## Complete Genome Sequence of *Lactobacillus salivarius* CECT 5713, a Probiotic Strain Isolated from Human Milk and Infant Feces<sup>∇</sup>

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Lactobacillus salivarius is a homofermentative lactic acid bacterium and is frequently isolated from mucosal surfaces of healthy humans. L. salivarius CECT 5713, a strain isolated simultaneously from breast milk and infant feces of a healthy mother-infant pair, has immunomodulatory, anti-inflammatory, and anti-infectious properties, as revealed by several *in vitro* and *in vivo* assays. Here, we report its complete and annotated genome sequence.

In the last years, culture-dependent and -independent analyses of the bacterial diversity of human milk and colostrum have revealed that these biological fluids are a source of live staphylococci, streptococci, lactic acid bacteria, and bifidobacteria in the infant gut (5, 6, 8, 9, 11, 13), where they play a key role in the initiation and development of the gut microbiota (12). In a previous study, we isolated L. salivarius CECT 5713 from human milk and infant feces of a mother-child pair (10). Subsequent studies revealed that this strain was a good probiotic candidate since it achieved high survival rates when exposed to the gastrointestinal tract conditions, showed a strong adherence to intestinal cells, stimulated the expression of mucin-encoding genes, produced antimicrobial compounds (lactate, acetate, and hydrogen peroxide), and displayed in vivo and in vitro immunomodulatory, anti-inflammatory, and antibacterial properties against pathogenic bacteria (2, 10, 15). Moreover, oral administration of L. salivarius CECT 5713 appears to be an efficient alternative for the treatment of infectious mastitis in lactating women (7). Similarly, studies with other L. salivarius strains in animal models and clinical trials have demonstrated their probiotic function and, particularly, their anti-inflammatory effects (3, 14, 16).

In order to interrogate the genome sequence of *L. salivarius* CECT 5713 with regard to its probiotic properties, the complete genome sequence was determined by a whole-genome shotgun strategy using pyrosequencing technology (454 Life Sciences, Banford, CT). The initial draft assembly provided by 454 Life Sciences was based on 444,604 high-quality pyrosequencing reads, which assembled into 59 contigs. The genome sequence of *L. salivarius* UCC118 (1), a well-characterized probiotic strain, was used to order these contigs into large scaffolds.

The genome of *L. salivarius* CECT 5713 consists of a circular chromosome of 1,828,169 bp, two plasmids (pHN1, 44,581 bp;

pHN2, 20,426 bp), and a megaplasmid (pHN3, 242,962 bp). The overall GC content of the chromosome is 32.93%, similar to that of the megaplasmid but lower than those of the plasmids (>38%). The entire genome of CECT 5713 contains 1,558 protein-, 87 tRNA-, and 51 rRNA-encoding genes. A comparison between the genomes of L. salivarius CECT 5713 and UCC118 revealed the presence of 52 protein-encoding genes that are exclusive for CECT 5713, including genes encoding a 6-phospho-β-glucosidase and three collagen-binding proteins, which may explain the high potential for competitive exclusion of pathogens displayed by this strain. The genes responsible for the bacteriocin activity of L. salivarius CECT 5713 are located in pHN3. This megaplasmid contains six open reading frames (ORFs) closely related, but not identical, to the genes responsible for the biosynthesis of salivaricin ABP-118, a two-component class II bacteriocin (4), in L. salivarius UCC118. Globally, several features of the L. salivarius CECT 5713 genome suggest a strong probiotic potential in humans.

**Nucleotide sequence accession numbers.** Genome information for the chromosome, the two plasmids, and the megaplasmid of *L. salivarius* CECT 5713 has been deposited in the EMBL and GenBank databases with accession numbers CP002034, CP002035 and CP002036, and CP002037, respectively.

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