

Genome Sequence of *Lactobacillus gastricus* PS3, a Strain Isolated from Human Milk

Virginia Martín,^a Nivia Cárdenas,^a Esther Jiménez,^a Antonio Maldonado,^b Juan Miguel Rodríguez,^a Leonides Fernández^a

Departamento de Nutrición, Bromatología y Tecnología de los Alimentos, Universidad Complutense de Madrid, Madrid, Spain^a; Departamento de Biotecnología de Alimentos, Instituto de la Grasa, Consejo Superior de Investigaciones Científicas–CSIC, Seville, Spain^b

V.M. and N.C. contributed equally to this work.

***Lactobacillus gastricus* is a mostly unknown lactobacilli species associated with mucosal surfaces. We present the draft annotated genome sequence of *L. gastricus* strain PS3, isolated from a human milk sample, to provide new insights into its biology and to characterize those genes related to advantageous technological and beneficial properties.**

Received 6 June 2013 Accepted 7 June 2013 Published 11 July 2013

Citation Martín V, Cárdenas N, Jiménez E, Maldonado A, Rodríguez JM, Fernández L. 2013. Genome sequence of *Lactobacillus gastricus* PS3, a strain isolated from human milk. *Genome Announc.* 1(4):e00489-13. doi:10.1128/genomeA.00489-13.

Copyright © 2013 Martín et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 3.0 Unported license](https://creativecommons.org/licenses/by/3.0/).

Address correspondence to Leonides Fernández, leonides@vet.ucm.es.

Lactobacillus gastricus was first isolated from gastric biopsy specimens from healthy humans and identified and described by Roos et al. (1, 2). This species has been associated with the *Lactobacillus reuteri* group according to its phylogenetic relatedness (3, 4). A recent investigation of breast milk from healthy women revealed the presence of *L. gastricus* strains in about one-third of 20 milk samples analyzed, together with other *Lactobacillus* species such as *Lactobacillus casei*, *Lactobacillus gasseri*, *Lactobacillus fermentum*, *Lactobacillus plantarum*, *L. reuteri*, *Lactobacillus salivarius*, and *Lactobacillus vaginalis* (5). The human milk microbiota has been regarded as essential for initiation and development of infant gut colonization (6–8). Furthermore, strains of *L. gasseri*, *L. fermentum*, and *L. salivarius* isolated from this biological fluid have demonstrated excellent probiotic potential (9, 10). After an initial characterization of technological and probiotic properties in a collection of *L. gastricus* strains isolated from human milk, strain PS3 was selected on the basis of some traits of probiotic and technological relevance and identified via 16S rRNA gene sequencing.

In order to get a deeper knowledge of the technological and probiotic properties of this strain, we performed whole-genome sequencing of *L. gastricus* PS3 by 454 pyrosequencing on a GS-FLX sequencer to 19.14-fold coverage (454 Life Sciences, Branford, CT). The initial draft assembly generated 93 contigs using the Newbler program version 2.3 (Roche Applied Science). The draft genome of *L. gastricus* PS3 consists of 1,904,872 bases with an average GC content of 41.8% and contains a total of 1,386 protein-encoding sequences and 43 RNA-encoding sequences (40 tRNAs and 3 rRNAs). Coding regions were predicted using the BG7 system (Era7 Technologies, Granada, Spain), which proceeds from protein similarity detection to open reading frame (ORF) prediction and is tolerant of sequencing errors in start and stop codons, frameshifts, and assembly or scaffolding errors (11). The semiautomatic annotation of the sequences resulted in 80 final contigs, 1,269 protein-coding genes, 40 tRNA-encoding genes, and 3 rRNAs.

More than 20 peptidases and proteases and several peptide transporter genes were predicted. This highly complex proteolytic system would allow the growth in milk and compensate for deficiencies in amino acid biosynthesis. Significantly, genes encoding three different putative glutamate-cysteine ligases (PS3_6383, PS3_8988, and PS3_21018) that synthesize γ -glutamylcysteine were detected. This is the major low-molecular-weight thiol that protects against oxidative stress in some lactic acid bacteria (12). Also, one putative gene coding glutamate decarboxylase (PS3_14606) was identified. This enzyme catalyzes the synthesis of γ -aminobutyric acid, an amino acid that contributes to bacterial acid resistance and has potential as a bioactive compound in humans (13). Putative genes coding for transport systems and enzymes related to utilization of diverse carbohydrates such as fructose, galactitol, mannose, lactose, cellobiose, sucrose, and β -glucosides were also identified, as well as genes encoding two putative esterases and a GDSL lipolytic enzyme.

Nucleotide sequence accession numbers. The results of this whole-genome shotgun project have been deposited at DDBJ/EMBL/GenBank under the accession number [AICN00000000](https://www.ncbi.nlm.nih.gov/nuccore/AICN00000000). The version described in this paper is the first version, [AICN01000000](https://www.ncbi.nlm.nih.gov/nuccore/AICN01000000).

ACKNOWLEDGMENTS

This study was partly supported by the FUN-C-FOOD (Consolider-Ingenio 2010) and AGL2010-15420 projects from the Ministerio de Economía y Competitividad (Spain).

REFERENCES

1. Endo A, Roos S, Satoh E, Morita H, Okada S. 2008. *Lactobacillus equigenerosi* sp. nov., a coccoid species isolated from faeces of thoroughbred racehorses. *Int. J. Syst. Evol. Microbiol.* 58:914–918.
2. Roos S, Engstrand L, Jonsson H. 2005. *Lactobacillus gastricus* sp. nov., *Lactobacillus antri* sp. nov., *Lactobacillus kalixensis* sp. nov. and *Lactobacillus ultunensis* sp. nov., isolated from human stomach mucosa. *Int. J. Syst. Evol. Microbiol.* 55:77–82.

3. Felis GE, Dellaglio F. 2007. Taxonomy of lactobacilli and bifidobacteria. *Curr. Issues Intest. Microbiol.* 8:44–61.
4. Naser SM, Dawyndt P, Hoste B, Gevers D, Vandemeulebroecke K, Cleenwerck I, Vancanneyt M, Swings J. 2007. Identification of lactobacilli by *pheS* and *rpoA* gene sequence analyses. *Int. J. Syst. Evol. Microbiol.* 57:2777–2789.
5. Martín V, Maldonado-Barragán A, Moles L, Rodríguez-Baños M, Campo RD, Fernández L, Rodríguez JM, Jiménez E. 2012. Sharing of bacterial strains between breast milk and infant feces. *J. Hum. Lact.* 28: 36–44.
6. Jiménez E, Delgado S, Maldonado A, Arroyo R, Albújar M, García N, Jariod M, Fernández L, Gómez A, Rodríguez JM. 2008. *Staphylococcus epidermidis*: a differential trait of the fecal microbiota of breast-fed infants. *BMC Microbiol.* 8:143.
7. Martín R, Langa S, Reviriego C, Jiménez E, Marín ML, Xaus J, Fernández L, Rodríguez JM. 2003. Human milk is a source of lactic acid bacteria for the infant gut. *J. Pediatr.* 143:754–758.
8. Martín R, Heilig GH, Zoetendal EG, Smidt H, Rodríguez JM. 2007. Diversity of the *Lactobacillus* group in breast milk and vagina of healthy women and potential role in the colonization of the infant gut. *J. Appl. Microbiol.* 103:2638–2644.
9. Martín R, Olivares M, Marín ML, Fernández L, Xaus J, Rodríguez JM. 2005. Probiotic potential of 3 lactobacilli strains isolated from breast milk. *J. Hum. Lact.* 21:8–17.
10. Martín R, Jiménez E, Olivares M, Marín ML, Fernández L, Xaus J, Rodríguez JM. 2006. *Lactobacillus salivarius* CECT 5713, a potential probiotic strain isolated from infant feces and breast milk of a mother-child pair. *Int. J. Food Microbiol.* 112:35–43.
11. Pareja-Tobes P, Manrique M, Pareja-Tobes E, Pareja E, Tobes R. 2012. BG7: A new approach for bacterial genome annotation designed for next generation sequencing data. *PLoS One* 7:e49239. doi:10.1371/journal.pone.0049239.
12. Kim EK, Cha CJ, Cho YJ, Cho YB, Roe JH. 2008. Synthesis of gamma-glutamylcysteine as a major low-molecular-weight thiol in lactic acid bacteria *Leuconostoc* spp. *Biochem. Biophys. Res. Commun.* 369:1047–1051.
13. Gobbetti M, Cagno RD, De Angelis M. 2010. Functional microorganisms for functional food quality. *Crit. Rev. Food Sci. Nutr.* 50:716–727.