



Draft Genome Sequence of *Lactobacillus fermentum* BFE 6620, a Potential Starter Culture for African Vegetable Foods, Isolated from Fermented Cassava

Eliud N. Wafula,^{a,b,c} Erik Brinks,^b Biserka Becker,^a Melanie Huch,^a Bernhard Trierweiler,^a Julius M. Mathara,^c Folarin A. Oguntoyinbo,^d Gyu-Sung Cho,^b Charles M. A. P. Franz^b

Department of Safety and Quality of Fruit and Vegetables, Federal Research Institute of Nutrition and Food, Kiel, Germany^a; Department of Microbiology and Biotechnology of the Max Rubner-Institut, Federal Research Institute of Nutrition and Food, Kiel, Germany^b; Jomo Kenyatta University of Agriculture and Technology, Food Science and Technology, Nairobi, Kenya^c; Department of Microbiology, Faculty of Science, University of Lagos, Akoka, Lagos, Nigeria^d

ABSTRACT We report the draft genome sequence of *Lactobacillus fermentum* BFE 6620 from fermented cassava used as a potential starter culture for African vegetable fermentation. Sequence analysis showed the assembled genome size to be 1,982,893 bp, encoding a predicted total of 2,003 protein-coding genes, 14 rRNAs, 54 tRNAs, and 3 noncoding RNAs (ncRNAs).

Lactobacillus fermentum is a heterofermentative lactic acid bacterium belonging to the *Bacilli* class of the phylum *Firmicutes* and the family *Lactobacillaceae*. This species occurs in diverse habitats, including the human gut, milk products, fermenting plant material, and animals (1). It is considered to be a good probiotic candidate, due to its ability to withstand gastrointestinal conditions (2), and was reported to have potential for prevention of community-acquired infections (3), modulation of the immune system, and production of antimicrobial compounds (4).

Lactobacillus fermentum BFE 6620 was isolated from fermented cassava for production of gari in Benin. This strain, together with *Lactobacillus plantarum* BFE 5092, was successfully used as a starter culture in the fermentation of African kale leaves (5). There are currently 25 *L. fermentum* genome sequences reported, of which 6 were completely sequenced. The genome of strain BFE 6620 was sequenced in order to assess its technological and functional properties for vegetable food fermentation and to compare its genome sequence with already sequenced *L. fermentum* strains from different sources.

The total genomic DNA of *L. fermentum* BFE 6620 was isolated using the peqGOLD bacterial DNA kit (Peqlab, Erlangen, Germany). The sequencing library was prepared with an Illumina Nextera XT library prep kit (Illumina, San Diego, CA, USA) and run on the MiSeq with 2 × 251 paired ends. In total, 2,429,489 paired-end sequence reads were obtained with an approximately 242-fold coverage, and the reads were assembled *de novo* using SPAdes version 3.10.1 (6). The draft genome assembly consisted of 149 scaffolds, and the N_{50} was 35,982. The genome size of *L. fermentum* BFE 6620 is 1,982,893 bp, with a 52.1 mol% G+C content. The genome sequence was annotated using the Rapid Annotations Subsystems Technology (RAST) and NCBI (7) servers. The sequence contained 2,003 protein-coding sequences, 14 rRNAs, 54 tRNAs, and 3 noncoding RNAs (ncRNAs). No acquired antibiotic resistance genes were found using ResFinder server (v. 2.1) (8). With the use of the RAST server, draft genome comparison with reference strain *L. fermentum* IFO3956 (GenBank accession no. AP008937) showed

Received 29 June 2017 Accepted 3 July 2017 Published 17 August 2017

Citation Wafula EN, Brinks E, Becker B, Huch M, Trierweiler B, Mathara JM, Oguntoyinbo FA, Cho G-S, Franz CMAP. 2017. Draft genome sequence of *Lactobacillus fermentum* BFE 6620, a potential starter culture for African vegetable foods, isolated from fermented cassava. *Genome Announc* 5:e00801-17. <https://doi.org/10.1128/genomeA.00801-17>.

Copyright © 2017 Wafula et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Gyu-Sung Cho, gyusung.cho@mri.bund.de.

that the *L. fermentum* BFE 6620 contained 88 coding genes for proteins involved in phosphoenolpyruvate/phosphotransferase (PEP/PTS) systems for utilization of trehalose, a malolactic enzyme, and a pyridoxamine 5'-phosphate oxidase (involved in vitamin B₆ biosynthesis), which were not present in the reference strain.

Accession number(s). This whole-genome shotgun project has been deposited at DDBJ/ENA/GenBank under the accession no. [NIWV00000000](https://doi.org/10.1093/nar/nwz000).

ACKNOWLEDGMENTS

The authors from the Max Rubner-Institut thank the German Federal Ministry for Education and Research for funding of the project Diversifying Food Systems, Horticultural Innovations and Learning for Improved Nutrition and Livelihood in East Africa (HORTINLEA) (project no. FKZ 031A248I), as well as the Alexander von Humboldt Foundation's Georg Foster Stipendium for Experienced Researcher for financing F. A. Oguntoyinbo. The responsibility for the article content lies solely with the authors.

REFERENCES

- Dellaglio F, Torriani S, Felis GE. 2004. Reclassification of *Lactobacillus cellobiosus* Rogosa et al. 1953 as a later synonym of *Lactobacillus fermentum* Beijerinck 1901. *Int J Syst Evol Microbiol* 54:809–812. <https://doi.org/10.1099/ijs.0.02947-0>.
- Jiménez E, Langa S, Martín V, Arroyo R, Martín R, Fernández L, Rodríguez JM. 2010. Complete genome sequence of *Lactobacillus fermentum* CECT 5716, a probiotic strain isolated from human milk. *J Bacteriol* 192:4800–4800. <https://doi.org/10.1128/JB.00702-10>.
- López-Huertas E. 2015. Safety and efficacy of human breast milk *Lactobacillus fermentum* CECT 5716. A mini-review of studies with infant formulae. *Benef Microbes* 6:219–224. <https://doi.org/10.3920/BM2014.0091>.
- Olivares M, Díaz-Ropero MP, Martín R, Rodríguez JM, Xaus J. 2006. Antimicrobial potential of four *Lactobacillus* strains isolated from breast milk. *J Appl Microbiol* 101:72–79. <https://doi.org/10.1111/j.1365-2672.2006.02981.x>.
- Oguntoyinbo FA, Cho GS, Trierweiler B, Kabisch J, Rösch N, Neve H, Bockelmann W, Frommherz L, Nielsen DS, Krych L, Franz CMAP. 2016. Fermentation of African kale (*Brassicacarinata*) using *L. plantarum* BFE 5092 and *L. fermentum* BFE 6620 starter strains. *Int J Food Microbiol* 238:103–112. <https://doi.org/10.1016/j.ijfoodmicro.2016.08.030>.
- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single cell sequencing. *J Comput Biol* 19:455–477. <https://doi.org/10.1089/cmb.2012.0021>.
- Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, Meyer F, Olsen GJ, Olson R, Osterman AL, Overbeek RA, McNeil LK, Paarmann D, Paczian T, Parrello B, Pusch GD, Reich C, Stevens R, Vassieva O, Vonstein V, Wilke A, Zagnitko O. 2008. The RAST server: rapid annotations using subsystems technology. *BMC Genomics* 9:75. <https://doi.org/10.1186/1471-2164-9-75>.
- Zankari E, Hasman H, Cosentino S, Vestergaard M, Rasmussen S, Lund O, Aarestrup FM, Larsen MV. 2012. Identification of acquired antimicrobial resistance genes. *J Antimicrob Chemother* 67:2640–2644. <https://doi.org/10.1093/jac/dks261>.