similarity. The main fatty acids of FI11369[T] were saturated $C_{16:0}$ (18.2%), unsaturated $C_{18:1}\omega9c$ (43.8%) and cyclopropane $C_{19:0}$ cyclo ($\omega10c$ and/or $\omega6$; 22.5%). Based on the genotypic and phenotypic data obtained in this study, a novel *Lactobacillus* species, *Lactobacillus garii* sp. nov., with the type strain FI11369[T] (=NCIMB 15148=DSM 108249), is proposed.

The genus *Lactobacillus*, which includes more than 200 species, is a taxonomically complex group due to the high level of phenotypic and genotypic diversity [1]. Lactobacilli are Gram-positive, mostly non-motile, catalase-negative, non-spore forming and rod-shaped bacteria. Their habitats are nutrient-rich environments such as food, soil, plants, animals and humans [2]. Lactobacilli dominate the microbiota of the vast majority of fermented foods and most studies have focused on their role in food fermentation and prevention of food spoilage [3], as well as their importance in the gut and their applications as probiotics [4].

Gari, a fermented food derived from cassava (*Manihot esculenta*), is widely consumed in West and Central Africa. The steps to obtain *gari* include washing and grating fresh cassava roots, followed by fermenting and dewatering at ambient temperature (ca. 30 °C) for up to 72 h. The fermented pressed cake is disintegrated and roasted into *gari* [5, 6]. In a previous study of the diversity of lactic acid bacteria in *gari*, *Lactobacillus plantarum* was the most frequently isolated species, followed by *Leuconostoc fallax* and *Lactobacillus fermentum* [6].

Strain FI11369[T] was isolated as part of a study to sequence the whole genomes of micro-organisms present in African fermented foods. In this study, a sample of *gari*, produced in the suburb of Pokuase, Accra (Ghana) was collected and preserved at 4 °C until further processing in the UK. The sample of *gari* was homogenized in PBS and 100 µl of a 10^{-5} dilution of the homogenate were plated on MRS (Oxoid) agar medium [7]. Plates were incubated for 48 h at 37 °C and 10 out of 194 colonies were picked based on different morphology and sub-cultured for three rounds until pure cultures were obtained. A pure culture of strain FI11369[T], the only selected colony with irregular edges, was obtained after three rounds of sub-culturing. DNA from strain FI11369[T] was extracted using the cetyltrimethylammonium bromide-based extraction protocol [8] and the genome was sequenced. Libraries were obtained using the Nextera XT DNA library Prep kit (Illumina) according to manufacturer instructions and sequenced for 150 cycles using the Illumina NextSeq platform at the Quadram Institute Bioscience (QIB; Norwich, UK). The 887369 reads generated were quality trimmed with BBDuk (version 38.68) to

Tree scale: 0.01 ⊢————┤

Leuconostoc mesenteroides subsp. *mesenteroides* ATCC 8293[T] (CP000414)

Lactobacillus delbrueckii subsp. *delbrueckii* DSM 20074[T] CP018615)

Lactobacillus fabifermentans DSM 21115[T] (NR 113339)

Lactobacillus herbarum TCF032-E4[T] (NR 145899)

Lactobacillus pingfangensis NCIMB 15187[T] (BJDG00000000)

Lactobacillus nangangensis NCIMB 15186[T] (BJDI00000000)

Lactobacillus daoliensis NCIMB 15181[T] (BJDH00000000)

Lactobacillus daowaiensis NCIMB 15183[T] (BJDJ00000000)

Lactobacillus mudanjiangensis DSM 28402[T] (NR 125561)

Lactobacillus dongliensis NCIMB 15184[T] (BJDK00000000)

Lactobacillus songbeiensis NCIMB 15189[T] (BJDL00000000)

Lactobacillus plantarum subsp. *argentoratensis* DSM 16365[T] (NR 042254)

Lactobacillus plantarum subsp. *plantarum* ATCC 14917[T] (NR 104573)

Lactobacillus pentosus DSM 20314[T] (NR 029133)

Lactobacillus paraplantarum DSM 10667[T] (NR 025447)

Lactobacillus garii FI11369[T] (MK011005)

Lactobacillus xiangfangensis LMG 26013[T] (NR 109000)

Lactobacillus plajomi NBRC 107333[T] (NR 136785)

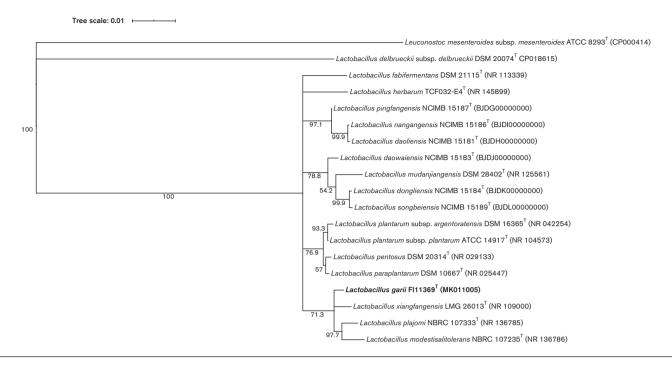*Lactobacillus modestisalitolerans* NBRC 107235[T] (NR 136786)

Fig. 1. Phylogenetic tree based on 16S rRNA gene sequences showing the relationship of strain FI11369[T] and the related *Lactobacillus* species. Tree reconstructed by the maximum-likelihood method. *Leuconostoc mesenteroides* subsp. *mesenteroides* ATCC 8293[T] was used as an outgroup. Bootstrap values are indicated at branch points based on 1000 replications. Bootstrap values below 62% are not shown. Bar, 0.01 substitutions per nucleotide position.

at least a quality of Q21. Cleaned reads were assembled with SPAdes (version 3.11.1) [9] and annotated using PATRIC [10]. The genome was assembled into 158 contigs, with an N50 of 57125 bp. The 16S rRNA gene sequence (accession no. MK011005) was extracted from the genome assembly (accession no. QWZQ00000000; size 2972171 bp). The sequence was compared with all the type strains present in the nucleotide collection database from the National Center for Biotechnology Information (NCBI). Sequence similarity with the closest type strains was calculated using the pairwise nucleotide sequence alignment for taxonomy tool from EzBioCloud [11]. The highest similarity was found with the type strains of *Lactobacillus xiangfangensis* (99.4% similarity), *Lactobacillus plajomi* (99.1%), *Lactobacillus paraplantarum* (99.1%), *Lactobacillus pentosus* (99.0%), *Lactobacillus plantarum* subsp. *plantarum* (99.0%), *Lactobacillus modestisalitolerans* (98.9%), *Lactobacillus plantarum* subsp. *argentoratensis* (98.9%), *Lactobacillus daowaiensis* (98.8%), *Lactobacillus fabifermentans* (98.6%), *Lactobacillus nangangensis* (98.6%), *Lactobacillus daoliensis* (98.5%), *Lactobacillus pingfangensis* (98.5%), *Lactobacillus herbarum* (98.5%), *Lactobacillus mudanjiangensis* (98.0%), *Lactobacillus dongliensis* (98.1%) and *Lactobacillus songbeiensis* (98.1%). All sequences were aligned using CLUSTAL_W version 2.1 [12]. A 16S rRNA gene phylogeny was reconstructed using the maximum-likelihood method with the Jukes–Cantor model [13] incorporated into Geneious version 11.1.3 (Biomatters) (Fig. 1). Branching support was estimated with 1000 bootstrap replicates. In addition,

the 16S rRNA gene was amplified by PCR using primers AMP_F (5′-GAG AGT TTG ATY CTG GCT CAG-3′) and AMP_R (5′-AAG GAG GTG ATC CAR CCG CA-3′) and sequenced by a Sanger sequencing service (Eurofins Genomics Germany GmbH, Germany). The partial 16S rRNA gene sequence (accession no. MN817919) obtained by Sanger sequencing was identical to the 16S rRNA gene sequence extracted from the genome.

To reconstruct a well-supported phylogenomic tree, we extracted the following single-copy marker genes from 18 type strains using AMPHORA2 [14]: *frr*, *infC*, *nusA*, *pgk*, *pyrG*, *rplA*, *rplC*, *rplD*, *rplE*, *rplF*, *rplK*, *rplL*, *rplM*, *rplN*, *rplP*, *rplS*, *rplT*, *rpmA*, *rpoB*, *rpsB*, *rpsC*, *rpsE*, *rpsI*, *rpsJ*, *rpsK*, *rpsM*, *rpsS*, *smpB* and *tsf*. Sequences were aligned using RAxML version 8.2.12 [15]. Single protein alignments were concatenated with the script from phylogenomics-tools (github.com/kbseah/phylogenomics-tools). Positions of the alignment that had more than 75% gaps were removed using Geneious version 11.1.3 (www.geneious.com). The resulting alignment was then used for maximum-likelihood phylogenomic reconstruction with FastTree version 2.1.11 [16] (Fig. 2).

The average nucleotide identity (ANI) of strain FI11369[T] was compared to the type strains that were closest based on our phylogenomic reconstruction, which included: *L. xiangfangensis*, *L. plajomi*, *L. paraplantarum*, *L. pentosus*, *L. plantarum* subsp. *plantarum*, *L. modestisalitolerans*, *L. plantarum* subsp. *argentoratensis*, *L. daowaiensis*, *L.*
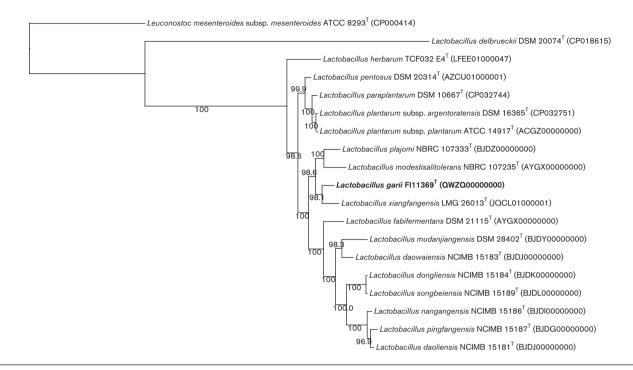
Tree scale: 0.01 ⊢⊣

Leuconostoc mesenteroides subsp. *mesenteroides* ATCC 8293[T] (CP000414)

Lactobacillus delbrueckii DSM 20074[T] (CP018615)

Lactobacillus herbarum TCF032 E4[T] (LFEE01000047)

Lactobacillus pentosus DSM 20314[T] (AZCU01000001)

Lactobacillus paraplantarum DSM 10667[T] (CP032744)

Lactobacillus plantarum subsp. *argentoratensis* DSM 16365[T] (CP032751)

Lactobacillus plantarum subsp. *plantarum* ATCC 14917[T] (ACGZ00000000)

Lactobacillus plajomi NBRC 107333[T] (BJDZ00000000)

Lactobacillus modestisalitolerans NBRC 107235[T] (AYGX00000000)

Lactobacillus garii FI11369[T] (QWZQ00000000)

Lactobacillus xiangfangensis LMG 26013[T] (JQCL01000001)

Lactobacillus fabifermentans DSM 21115[T] (AYGX00000000)

Lactobacillus mudanjiangensis DSM 28402[T] (BJDY00000000)

Lactobacillus daowaiensis NCIMB 15183[T] (BJDJ00000000)

Lactobacillus dongliensis NCIMB 15184[T] (BJDK00000000)

Lactobacillus songbeiensis NCIMB 15189[T] (BJDL00000000)

Lactobacillus nangangensis NCIMB 15186[T] (BJDI00000000)

Lactobacillus pingfangensis NCIMB 15187[T] (BJDG00000000)

Lactobacillus daoliensis NCIMB 15181[T] (BJDJ00000000)

Fig. 2. Phylogenomic tree based on the concatenated *frr*, *infC*, *nusA*, *pgk*, *pyrG*, *rplA*, *rplC*, *rplD*, *rplE*, *rplF*, *rplK*, *rplL*, *rplM*, *rplN*, *rplP*, *rplS*, *rplT*, *rpmA*, *rpoB*, *rpsB*, *rpsC*, *rpsE*, *rpsI*, *rpsJ*, *rpsK*, *rpsM*, *rpsS*, *smpB* and *tsf* gene sequences showing the relationship of strain FI11369[T] and the related *Lactobacillus* species. Tree reconstructed with an alignment spanning 6923 aa using the approximately-maximum-likelihood method implemented in FastTree 2.1.11. *Leuconostoc mesenteroides* subsp. *mesenteroides* ATCC 8293[T] was used as an outgroup. The node labels represent SH-like branching support values. Bar, 0.01 substitutions per aa position.

fabifermentans, *L. nangangensis*, *L. daoliensis*, *L. pingfangensis*, *L. herbarum*, *L. mudanjiangensis*, *L. dongliensis* and *L. songbeiensis*. The ANI and digital DNA–DNA hybridization (dDDH) parameters were calculated using FastANI [17] and formula 2 from the Genome-to-Genome Distance Calculator 2.1 (GGDC 2.1) software [18] respectively using the default settings. The ANI values ranged from 82.4 to 79.7% and the dDDH values ranged from 38.8 to 19.6% (Table 1), which are well below the generally accepted threshold to identify the same species (95–96% for ANI and 70% for dDDH [18–20]). Thus, we confirmed that strain FI11369[T] represents a novel species. The DNA G+C content of FI11369[T] is 48.3 mol%, similar to the 46.6 mol% G+C of its closest relative *L. xiangfangensis* DSM 27103[T] [21].

Morphological and biochemical tests were performed after growth of FI11369[T] at 30 °C in aerobic conditions, unless otherwise stated. Colony morphology was observed after incubation on MRS agar plates. Cell morphology was investigated using scanning electron microscopy (Zeiss Supra 55 VP) as previously described [22]. The cells of FI11369[T] were rod-shaped, 1–2 μm in length and formed short chains or pairs (Fig. 3). Motility and spore formation were determined by phase-contrast microscopy using Olympus microscope CX41 using cells grown on MRS agar and broth for 2 days. Gram-staining was performed using

a commercial Gram-staining kit (Remel, Thermo Fisher Scientific). Catalase activity was determined by treating the cells with 3% H_2O_2. Gas production from glucose was analysed in MRS broth using inverted Durham tubes. Relationship with oxygen was determined on MRS broth or MRS plates under aerobic and anaerobic conditions. The latter were achieved by providing anaerobic gas mix (85% N_2, 5% CO_2 and 10% H_2) in an anaerobic cabinet (Whitley A95 Workstation). Growth at different temperatures (6, 12, 20, 25, 28, 30, 37, 40 and 42 and 55 °C), pH (9, 8.8, 8.4, 8, 7.8, 7.4, 7, 6, 5, 4.0, 3.8, 3.4, 3.2 or 3.0) and salt levels (commercial MRS with 2, 4, 6, 8, 10, 14 or 16% additional NaCl) was measured using OD_{600} after 3 days of incubation. Production of D- and L-lactic acid was analysed using the K-DLATE assay kit (Megazyme). Fermentation products from glucose were analysed by HPLC as previously described [23]. Nitrate and nitrite reduction, indole production, Voges–Proskauer reaction, H_2S production, deamination of arginine, hydrolysis of gelatin and urease production were determined using the API 20E system. Hydrolysis of hippurate was tested using hippurate discs (Sigma-Aldrich). Pyrrolidonylarylamidase production was tested using pyrase strips (Sigma-Aldrich). Bile–aesculin tolerance was tested by growing the isolates on bile aesculin agar (Sigma-Aldrich). Haemolytic activity was checked

Table 1. ANI values (%) and dDDH prediction values (%) between *Lactobacillus garii* sp. nov. and its closely related species

Species	Strain	Accession No.	ANI	dDDH
L. xiangfangensis	LMG 26013	JQCL00000000	82.4	23.8
L. plajomi	NBRC 107333	BJDZ00000000	81.6	22.1
L. plantarum subsp. *argentoratensis*	DSM 16365	CP032751	81.1	21.2
L. paraplantarum	DSM 10667	CP032744	80.8	20.9
L. modestisalitolerans	NBRC 107235	AYGX00000000	80.6	21.2
L. pentosus	DSM 20314	AZCU01000001	80.6	25.1
L. plantarum subsp. *plantarum*	ATCC 14917	ACGZ00000000	80.4	20.6
L. herbarum	TCF032-E4	LFEE01000047	80.1	38.8
L. fabifermentans	DSM 21115	AYGX00000000	80.1	21.2
L. daoliensis	NCIMB 15181	BJDJ00000000	80.0	20.4
L. dongliensis	NCIMB 15184	BJDK00000000	80.0	20.4
L. nangangensis	NCIMB 15186	BJDI00000000	79.9	20.0
L. daowaiensis	NCIMB 15183	BJDJ00000000	79.9	20.4
L. songbeiensis	NCIMB 15189	BJDL00000000	79.7	20.4
L. mudanjiangensis	DSM 28402	BJDY00000000	79.7	19.6
L. pingfangensis	NCIMB 15187	BJDG00000000	79.7	20.2

on Columbia blood agar with 5% sheep blood. Pyruvate utilization was tested in broth medium as previously described [24]. Tellurite tolerance was tested on MRS agar supplemented with 0.02% of potassium tellurite. Carbon source utilization by strain FI11369^T and the closest relative *L. xiangfangensis* DSM 27103^T (Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH) was examined using the API 50 CH system (bioMérieux) following the manufacturer's instructions. Strips were incubated at 30 °C and readings were made after 48 h (Table 2). Whole-cell fatty acids were analysed by gas chromatography of fatty

Fig. 3. Scanning electron microscope image of strain FI11369^T cells after 48 h incubation at 30 °C in MRS broth. Bar, 1 μm.

Table 2. Distinctive features of the carbohydrate fermentation profiles of strain FI11369^T and closest phylogenetically related species

Strains: 1, FI11369^T (data from this study); 2, *Lactobacillus xiangfangensis* DSM 27103^T (data from this study); 3, *Lactobacillus plajomi* NBRC107333^T [22]; 4, *Lactobacillus modestisalitolerans* NBRC 107235^T [22]. +, Positive; –, negative; w, weak reaction; d, delayed (>72h).

Carbon source	Strain			
	1	2	3	4
D-Xylose	w	w	–	–
D-Adonitol	–	w	–	–
D-Galactose	w	–	+	+
Amygdalin	+	–	–	–
Arbutin	+	–	–	–
Aesculin	+	d	w	w
Salicin	+	–	+	+
Lactose	–	–	–	+
Melibiose	–	–	–	+
Raffinose	–	–	–	+
L-Arabitol	–	–	+	–
Gluconate	–	–	+	+

Table 3. Comparative fatty acid compositions of strains FI11369[T] and the closely related *Lactobacillus* species

Strains: 1, FI11369[T] (data from this study); 2, *Lactobacillus xiangfangensis* DSM 27103[T] (data from this study); 3, *Lactobacillus plajomi* NBRC107333[T] [24]; 4, *Lactobacillus modestisalitolerans* NBRC 107235[T] [22]. Values are percentages of total fatty acids. Fatty acids amounting to less than 0.5% of the total fatty acids are not shown. ND, not detected. The major components of cellular fatty acid are highlighted in bold.

Fatty acid	Strain			
	1	**2**	**3**	**4**
Saturated:				
$C_{14:0}$	0.7	0.8	6.6	ND
$C_{16:0}$	**18.2**	**20.0**	**11.4**	**13.0**
$C_{18:0}$	2.9	2.5	ND	ND
$C_{19:0}$ iso	2.0	ND	ND	ND
Unsaturated:				
$C_{16:1}$ $\omega7c$ and/or $\omega6c$	1.1	0.9	2.6	ND
$C_{18:1}$ $\omega9c$	**43.8**	**28.1**	**36.1**	**51.5**
$C_{18:1}$ $\omega6c$ and/or $\omega7c$	8.8	6.9	9.7	ND
Cyclopropane:				
$C_{19:0}$ cyclo $\omega10c$ and/or $\omega6$	**22.5**	**40.8**	**33.7**	**35.6**

acid methyl esters (GC-FAME) using the Sherlock microbial identification method [25, 26] after growth of the strain in trypticase soy medium for 2 days at 28 °C. Peptidoglycan was isolated from cells grown in MRS broth for 10 h at 30 °C and its structure and cell-wall sugar composition was studied after total hydrolysis (100 °C, 4 N HCl, 16 h) and partial hydrolysis (4 N HCl, 45 min, 100 °C) as previously described [27]. Both analyses were performed by the Leibniz Institute DSMZ-German Collection of Microorganisms and Cell Cultures identification service. The main fatty acids (>10%) of FI11369[T] were saturated $C_{16:0}$ (18.2%), unsaturated $C_{18:1}$ $\omega9c$ (43.8%) and cyclopropane $C_{19:0}$ cyclo ($\omega10c$ and/or $\omega6$; 22.5%) (Table 3). The total hydrolysate of the peptidoglycan of strain FI11369 contained muramic acid (Mur) and the amino acids diaminopimelic acid (Dpm), alanine (Ala) and glutamic acid (Glu) in a molar ration of 1.6:0.9:1.6:1. Enantiomeric analysis of the peptidoglycan amino acids revealed the presence of meso-Dpm. The partial hydrolysate contained peptides M-Glu, Ala-Glu, Glu-Dpm, Dpm-Ala, Glu-Dpm-Ala and Glu-Dpm-Ala-Dpm. These data strongly suggested that the peptidoglycan type of the strain was A1γ meso-Dpm direct.

DESCRIPTION OF *LACTOBACILLUS GARII* SP. NOV.

Lactobacillus garii (ga'ri.i. N.L. gen. n. *garii* of *gari*).

Cells of strain FI11369[T] are Gram-positive, non-motile, non-spore-forming, catalase-negative, straight rod-shaped,

1–2 µm long, and usually occur in pairs or in short chains. They are facultative anaerobes. Colonies grown aerobically on MRS agar at 30 °C for 48 h are irregular and umbonate. Strain FI11369[T] grows at 12–40 °C (weakly at 6 and 42 °C, optimum at 30–37 °C), at pH range 4.0–8.8 (optimum at pH 6) and with 0–8% NaCl (delayed growth at 10%, optimum in the absence of NaCl supplementation). Gas is not produced from glucose. Only D-lactate is synthesized from glucose. Acid is produced from D-ribose, D-xylose, D-galactose, D-glucose, D-fructose, D-mannose, D-mannitol, D-sorbitol, *N*-acetyl glucosamine, amygdalin, arbutin, aesculin, salicin, cellobiose, maltose, sucrose, trehalose and gentiobiose. Acid is not produced from glycerol, erythritol, D-arabinose, L-arabinose, L-xylose, D-adonitol, ß-D-xylopyranoside, L-sorbose, L-rhamnose, dulcitol, inositol, α-D-mannopyranoside, α-D-glucopyranoside, lactose, melibiose, inulin, melezitose, raffinose, starch, glycogen, xylitol, turanose, D-lyxose, D-tagatose, D-fucose, L-fucose, D-arabitol, L-arabitol, gluconate and 2- or 5-keto-gluconate. Strain FI11369[T] is positive for α-haemolytic activity, bile–aesculin test and Voges–Proskauer test and negative for pyruvate utilization, tellurite tolerance, hippurate hydrolysis, pyrrolidonylarylamidase production, deamination of arginine, H_2S production, nitrate and nitrite reduction, urease production and gelatin hydrolysis. Cellular fatty acids mainly comprised saturated $C_{16:0}$, unsaturated $C_{18:1}$ $\omega9c$ and cyclopropane $C_{19:0}$ cyclo ($\omega10c$ and/or $\omega6$). Cells contain *meso*-diaminopimelic acid in their cell-wall peptidoglycan. The peptidoglycan type is A1γ meso-Dpm

direct. The genome size of the type strain is 2972171 bp and the DNA G+C content is 48.3 mol%.

The type strain, FI11369[T] (=NCIMB 15148=DSM 108249), was isolated in the UK from *gari* produced in Ghana. The GenBank accession numbers of the 16S rRNA gene and the genome sequence of FI11369[T] are MN817919 and QWZQ00000000, respectively.

Author contributions
Maria Diaz: Conceptualization, Investigation, Visualization, Writing - original draft. Lizbeth Sayavedra: Writing – review and editing. Amy Atter: Resources. Shikha Saha: Investigation. Melinda J Mayer: Funding acquisition, Writing - review and editing. Wisdom Amoa-Awa: Resources, Writing - review and editing. Arjan Narbad: Funding acquisition, Writing - review and editing.

Conflicts of interest
The authors declare no conflicts of interest.

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