P40 and P75 Are Singular Functional Muramidases Present in the Lactobacillus casei/paracasei/rhamnosus Taxon

Christine Bäuerl1†, Gulyaim Abitayeva2,3†, Sebastián Sosa-Carrillo4, Ana Mencher-Beltrán5, Noemí Navarro-Lleó†, José M. Coll-Marqués1, Manuel Zúñiga-Cabrera1, Serik Shaikhin2,3 and Gaspar Pérez-Martínez1∗

1 Departamento de Biotecnología, Instituto de Agroquímica y Tecnología de Alimentos (C.S.I.C.), Valencia, Spain, 2 Department of Microbiology, L.N. Gumilyov Eurasian National University, Astana, Kazakhstan, 3 Laboratory of Genetics and Biochemistry of Microorganisms, Republican Collection of Microorganisms at Science Committee of Ministry of Education and Science RK, Astana, Kazakhstan, 4 Computational Biology Department, Inria, Institut Pasteur and Université Paris Diderot, Paris, France, 5 Instituto de Ciencias de la Vid y del Vino (CSIC), Logroño, Spain

Lactobacillus casei and Lactobacillus rhamnosus proteins P40 and P75 belong to a large family of secreted cell wall proteins that contain a carboxy(C)-terminal CHAP or NlpC/P60 superfamily domains. In addition to their peptidoglycan hydrolases activity, proteins in this family are specific antigens of pathogens, frequently responsible of interactions with the host. L. rhamnosus GG and L. casei BL23 purified P40 and P75 proteins have antiapoptotic activity by inducing the EGF/Akt pathway. The aim of this work was to study the genetics, phylogeny and dissemination of this family of proteins in the genus Lactobacillus as well as their characteristics and likely function. The scrutiny of their DNA encoding sequences revealed the presence of minisatellite DNA in the P75 encoding gene of L. casei/paracasei strains (cmuB) with intraspecific indels that gave rise to four different alleles (cmuB1–4), which are exclusive of this species. Phylogenic analyses suggest that both proteins are present mainly in the L. casei and Lactobacillus sakei phylogenomic groups. A P40 ancestral gene was possibly present in the common ancestor of Enterococcaceae, Lactobacillaceae and Streptococcaceae. P75 is also present in L. casei and L. sakei groups, but its evolution is difficult to explain only by vertical transmission. Antibodies raised against the N-terminal regions of P40 and P75 improved their immunological detection in culture supernatants as they recognized almost exclusively proteins of L. casei/paracasei/rhamnosus strains, highlighting their structural similarity, that allowed to detect them in different fermented dairy products that contained probiotic L. casei strains. Purified P40 and P75 proteins showed no evident lytic activity but they complemented L. casei BL23 cmuA and cmuB defective mutants, respectively, thus proving that they actively participate in cell division.

Keywords: P40, P75, secreted muramidases, genetic analysis, minisatellite, phylogeny, Lactobacillus casei, Lactobacillus rhamnosus
INTRODUCTION

Proteins carrying an NlpC/P60 domain constitute a large and widely distributed superfamily of bacterial proteins. However, they remain largely unexplored. An important subgroup contains the NlpC/P60 domain at the C-terminal region, and it includes phage and bacterial surface bound/secreted peptidoglycan (PGN) hydrolases, muramidases or lysins (Anantharaman and Aravind, 2003). They are also typical antigens or pathogenicity factors in bacterial pathogens (Hess et al., 1995; Park et al., 2000; Teng et al., 2003; Oechslin et al., 2013). Their enzymatic activity is determined by this conserved domain and relies on a catalytic triad formed by a cysteine that acts as a nucleophile, a catalytic histidine and a charged amino acid, frequently another histidine – but also glutamic acid, aspartic acid or asparagine-. For this reason, these domains were also called cysteine, histidine-dependent amidohydrolases/peptidases (CHAP) (Bateman and Rawlings, 2003; Aramini et al., 2008; Sanz-Gaitero et al., 2014).

In Firmicutes, cell wall maintenance and reshaping during cell division involves a wide variety of enzymes (Vermassen et al., 2019); among them, CHAP-containing proteins have been associated to cell wall degradation and cell division (Layec et al., 2008). NlpC/P60 C-terminus domain proteins are highly represented in most bacterial genomes and it is very frequent to find between three to five genes encoding CHAP-containing proteins (see below). Their amino(N)-terminal regions have been related to substrate recognition, as in the case of domains LysM (Visweswaran et al., 2014) and SH3 (Gründling and Schneeewind, 2006), however, there are exceptions like the Var-Cse region is necessary for the species. Furthermore, this Var-Cse region is necessary for normal cell division (Borges et al., 2006).

Cysteine, histidine-dependent amidohydrolases/peptidases-containing proteins are also involved in bacteria-host interactions. In Lactobacillus rhamnosus GG, P40 and P75 were found to inhibit epithelial cell apoptosis and to promote cell growth (Yan et al., 2007). Their homologous counterparts in L. casei also displayed pro-proliferative and anti-apoptotic activity that was demonstrated in vitro and in vivo (Wang et al., 2014, 2016; Shen et al., 2018). These proteins induced the phosphorylation of the epidermal growth factor receptor (EGFR) and other intermediates in the EGFR/Akt signal transduction pathway (Bäuerl et al., 2010; Yan et al., 2011, 2013; Yan and Polk, 2012). In addition, it has been proposed that the functional activity of some PGN hydrolases may be explained by their ability to hydrolyse PGN ligands that induce NOD2 signal transduction pathways, thus subverting host innate immune response (Humann and Lenz, 2009). Furthermore, P40 biological function has been extended to the induction of IgA synthesis (Wang et al., 2016).

Until present, research on P40 and P75 has shown that they are secreted cell wall muramidases encoded by cmuA and cmuB in L. casei and mspB and mspA in L. rhamnosus GG (LLG) and have muramidase activity. More specifically LGG P75 has γ-D-glutamyl-L-lysyl-endopeptidase activity. The observation of knock out mutants indicated that these proteins are possibly required for the normal conformation of the cell wall or separation of bacterial daughter cells in both, L. casei and L. rhamnosus (Bäuerl et al., 2010; Regulski et al., 2012). P40 and P75 might share high similarity within L. casei/paracasei species, as anti-P40 antibodies recognized P40 and P75 in a large number of strains (Bäuerl et al., 2010), however, no further characterization and comparative studies have been carried out. The purpose of this work was to study the structure, physiological function, phylogenetic and genetic features in these biologically interesting proteins, in order to highlight links among strains in the L. casei/paracasei/rhamnosus taxon and to determine if they are exclusively present in this bacterial group.

Taxonomy remark: In this work, we respected the existing species name in the annotated sequence databases. L. casei BL23 and L. rhamnosus GG belong to the L. casei/paracasei/rhamnosus group, recently included in the L. casei phylogenomic group (Salvetti et al., 2018), a group that has some taxonomical controversy regarding L. casei and L. paracasei strains. L. casei BL23 is taxonomically different to the type strain L. casei ATCC 393T (Acedo Felix and Perez Martinez, 2003) and more similar to L. paracasei, as well as a very large number of strains that are still designated as L. casei. Consequently and due to the low number of strains homologous to the standing type strain (ATCC 393T), L. casei will be equivalent to L. paracasei with the exception of L. casei ATCC 393T.

MATERIALS AND METHODS

Bacterial Strains and Culture Conditions

Lactobacillus strains were grown on MRS medium (DIFCO) at 37°C, except L. sakei, L. pentosus and L. curvatus strains that were grown at 30°C. All lactobacilli were stored at −80°C in 15% glycerol in the laboratory’s collection and their respective origins are listed in Table 1. Listeria monocytogenes BL1001 (CECT 932) and Staphylococcus aureus BL102 (CECT 86) were grown on Brain Heart Infusion (Conda-Prondasida) and Enterococcus faecalis BL141 (laboratory isolate) was cultured in Bacto™ Todd Hewitt broth (Becton Dickinson) static at 37°C. The cloning hosts were Escherichia coli DH5α and DH10B and pQE80e derived plasmids were introduced into E. coli BL21(DE3)-pLysS for protein expression and purification. They were grown in LB medium at 37°C under agitation. Recombinant plasmids in E. coli were selected with ampicillin at 100 μg/ml and chloramphenicol at 20 μg/ml. Solid medium was prepared by adding 1.8% (w/v) agar. Strains were identified by PCR amplification and 16S rDNA sequencing using standard primers...
TABLE 1 | List of *Lactobacillus* strains used in this study.

| Collection number | Species | Collection number or source (Origin) |
|-------------------|---------|-------------------------------------|
| BL6               | *L. casei* | ATCC393<sup>T</sup> (Dairy product) |
| BL23              | *L. casei* | [Bruce Chassy, U. Urbana (Illinois, United States)] |
| BL28              | *L. casei* | 64H (C.-A. Alpert, U. Ösnabrück, Germany) |
| BL32              | *L. casei* | CECT 4040 (Majoreiro Cheese) |
| BL81              | *L. casei* | Commercial Dairy product* |
| BL82              | *L. casei* | CECT 277 (Sour milk) |
| BL83              | *L. casei* | CECT 4043 (Majoreiro Cheese) |
| BL86              | *L. casei* | CECT 4045 (Majoreiro Cheese) |
| BL87              | *L. casei* | ATCC 11578 (Oral Cavity) |
| BL90              | *L. casei* | ATCC 334 (Emmental Cheese) |
| BL101             | *L. casei* | Commercial Dairy product* |
| BL193             | *L. casei* | Commercial Dairy product* |
| BL199             | *L. casei* | CERELA 87 (Infant feces) |
| BL201             | *E. faecalis* | Laboratory isolate; adult feces |
| BL202             | *E. faecium* | Laboratory isolate; adult feces |
| BL203             | *L. casei* | Laboratory isolate; adult feces |
| BL205             | *L. casei* | Nordic fermented product* |
| BL206             | *L. casei* | Nordic fermented product* |
| BL212             | *L. casei* | ADNOX 95 (PROIMI-CERELA) |
| BL216             | *L. casei* | CECT 5289 (Unknown)** |
| BL227             | *L. casei* | Commercial Dairy product* |
| BL229             | *L. casei* | Commercial Dairy product* |
| BL312             | *L. paracasei* | Commercial Dairy product* |
| BL358             | *L. casei* | Commercial Dairy product* |
| BL1               | *L. rhamnosus* | CECT 278; ATCC 7469<sup>T</sup> |
| BL2               | *L. rhamnosus* | CECT 276; NCIB 8651 |
| BL3               | *L. rhamnosus* | CECT 275; NCIB 8963 |
| BL102             | *L. rhamnosus* | Laboratory isolate |
| BL103             | *L. rhamnosus* | Laboratory isolate (MRS contaminant) |
| BL327             | *L. rhamnosus* | CECT 288; ATCC 11979(9695) (Unknown) |
| BL367             | *L. rhamnosus* | Commercial Dairy product* |
| BL376             | *L. rhamnosus* | Commercial Dairy product* |
| BL377             | *L. rhamnosus* | ATCC 53103 (Human Feces) |
| BL378             | *L. rhamnosus* | Commercial Dairy product* |
| BL305             | *L. casei* | (cmuA<sup>-</sup>) (Bäuerl et al., 2010) |
| BL306             | *L. casei* | (cmuB<sup>-</sup>) (Bäuerl et al., 2010) |
| BL5               | *L. acidophilus* | CECT 289 (Unknown)** |
| BL7               | *L. brevis* | CECT 216 (Beer) |
| BL8               | *L. plantarum* | CECT 748 (Pickled cabbage) |
| BL13              | *L. sakei* | CECT 906 (Starter of sake, moto) |
| BL14              | *L. curvatus* | CECT 904 (Mik) |
| BL15              | *L. alimentarius* | CECT 570<sup>T</sup> (Unknown)** |
| BL26              | *L. plantarum* | CECT 221 (Grass silage) |
| BL33              | *L. delbrueckii subsp. bulgaricus* | CECT 4005<sup>T</sup> (Bulgarian Yogurt) |
| BL34              | *L. fermentum* | CECT 4007<sup>T</sup> (Fermented Beet) |
| BL35              | *L. pentosus* | CECT 4023<sup>T</sup> (Unknown)** |
| BL36              | *L. brevis* | CECT 4121<sup>T</sup> (Human feces) |
| BL47              | *L. pentosus* | MD533 (University of Amsterdam) |
| BL73              | *L. acidophilus* | CNRZ 55 (INRA Collection) |
| BL74              | *L. acidophilus* | CNRZ 216 (INRA Collection) |
| BL75              | *L. acidophilus* | CNRZ 217 (INRA Collection) |
| BL76              | *L. murinus* | CNRZ 220 (INRA Collection) |

(Continued)
TABLE 1 | Continued

| Collection number | Species | Collection number or source (Origin) |
|-------------------|---------|-------------------------------------|
| BL80              | L. sakei| 23K (INRA Collection, Meat product) |
| BL84              | L. brevis| Laboratory Isolate (MRS contaminant) |
| BL89              | L. delbrueckii subsp. lactis| DSM 7290 (Unknown)** |
| BL95              | L. zeae| ATCC15820 (Corn Step Liquor) |
| BL130             | L. corynformis subsp. corynformis| CECT 9827 (Silage) |
| BL131             | L. corynformis subsp. torquiens| CECT 4129T (Air of cowshed (dairy barn)) |
| BL132             | L. plantarum| CECT 748† (Pickled cabbage) |
| BL134             | L. paracasei subsp. tolerans| CECT 4175† (Pasteurized Milk) |
| BL135             | L. agilis| CECT 4131T (Municipal sewage) |
| BL156             | L. farciniminis| CECT 571† (Sausage) |
| BL157             | L. ruminis| CECT 4061† (Saliva) |
| BL158             | L. salivarius| CECT 4063† (Saliva) |
| BL159             | L. salivarius| CECT 4062† (Saliva) |
| BL160             | L. delbrueckii subsp. bulgaricus| Commercial Dairy product* |
| BL166             | L. plantarum| WCFS1 (Univ. Wageningen) |
| BL202             | L. acidophilus| Laboratory isolate; feces |
| BL207             | L. helveticus| Commercial Dairy product* |
| BL221             | L. crispatus| M247- Ma_Luisa Callegari (I. Microbiologia Piacenza) |
| BL223             | L. gasseri| 4B2- Ma_Luisa Callegari (I. Microbiologia Piacenza) |
| BL228             | L. acidophilus| Commercial probiotic* |
| BL231             | L. acidophilus| Commercial probiotic* |
| BL277             | L. gasseri| DSMZ 20243† (Human) |
| BL278             | L. crispatus| DSMZ 20584† (Eye) |
| BL279             | L. acidophilus| CECT 4529 (Unknown)** |
| BL280             | L. acidophilus| CECT 4179 (White rat feces) |
| BL281             | L. johnsonii| CECT 289 (Unknown)** |
| BL287             | L. johnsonii| DSMZ 10533† (Human Blood) |
| BL288             | L. intestinalis| DSMZ 6629† (Rat intestine) |

Symbols in the table: (*) Type Strain; (**) Commercial Brands are avoided in this work; (***) Strains with undefined origin or not found in current culture collections; CECT, Colección Española de Cultivos Tipo (Spain); ATCC, American Type Culture Collection (United States); DSMZ, Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH (Germany); CERELA, Centro de Referencia para Lactobacilos (Argentina).

27f and 1493r (Table 2) at the Genomics Service of the University of Valencia.

Cloning, Expression and Purification of P40 and P75

DNA fragments were amplified from L. casei BL23 chromosomal DNA with proofreading Expand blend (Expand High Fidelity PCR System, Roche, Mannheim, Germany). Specially designed primers were used to amplify the mature protein encoding sequences of P40 (LcasP40N-for and LcasP40C-rev) and P75 (LcasP75N-for and LcasP75C-rev), as well as the NH-terminal domains (N-domains) of P40 (LcasP40N-for and LcasP40N-rev) and P75 (LcasP75N-for and LcasP75N-rev), respectively. Primers had additional restriction sites to facilitate subsequent cloning. Sequence details are described in Table 2. PCR amplification conditions were as follows: 94°C, 3 min; 30 cycles of 94°C, 30 s; 50–55°C, 30 s; 72°C, 1.5 min; 72°C, 7 min for final extension. Amplicons were digested with BamHI and SmalI, ligated to pQE80e carrying the RGS-His epitope encoding region and transformed in E. coli DH10B in order to check inserts. Fragments with the correct DNA sequence were subcloned in E. coli BL21(BE3)-[pLysS] to induce expression with IPTG (see below). Restriction endonucleases and ligase were purchased from Gibco BRL (Invitrogen Corp., Carslab, CA, United States) and used as recommended by the manufacturer. Nucleic acid manipulation and general cloning procedures were performed according to laboratory manuals (Sambrook and Russell, 2001).

For protein purification, recombinant bacteria were grown in 500 ml of LB supplemented with 100 µg/ml ampicillin and 20 µg/ml chloramphenicol at 37°C. When bacterial cells reached an OD600 nm of 0.4, 1 mM IPTG was added and growth continued for 3 h. The bacterial pellets were recovered by centrifugation and resuspended in 30 ml of Binding Buffer A-His (50 mM Tris–Cl pH 7.5, 100 mM NaCl, 50 mM Na2SO4) containing 0.5 mM PMSF and 0.5 mg/ml lysozyme. Bacteria were lysed by sonication, and cell debris was removed by centrifugation at 10000 × g for 5 min at 4°C. The resulting supernatant was filtered (pore-size 0.45 µm) and loaded onto a HisTrap™ FF Crude Column (GE Healthcare Bio-Sciences AB, Uppsala, Sweden) using an ÄktaPrime™ Plus chromatography system (GE Healthcare Bio-Sciences AB, Uppsala, Sweden), and His-tagged proteins were separated according to the instructions of the manufacturer. Briefly, after washing of the column with
TABLE 2 | List of Oligonucleotides used in this study and their purposes.

| Name | Oligonucleotides | Purpose | References |
|------|------------------|---------|------------|
| 27f  | AGA7TGTGATCMTGGCTCACG | 16S rDNA sequencing | Lane, 1991 |
| 1439r | TACGCTTACTTTCACGACTTT | 16S rDNA sequencing | Lane, 1991 |
| LcaP40N-for | GTGAGATCAGCATAAGGCGACAG (BamHI) | Cloning | Bäuerl et al., 2010 |
| LcaP40N-rev | GTAGGGCCCTTTAGTGTGATATATC (SmaI) | Cloning | This work |
| LcaP40C-rev | GTAGGCGCTTCACGCGTGATATTA (SmaI) | Cloning | This work |
| LcaP75N-for | GTTGACATTCAGCGCGAGAACG (Bgl II) | Cloning | Bäuerl et al., 2010 |
| LcaP75N-rev | TAAAGGCGCTTTAGTGAAAGGAGC (SmaI) | Cloning | This work |
| LcaP75C-rev | TAAAGGCGCTTTAGTGAAAGGACG (SmaI) | Cloning | This work |
| P40carh-f1 | CACTTGAGGTCTATGACMTGGTAGACG (BL23 genome position 22495-22523) | PCR detection | This work |
| P40carh-r | CGGCTTGTGATGCTGAAGGCC (BL23 genome position 22296-22275) | PCR detection | This work |
| P75carh-f | GACCAACGTGATGGAAAGGCTCC (BL23 genome position 277183-277204) | PCR detection | This work |
| P75carh-r | GGRATCCACTGTGCTTGCGC (BL23 genome position 277321-277301) | PCR detection | This work |

Underlined sequences in cloning primers indicate the restriction enzyme site in brackets.

10 volumes of Buffer A-His and 10 volumes of Buffer A-His supplemented with 40 mM imidazole, the recombinant proteins were eluted using a 20 ml linear gradient from 40 to 500 mM imidazole in Buffer A-His (Buffer B). A constant flow rate of 1 ml/min was maintained in all the purification steps (binding-washing-elution).

Protein concentration was determined using the Bio-Rad Protein Assay Dye Reagent Concentrate (Bio-Rad, CA, United States) based on the Bradford method. All purified proteins were buffer-exchanged to Tris 50 mM pH 8.0, NaCl 100 mM, 1 mM EDTA and glycerol (15% v/v) was added before storage at −80°C.

Detection of P40 and P75 by PCR

DNA isolation of lactobacilli was performed using the REAL-pure SSS DNA™ extraction kit (Durviz, Spain) according to the manufacturer's instructions with some modifications. An aliquot of 1.5 ml of cell culture was centrifuged, the pellet was resuspended in 150 µl of RBC buffer (REAL-pure SSS DNA™ extraction kit), lysozyme (15 µl of a 20 mg/ml solution) and 15 U of mutanolysin were added and the cell suspension was incubated for 30 min at 37°C. Cells were centrifuged at 11000 x g for 1 min, the supernatant was removed and the pellet was vortexed to resuspend in 10–20 µl of residual liquid. Then, 300 µl of lysis solution and 10 µl RNase (10 mg/ml) were added. The remaining steps of the extraction were followed according to the instructions of the manufacturer. Quantitation of DNA was carried out in a Nanodrop™ spectrophotometer (Thermo Fisher Scientific, Waltham, MA, United States). Primers used for the detection tests of P40 (P40_casrham_for1, P40_casrham_rev) and P75 (P75_casrham_for, P75_casrham_rev) were developed as described in the section “Results” (Table 2). PCR reactions contained 70 ng of total bacterial DNA as a template and amplification conditions were 95°C, 4 min; 30 cycles at 95°C, 30 s, 59°C (P40) – 52°C (P75), 30 s; 72°C, 30 s; and a final extension of 72°C, 5 min.

Antagonistic/Lytic Activity of P40 and P75

Target bacterial strains (L. casei BL23, L. monocytogenes BL1001, E. faecalis BL143 and S. aureus BS102) were grown to OD550 0.4–0.6. Aliquots of 1 ml of these cultures were centrifuged, washed twice with 50 mM Tris Buffer pH 7.5, MgCl2 2.5 mM and adjusted to an OD550 0.1 in sterile peptone (10 g/l peptone, pH 6.3). Then, 0.9 ml of the cell suspensions were transferred to Eppendorf tubes and 100 µl of a concentrated solution of P40 and P75, individually or mixed, was added to obtain a final concentration of 50 µg/ml of each of them. Tubes were incubated at 37°C for 0 to 2 h. To test cell lysis, 55 µl of 1.0% Triton x100 was added to 445 µl of the samples taken at different times, mixed by inversion and kept on ice. When all samples were collected, OD550 of non-treated cell suspensions was measured against a blank with Triton x100. Antagonistic activity of P40 and P75 was evaluated by differences in OD550 values of peptone, and that of Triton x100-treated samples against a blank with Triton x100.

Isolation of Antibodies and Immunological Detection

Rabbit polyclonal antibodies were obtained by standard procedures (Leenaars and Hendriksen, 2005). Briefly, 0.5 to 1.0 mg of purified N-domains of P40 and P75 were dissolved in 0.5 ml of phosphate buffered saline (PBS) and filtered through 0.22 µm pore size membranes. Before injections, they were mixed with 0.5 ml of Complete Freund’s Adjuvant for the first injection (incomplete Freund’s Adjuvant in subsequent injections) and the mixes were administered subcutaneously in four to six sites on the back of the animals. Injections were repeated at 21 days intervals and antibody titer was tested after the second injection (ear bleeding) to decide if another boost was needed. Immunization experiments in rabbits had the approval of the Ethical Committee of University of Valencia, where the experiments were carried out, and the corresponding authorization of the Government of the
Comunitat Valenciana (2014/VSC/PEA/00197). Presence of P40 and P75 was analyzed both by Dot-Blot and Western-Blot as previously described (Bäuerl et al., 2010). Briefly, protein extracts or supernatants were analyzed by SDS-PAGE and electro-transferred to a Hybond nitrocellulose membrane (GE Healthcare Life Sciences, Uppsala, Sweden). For Dot-blot analysis, 2 µl of bacterial cell extract or cell culture supernatant were manually spotted on the nitrocellulose membrane. Then the membranes were blocked in 5 % non-fat dry milk solution and incubated either with rabbit polyclonal anti-P40N or anti-P75N (1:5000 in 5% non-fat dry milk) or when indicated with rabbit polyclonal anti-P40T and anti-P75T (Yan et al., 2007) (1:10000 in 5% non-fat dry milk). After incubation with HRP-conjugated goat anti-rabbit antibody (GE-Healthcare), detection was performed using the chemiluminescent ECL kit (GE HealthCare). Light emission from the blots was detected and quantified with a LAS-1000 image analysis system (Fuji, Tokyo, Japan).

Microscopic and Immunohistochemistry Techniques

The cellular distribution of P40 and P75 in L. casei BL23 and L. rhamnosus GG was first determined by microscopic epifluorescence observation. To this purpose, bacterial suspensions were fixed and treated with rabbit anti-P40N and anti-P75N polyclonal antibodies. Subsequently, the preparations were treated with fluorescein-coupled goat anti-rabbit antibodies following a standard protocol. Briefly, bacteria were grown overnight, and then diluted 1:100 in MRS and incubated for 1 h. Bacteria were fixed by placing 5 µl of the bacterial suspension on a slide, where they were mixed with 10 µl of 4% para-formaldehyde (PFA), air dried and then gently heated. Bacteria were subsequently covered with 50 µl blocking solution (4% BSA in PBS) for 1 h at room temperature (RT). The liquid was removed gently with a pipette and bacteria were covered with 50 µl of the primary antibody solution (anti-P40N and anti-P75N) diluted 1:400 in PBS supplemented with 1% BSA and incubated for 1 h at RT. Then bacteria were washed six times with 50 µl PBS, covered with 50 µl diluted (1:500) Alexa Fluor 488 cross-absorbed goat anti-rabbit secondary antibody (Invitrogen, United States) solution in 1% BSA/PBS and incubated for 1 h at RT. Cells were washed 6 times with 50 µl PBS in the dark and fixed again with 50 µl 4% PFA for 10 min at RT in the dark. Immunolabelled bacteria were dried and covered with a drop (5 µl) of mounting medium (PBS/glycerol 1:1) before lying on the cover slide. Samples could be stored at 4°C in the dark for several weeks. Epifluorescence digital images were acquired with an Eclipse 90i Nikon microscope (Nikon Corporation, Japan) at 1000-fold magnification, equipped with a digital camera (Nikon DS-5Mc). Bright field and phase contrast digital images were also acquired with the same Eclipse 90i Nikon microscope (Nikon Corporation, Japan) and camera (Nikon DS-5Mc).

For immunohistochemistry in transmission electron microscopy, bacterial suspensions in PBS were included in 1% low-gelling temperature agarose and fixed overnight with Karnovsky fixative (2.5% paraformaldehyde, 0.5% glutaraldehyde in 0.2M cacodylate buffer pH 7.4). Samples were subsequently washed three times with PBS, post-fixed with 2% osmium tetroxide for 2 h, washed and dehydrated in ascending ethanol concentrations from 30 to 90% and embedded in LR-White resin. After polymerization of the resin, ultrathin sections were obtained with an UC6 Ultramicrotome (Leica Microsystems, Germany) with a diamond knife (Diatome, Switzerland), and placed in nickel grids. Grids were incubated for 3 h with anti-P40N and anti-P75N antibodies diluted 1:50 in PBS for 2 h. After washing in Tris 50 mM pH 8.0 containing 1% gelatin and 0.5% Tween-20 for 10 min, grids were incubated for 1 h with a 10 nm gold conjugate goat anti-rabbit antibody diluted in Tris 50 mM pH 8.0 with 1% BSA and 0.5% Tween-20 for 2 h. Subsequently, grids were washed with PBS, fixed with 2.5% glutaraldehyde in PBS for 2 min and washed in distilled water. Finally, samples were stained with 2% uranyl acetate and lead citrate for 25 and 12 min, respectively. Samples were examined in a transmission electron microscope Jeol JEM-1010 (80 kV) with an AMT RX80 (8 Mpx) digital camera.

Sequence Analysis

Sequences of P40 and P75 homologs were retrieved from the microbial genome repository at the NCBI by PSI-BLAST (Altschul et al., 1997) using as query sequences the L. casei BL23 P40 (Acc. No. CAQ65155) and P75 (Acc. No. CAQ65403) protein sequences. Domain analysis was performed using the tools implemented at the NCBI Blast site1. The resulting datasets were aligned with M-Coffee (Notredame et al., 2000) at the T-Coffee server2 with default settings. Positions of uncertain homology and gaps were removed using GBLOCKS (Castresana, 2000) at the Gblocks server3 allowing smaller final blocks and less strict flanking positions. Redundant sequences were removed by using the EMBOSS suite Skipredundant tool (Rice et al., 2000) with a percentage sequence identity redundancy threshold of 98%. The best-fit models of amino acid substitution were selected by using the tool implemented in MEGA (ver. 7.0.18) (Kumar et al., 2016). The models selected were used to obtain maximum likelihood trees using PhyML ver. 3.0 (Guindon and Gascuel, 2003) at the PhyML server4. Bootstrap support values were obtained from 1000 pseudorandom replicates.

---

1https://blast.ncbi.nlm.nih.gov/Blast.cgi
2http://tcoffee.crg.cat
3http://molevol.cmima.csic.es/castresana/Gblocks_server.html
4http://www.atgc-montpellier.fr/phyml
RESULTS

Features of L. casei BL23 P40 and P75 Encoding Sequences

P40 and P75 proteins belong to a family of proteins carrying a C-terminal CHAP (NlpC/P60 superfamily) domain. At the N-terminal region, P40 has abundant positively charged amino acids included in a CwI0 (COG3883) domain of the cl25603 superfamily characteristic of PGN hydrolases. Between both domains there is a serine, alanine, threonine rich spacer region with undefined similarity (Supplementary Figures S1A,B). Domains search of P75 also rendered a C-terminal CHAP region, but the N-terminus is rich in polar amino acids without known conserved domains. This protein also has a middle region with abundant serine and alanine residues and, in addition, threonine and proline rich areas (Figure 1A).

Many of the protein sequences analyzed below show variable intermediate regions next to the CHAP domain. Tandem repeat (TR) search in the genes encoding P40 and P75 in L. casei BL23 (cmuA, cmuB) and L. rhamnosus GG (mspB, mspA) resulted in the localization of two consecutive 18 nt repeats or minisatellite DNA regions (AGTGCTGCCGCAGAATCC and ACCCGACTCCTGCACCA) in the middle region of L. casei BL23 cmuB. These TR are not present in L. rhamnosus mspA (Supplementary Figure S2), neither in L. zae (AZCT01000007.1:13555-14967) or L. casei subsp casei ATCC 393 (AP012544.1:225479-226891) homologous genes. These TR, or microsatellites, were found in a large number of homologous genes in L. paracasei strains, including L. casei ATCC 334 and BL23. The vast majority of strains included in the L. casei/paracasei group had identical TR structure as BL23 with a few interesting exceptions. Three instead of two copies of the first TR were found in the strains L. casei Zhang, L. casei LOCK919, L. paracasei ZFM54 and L. paracasei NNRRL B, and a total deletion in L. paracasei JCM 8130. Two sequence divergences for the downstream second microsatellite were found showing a total deletion in L. paracasei TMW 1.1434 and an additional third copy in L. paracasei JCM 8130 (Figure 1B and Supplementary Table S3). This variability in the intermediate region of P75 in L. casei/paracasei conformed four alleles, cmuB1, cmuB2, cmuB3, and cmuB4 (Figure 1B). These genetic variations did not affect the phylogeny built with protein sequences shown below.

P40 and P75 Phylogeny

As in other bacterial groups, Lactobacillus and closely related bacteria host several proteins with a conserved C-terminal NlpC/P60 domain, so that analyzing the annotated genomes of representative strains of Lactobacillus and Lactococcus four sequences were found in L. casei BL23, three in L. rhamnosus GG, three in Lactococcus lactis subsp. cremoris MG1363, five in L. plantarum WCF51, five in Lactobacillus delbrueckii subsp. bulgaricus ATCC BAA-365, three in Lactobacillus brevis ATCC BAA-365 and just one in Lactobacillus gasseri ATCC 33323 (Supplementary Table S1).

Frequently, the conserved domains of the proteins are selected to obtain sound phylogenetic relationships as shown by other studies already performed with proteins of this family (Anantharaman and Aravind, 2003; Bateman and Rawlings, 2003; Layec et al., 2008). However, the NlpC/P60 superfamily domain covers a small part of the protein at the C-terminus, for which important information could be missed. This is particularly important if we try to find the most similar protein sequences -with muramidase activity-, but also aim to investigate other biological functions.

Proteins of the P40 cluster were identified as such when having the same domain composition and a conserved sequence homology on at least 80% of the whole protein length. In general, we found similar proteins in species closely related to the L. casei, L. paracasei, L. zae and L. rhamnosus taxonomic group (Figure 2A). P40 homologs were also found in species of the families Enterococcaceae and Streptococcaceae. It seems interesting that the NlpC/P60 domain of P40 homologs shared similarity to the corresponding domain of the surface antigen of E. faecium SagA (Teng et al., 2003). In Lactobacillus, true P40 proteins are found in strains of the L. casei, L. sakei and Lactobacillus salivarius groups as defined by Salvetti et al. (2018). Within the L. casei group, P40 homologs are absent in Lactobacillus brentiae, Lactobacillus cameliae, Lactobacillus pantheris, Lactobacillus saniviri, Lactobacillus sharpeae, and Lactobacillus thailandensis. The genes harbored by strains of L. casei, L. paracasei, L. rhamnosus and L. zae constitute a well supported group suggesting that a P40 encoding gene was present in the common ancestor of these species. Apart from those, Lactobacillus nasuensis and Lactobacillus manihotivorans, which also belong to the L. casei group, encode P40 homologs but the lack of support of their phylogenetic positions make it difficult to determine their origin. The four species of the L. sakei group encode P40 homologs whereas only two species of the L. salivarius group encode P40 homologs. Remarkably, many strains of L. sakei encode two paralog proteins, one of them clustering with the other sequences from the L. sakei group, as expected, but the other one in a well-supported cluster with the sequences of Lactobacillus murinus and Lactobacillus animalis. The limited distribution of P40 encoding genes within species of the L. salivarius group may suggest a possible horizontal gene transfer of this gene from L. sakei to the last common ancestor of these two closely related species. However, the presence of two paralogs encoding P40 is quite anomalous, since no other species harbor two P40 encoding genes. Additional data would be required to settle this point. The shape of this tree does not suggest that P40 could derive from horizontal transfer from streptococci or enterococci to lactobacilli or vice versa. As it is, we hypothesize that a P40 encoding gene was present in the last common ancestor of Enterococcaceae, Lactobacillaceae and Streptococcaceae. The current distribution of these genes would be mostly explained by lineage-specific gene losses.

When applied to P75 proteins, the criterion of selection used above for P40 proteins was only fulfilled by P75 homologs of the L. casei and L. sakei groups. Therefore, to gain insight into the phylogenetic position of P75, all sequences retrieved in the PSI-BLAST search from species belonging to Lactobacillaceae were included in the analysis. Proteins like P75 do not have a defined consensus domain at the amino terminal region.
and, in fact, there is a great dispersion in sequence among them, hence, the C-terminal NlpC/P60 domain will have a fundamental weight in the phylogenetic analysis. On the basis of the homology of the NlpC/P60 domain, many other proteins could have been included in the comparison, like some enterococcal proteins, P45 and TrsG from *L. casei* and other species, but this did not add any significant information to existing trees (Anantharaman and Aravind, 2003). As previously observed for P40, P75 was found within the *L. casei* group in *L. casei*, *L. paracasei*, *L. rhamnosus* and *L. zeae*.* L. nasuensis* and *L. cameliae* also encoded putative P75 proteins (although similarity is limited to the N-terminal and C-terminal parts) whereas the corresponding gene is absent in *L. brantae*, *L. pantheris*, *L. saniviri*, *L. sharpeae*, and *L. thailandensis*.

![FIGURE 1](https://www.frontiersin.org/wit...2.png)
FIGURE 2 | Maximum likelihood phylogenetic trees of representative P40 (A) and P75 (B) protein sequences. The trees have been arbitrarily rooted to facilitate its visualization. Support values higher than 750 for the bootstrap analysis are indicated. Protein sequence codes are supplied in Supplementary Tables S3.1, S3.2.
Cloning and Purification of P40, P75 and N-Terminal Domains

The complete mature proteins (P40T and P75T) were overexpressed in *E. coli* BL21 by cloning the corresponding genes (LCABL_00230, LCABL_02770) of *L. casei* BL23 in the expression vector pQE80E. PCR oligonucleotides were designed to allow in frame fusion of the gene encoding the mature protein (excluding the secretion signal) to the RGS-His epitope encoding region in pQE80E.

NlpC/P60 domains are conserved in proteins of different bacterial taxa. Hence, in order to obtain highly specific polyclonal antibodies recognizing P40 and P75, N-domains of both proteins were expressed in *E. coli* and purified. P40 N-domain (P40N) included the Cwl0(COG3883) domain (residues 74 to 211) and in the case of P75, the N-domain spanned the complete peptide sequence upstream NlpC/P60 domain. Because there was no domain description in databases associated to that region, the N-terminal region, as suggested by structural predictions (Supplementary Figure S4). The homologous P75 protein of *L. rhamnosus* GG was shown to be glycosylated, which could influence its migration in PAGE, however, *L. casei* BL23 was shown to be glycosylated, which could influence its migration in PAGE, however, *L. casei* BL23 homologs were also present in all species of the *L. sakei* group, as previously observed for P40 (Figure 2A). Both groups form a well-supported cluster. However, phylogenomic analyses (Sun et al., 2015; Salvetti et al., 2018) do not place these two groups as sister groups and the presence of more distantly related P75 proteins in other groups of lactobacilli makes it difficult to accept that evolution of P75 in lactobacilli is explained by vertical inheritance and lineage-specific gene losses. For example, the *L. perolens* group is a sister group of the *L. casei* group, however, their corresponding sequences are more distantly related than those of the *L. sakei* group (Figure 2B). Unfortunately, lack of support in deeper nodes of the tree precludes establishing reliably the relationships between these clusters. Interestingly, *L. paracasei* Lpp122 harbors a paralog encoding a putative P75 protein that clusters with the corresponding gene of *L. manihotivorans*. These two sequences form a well-supported cluster with sequences of species of the *L. perolens* group (Figure 2B). This arrangement suggests a horizontal gene transfer from a member of the *L. perolens* group. In summary, the analysis indicates that P75 proteins of *L. casei* and *L. sakei* groups share a common origin but it cannot be ascertained whether vertical inheritance or horizontal gene transfer would explain the current distribution of these genes.

Localization, Mutant Complementation and Lytic Activity of P40 and P75

Immunofluorescence observations showed that P40 and P75 were predominantly detected at the septum and polar regions in *L. casei* BL23 (Figure 4). When we tested P40 in the closely related bacterium *L. rhamnosus* GG, it was localized apically but a clear observation was difficult as it was possibly associated to the exopolysaccharide secreted by this strain, as can be observed both in epifluorescence and TEM (Figure 4).

In a previous work, P40 (BL305) and P75 (BL306) defective *L. casei* BL23 mutant strains were obtained (Bäuerl et al., 2010). Both mutants had distinctive morphologies associated to a possibly abnormal cell division process. Cell suspensions of both mutants were incubated with P40 and P75 (final concentration of 50 µg/ml) for 0 to 3 h and observed under the microscope (Figure 5). After only 3 h incubation with P40, apparent cell morphology of mutant BL305 reverted significantly to the wild type, and although BL306 typical extremely long rods persisted after 3 h incubation with purified P75, phase contrast evidenced the primordial formation of septa. These complementation assays and their localization suggest that P40 and P75 are involved in septation and separation of daughter cells in *L. casei* BL23.

Given the relative similarity of P40 and P75 to putative bacterial lysins (see below) and since these two proteins are found in the extracellular medium, their (auto)lytic activity was assayed separately and together in *L. casei* BL23 and against gram positive competitors/pathogens such as *ENTEROCOCCUS FAECALIS*, *LISTERIA MONOCYTGENES* and *STAPHYLOCOCCUS AUREUS*. Different conditions were used but no significant effect could be found in cell suspensions adjusted at different pH values even when P40 and P75 treatment was followed by a detergent (Triton x100) treatment (Supplementary Figure S5). The only noticeable effect...
was an increase in OD promoted by P75 in *L. casei* BL23 and by both enzymes together in *L. casei* and *E. faecalis*, suggesting that they may promote cell division, in agreement with the microscopic observations (Figures 4, 5).

### Detection of P40 and P75 in the *L. casei/paracasei/rhamnosus* Taxon and Other Lactobacilli

Homology searches indicated that most Firmicutes, also the *Lactobacillaceae* family, normally harbor more than one gene encoding proteins containing the NlpC/P60(CHAP) C-terminal domain (Layec et al., 2008). Our protein comparisons suggested that amino acid sequences of this group rendered distinctive clusters that grouped P40 and P75 separately from other CHAP-containing proteins and also found that the C-terminal NlpC/P60 domains were better conserved than the N-terminal part (see above). For this reason, N-terminal domains of both proteins were cloned as His-tag fusions and purified to obtain antibodies that were used in order to specifically recognize P40 and P75 from *L. casei/rhamnosus* strains. Initial results were very encouraging because a single band was detected with each antibody in culture supernatants of *L. casei* BL23 (Figure 6A). Then, culture supernatants of the *Lactobacillus* laboratory’s collection (Table 1) were tested by Western blot with anti-P40N and anti-P75N antibodies. Most lactobacilli belonging to *L. casei*, *L. zeae* and *L. rhamnosus* species showed Western blot bands that coincided with the P40 and P75 migrations in SDS-PAGE – with the exception of *L. casei* subsp. *tolerans* BL134-. In a few cases, more than one band or a band of smaller molecular size were detected with anti-P40N (BL87, BL90) or anti-P75N (BL199, BL212). Strain *L. casei* subsp. *pseudoplantarum* BL205 did not show any band and P40 was missing under the conditions assayed in *L. casei* BL6 and *L. rhamnosus* BL103. Inspection of the genome of *L. casei* BL6 (ATCC393) reveals however, that this strain harbors a gene encoding a “CHAP domain-containing protein” (WP_025013825.1) with 85% identity to BL23 P40.

---

**FIGURE 4** | Localization of P40 in *L. casei* BL23 and *L. rhamnosus* GG with anti-P40N antibodies, and P75 in *L. casei* BL23 with anti-P75N antibodies. (A–C) Show random epifluorescence microscopic fields of Alexa immune labeled P40 on *L. casei* BL23 and *L. rhamnosus* BL376 (LGG); (E) of P75 in *L. casei* BL23; and (B,D) show TEM of *L. casei* BL23 and *L. rhamnosus* BL376 (LGG) subject to immunogold staining of P40.
FIGURE 5 | Mutant complementation with purified P40 and P75. Morphological observation by phase contrast transmission microscopy (objective × 100) of L. casei BL23 (A), L. casei P40 knock down mutant, cmuA (BL305) incubated for 0 h (B) and 3 h (D) with purified P40 (final concentration of 50 µg/ml), and L. casei P75 knock down mutant, cmuB (BL306) incubated for 0 h (C) and 3 h (E) with purified P75 (final concentration of 50 µg/ml) and as indicated in the panels. (C,E) Show enlarged details for a better distinction of nascent septa.

This confirmed that the antibodies obtained here against the N-terminal domain of P40 and P75 gained specificity.

PCR Detection of P40 and P75 Encoding Genes in Lactobacilli

The similarity at DNA level of the corresponding genes in the species of L. casei, L. paracasei, L. zeae and L. rhamnosus allowed the design of oligonucleotides targeted to different conserved sequences (Supplementary Tables S4, S5 and Figures 6A,B) that were used to test the specific amplification of cmuA and cmuB by PCR. The oligonucleotide combinations that rendered reproducible positive bands in all L. casei/paracasei/rhamnosus strains of the laboratory’s collection are described
in Table 2. The only exception was strain \textit{L. casei} subsp tolerans BL134 as occurred in Western blots. P40-encoding gene could be unequivocally amplified with primers P40carhf1/P40carhr yielding amplicons of about 278 nt (Table 2). The P75 encoding gene was identified by PCR with P75carhf and P75carhr obtaining a band of 138 nt. In other \textit{Lactobacillus} species both genes were only amplified in \textit{L. sakei} BL80. The rest of \textit{Lactobacillus} strains tested yielded no amplification bands, excepting the P75 specific PCR in \textit{L. plantarum} BL35 and \textit{Lactobacillus delbrueckii} subsp \textit{lactis} BL89.

### Detection of P40 and P75 in Fermented Dairy Products

Different commercial products found in local supermarkets that contained in their ingredients strains of \textit{L. casei} were used to detect P40 and P75. In order to avoid possible interference of gelling agents, five drinkable products were selected (D, C, J, H, N) and their clear fraction and pellets were subjected to Western blot (Figure 7). P40 could only be detected in the supernatant of one of the products (H) while P75 was found in four of them (D, J, H, N). Four other fermented dairy products that did not contain \textit{L. casei} were also tested but did not show any band on both assays (data not shown). Binding to the food matrix components or instability of P40 during the long shelf life of those products (at least 28 days) could explain why it was hardly detected.

### DISCUSSION

Proteins P40 and P75 were first described in the biologically active fraction of \textit{L. rhamnosus} GG conditioned medium that

---

**FIGURE 6** | Diagram summarizing immunoblot signals (Western blot) obtained with anti-P40N and anti-P75N from overnight culture supernatants of the designated strains. (A) Shows a group of strains belonging to the taxon \textit{Lactobacillus} casei/paracasei/rhamnosus and (B) representative strains of the genus \textit{Lactobacillus} from the laboratory’s collection.

| A | Western blot | PCR detection | B | Western blot | PCR detection |
|---|---|---|---|---|---|
| P40 | P75 | P40 | P75 | P40 | P75 | P40 | P75 |
| BL6 \textit{L. casei} | + | + | BL5 \textit{L. acidophilus} | | | |
| BL23 \textit{L. casei} | + | + | BL7 \textit{L. brevis} | | | |
| BL28 \textit{L. casei} | + | + | BL8 \textit{L. plantarum} | | | |
| BL32 \textit{L. casei} | + | + | BL13 \textit{L. sakei} | | | |
| BL81 \textit{L. casei} | + | + | BL14 \textit{L. curvatus} | | | |
| BL82 \textit{L. casei} | + | + | BL15 \textit{L. alimentarius} | | | |
| BL83 \textit{L. casei} | + | + | BL26 \textit{L. plantarum} | | | |
| BL86 \textit{L. casei} | + | + | BL33 \textit{L. delbrueckii} subsp \textit{bulgaricus} | | | |
| BL87 \textit{L. casei} | + | + | BL34 \textit{L. fermentum} | | | |
| BL90 \textit{L. casei} | + | + | BL35 \textit{L. pentosus} | | | |
| BL91 \textit{L. casei} | + | + | BL36 \textit{L. brevis} | | | |
| BL95 \textit{L. zea} | + | + | BL47 \textit{L. pentosus} | | | |
| BL101 \textit{L. casei} | + | + | BL73 \textit{L. acidophilus} | | | |
| BL134 \textit{L. paracasei} subsp tolerans | + | + | BL74 \textit{L. acidophilus} | | | |
| BL139 \textit{L. casei} | + | + | BL75 \textit{L. acidophilus} | | | |
| BL199 \textit{L. casei} | + | + | BL76 \textit{L. murinus} | | | |
| BL203 \textit{L. casei} | + | + | BL80 \textit{L. sakei} | + | + | |
| BL206 \textit{L. casei} | + | + | BL84 \textit{L. brevis} | + | + | |
| BL212 \textit{L. casei} | + | + | BL89 \textit{L. delbrueckii} subsp \textit{lactis} | | | |
| BL216 \textit{L. casei} | + | + | BL130 \textit{L. coryniformis} | | | |
| BL227 \textit{L. casei} | + | + | BL131 \textit{L. coryniformis} | | | |
| BL229 \textit{L. casei} | + | + | BL132 \textit{L. plantarum} | | | |
| BL312 \textit{L. paracasei} | + | + | BL135 \textit{L. agilis} | | | |
| BL358 \textit{L. casei} | + | + | BL156 \textit{L. jaccarina} | | | |
| BL1 \textit{L. rhamnosus} | + | + | BL157 \textit{L. ruminis} | | | |
| BL2 \textit{L. rhamnosus} | + | + | BL158 \textit{L. salivarius} | | | |
| BL3 \textit{L. rhamnosus} | + | + | BL159 \textit{L. salivarius} | | | |
| BL102 \textit{L. rhamnosus} | + | + | BL160 \textit{L. delbrueckii} subsp \textit{bulgaricus} | | | |
| BL103 \textit{L. rhamnosus} | + | + | BL166 \textit{L. plantarum} | | | |
| BL327 \textit{L. rhamnosus} | + | + | BL202 \textit{L. acidophilus} | | | |
| BL367 \textit{L. rhamnosus} | + | + | BL207 \textit{L. helveticus} | | | |
| BL376 \textit{L. rhamnosus} | + | + | BL221 \textit{L. crispatus} | | | |
| BL377 \textit{L. rhamnosus} | + | + | BL223 \textit{L. gasseri} | | | |
| BL378 \textit{L. rhamnosus} | + | + | BL228 \textit{L. acidophilus} | | | |
| | | | BL231 \textit{L. acidophilus} | | | |
P75 homologs, so the NlpC/P60 domain had a greater weight in loss explaining its current distribution. The case of P75 was very... and Lactobacillaceae gene was present in the last common ancestor of phylogenetic analysis suggests that an ancestral P40 encoding...

P40 and P75 belong to an important group of cell wall degrading enzymes with different N-terminal activities may constitute an additional factor contributing to their functional properties in vivo. Both proteins have a C-terminal NlpC/P60 domain, a widely distributed domain in bacteria and bacteriophages, which is also present even in eukaryotic regulators, proteins of poxviruses and animal RNA viruses [for reviews, see (Anantharaman and Aravind, 2003; Bateman and Rawlings, 2003)]. P40 and P75 belong to an important group of cell wall degrading enzymes with different N-terminal regions. Layec et al. (2008) described in Firmicutes 12 different subgroups. P40 proteins found in L. casei, L. paracasei, L. zae and L. rhamnosus constitute a well-supported cluster. Taxonomically close species, like those included in the L. sakei and L. salivarius clusters (Salvetti et al., 2018) also encode P40 homologs. The phylogenetic analysis suggests that an ancestral P40 encoding gene was present in the last common ancestor of Enterococcaceae, Lactobacillaceae and Streptococccae, with lineage-specific gene loss explaining its current distribution. The case of P75 was very different, as there is not a clearly defined N-terminal domain in P75 homologs, so the NlpC/P60 domain had a greater weight in the analysis. P75 related proteins were only present in the L. casei and L. sakei groups, but there was lack of support for deep nodes in those taxonomically close bacterial groups and in addition, L. perolens P75 was a sister group of the L. casei group. This analysis indicated that P75 proteins of L. casei and L. sakei groups share a common origin, but the results obtained did not allow obtaining a clear picture of their evolutionary history. When analyzing the nucleotide sequence of cmuB two highly conserved tandem repeats of 18 nucleotides were clearly identified with four strain-specific copy number variations (alleles: cmuB1, cmuB2, cmuB3, cmuB4). A similar case of TR in a PGN hydrolase was described in the cse gene of S. thermophilus (Borges et al., 2006), although this was a more complex scenario with a mosaic of 11 different TR sequences grouped in larger repeated sequence blocks, with additional degrees of sequence degeneration. As in the case of S. thermophilus cse, all alleles are functional suggesting that TR protein sequences in CmuB are not essential for the activity of the enzyme, acting as spacers between the C-terminal NlpC/P60 domain and other so far uncharacterized domain(s) at the N-terminal region. TR sequences are prone to insertions and deletions (indels) providing flexibility for functional adaptation. They are frequent in surface exposed proteins that interact directly with host structures (Moxon et al., 2006), like epitopes or adhesion/invasion elements of epithelial pathogens (Zhou et al., 2014). This genetic variability translated to protein sequences only affected strains within the L. casei/paracasei groups and had no relevance in the phylogenetic analysis performed. The presence of the microsatellites in cmuB was only detected in strains considered as L. paracasei, this discovery constitutes the basis of an interesting taxonomical feature to be further explored.

In previous studies, P40 and P75 were recognized by anti-P40 antibodies in a large number of L. casei/paracasei/rhamnosus strains (Yan et al., 2007; Bäuerl et al., 2010) suggesting those antibodies had a broad specificity possibly recognizing P40 and P75 similar surface epitopes. It is shown here that antibody specificity was much improved by immunizing rabbits with the purified N-terminal regions of both proteins. The specific detection of P40 and P75 in L. casei/paracasei/rhamnosus strains was successfully achieved using polyclonal anti-P40N and anti-P75N rabbit antibodies obtained, and not in other Lactobacillus species, with only a few exceptions. Absence of Western blot signal in some L. casei/paracasei/rhamnosus strains may be due to the culture conditions used, or to the fact that homologous genes are not found in some strains, possibly because the activity of those proteins may be efficiently replaced and complemented by other NlpC/P60 proteins. Frequently, Firmicutes genomes contain more than two genes encoding NlpC/P60 PGN hydrolases. In addition to P40 and P75 (WP_003572828.1, WP_012490875.1), L. casei BL23 genome contains genes encoding at least two additional NlpC/P60 C-terminal domain proteins (WP_012491782.1, WP_012491062.1) not mentioned in our study. Lactobacillus plantarum WCFS1, another well known probiotic strain, encodes five NlpC/P60 C-terminal putative proteins that do not belong to the P40 or P75 families (Supplementary Table S1), four of them (LytA, LytB, LytC, LytD) recently studied in detail (Duchêne et al., 2019).
In the case of P75, subcloning of the N-terminal domain showed that this region is responsible for the unusual migration in SDS-PAGE of the whole protein. P40 and P75 have always been detected in the culture supernatants. This work showed that P40 and P75 are localized or at least bind to the polar regions in BL23, and that purified proteins would complement specific mutants where cmuA and cmuB had been inactivated. Microscopic images (Figures 4, 5) suggest that P40 may participate in early cell division stages and perimetric cell growth and P75 may be required in septum formation and cell separation. Lytic activity has been described in other NlpC/P60 proteins in lactobacilli (Liu et al., 2015), but P40 and P75 did not show such activity in our experiments suggesting that cell wall remodeling does not necessarily imply a lytic or autolytic activity (Vermassen et al., 2019).

CONCLUSION
This work has defined genetic characteristics of the genes encoding P40 and P75 in L. casei/paracasei strains. These antiapoptotic proteins are actually required by the bacterial cells for growth and cell division and the phylogenetic analysis suggested that they are found in L. casei/paracasei/rhamnosus (L. casei phylogenomic group) and close species like L. sakei. P40 likely evolved by vertical inheritance, while in some cases horizontal transfer may have also participated in the inheritance of P75 proteins. In addition to these findings, we have developed tools that specifically recognize L. casei phylogenomic group, such as the anti-P40N and anti-P75N antibodies detecting P40 and P75 in L. casei/paracasei/rhamnosus strains, also suitable for commercial fermented dairy products, and PCR primers that specifically recognize L. casei/paracasei/rhamnosus strains.

DATA AVAILABILITY
All datasets for this study are included in the manuscript and the Supplementary Files.

ETHICS STATEMENT
This study was carried out in accordance with the recommendations of the European Union Law and 2010/63/EU and the Spanish Government RD 53/2013 on the protection of animals used for scientific purposes, name of committee. The protocol was approved by the Ethical Committee of University of Valencia and had the corresponding authorization of the Government of the Comunitat Valenciana (2014/VSC/PEA/00197).

AUTHOR CONTRIBUTIONS
All authors have contributed significantly to the development of this work. CB cloned, purified, and produced the antibodies and initiated the western blot series. GA performed the western blots, microscopic preparations, and DNA sequence alignments. SS-C, AM-B, and NN-L initiated all the cloning series until the proteins could be expressed in E. coli and initiated the phylogenetic analysis and western blots. JC-M contribution was crucial to achieve cloning and microscope preparations. MZ-C reorganized and performed the final phylogenetic analysis and evolutionary conclusions. SS contributed to the supervision and manuscript preparation. GP-M supervised and financed all the experiments and carried out the genetic analysis, writing up and figure preparation for the submission.

FUNDING
We wish to thank the Grants of the Spanish Ministry of Science and Universities AGL2013-47420-R and AGL2015-70487-P. GA and SS wish to thank the grant from the L. N. Gumilyov Eurasian National University. We acknowledge support of the publication fee by the CSIC Open Access Publication Support Initiative through its Unit of Information Resources for Research (URICI).

ACKNOWLEDGMENTS
We would like to acknowledge M. T. Mínguez for her valuable technical assistance in the preparation of samples and use of the transmission electron microscope at the S.C.S.I.E., University of Valencia.

SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.01420/full#supplementary-material

REFERENCES
Acedo Felix, E., and Perez Martinez, G. (2003). Significant differences between Lactobacillus casei subsp. casei ATCC 393T and a commonly used plasmid-cured derivative revealed by a polyphasic study. Int. J. Syst. Evol. Microbiol. 53, 67–75. doi: 10.1099/ijs.0.02325-0
Altschul, S. F., Madden, T. L., Schäffer, A. A., Zhang, J., Zhang, Z., Miller, W., et al. (1997). Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. Nucleic Acids Res. 25, 3389–3402. doi: 10.1093/nar/25.17.3389

Anantharaman, V., and Aravind, L. (2003). Evolutionary history, structural features and biochemical diversity of the NlpC/P60 superfamily of enzymes. Genome Biol. 4, 1–12.
Aramini, J. M., Rossi, P., Huang, Y. J., Zhao, L., Jiang, M., Maglaqui, M., et al. (2008). Solution NMR structure of the NlpC/P60 domain of lipoprotein Spr from Escherichia coli: structural evidence for a novel cysteine peptidase catalytic triad. Biochemistry 47, 9715–9717. doi: 10.1021/bi8010779
Bateman, A., and Rawlings, N. D. (2003). The CHAP domain: a large family of amidas including GSP amidase and peptidoglycan hydrolases.
Nucleic acid Techniques in

Lane, D. J. (1991). “16S/23S rRNA sequencing,” in

Kumar, S., Stecher, G., and Tamura, K. (2016). MEGA7: molecular evolutionary

Leenaars, M., and Hendriksen, C. F. M. (2005). Critical steps in the production of

Castresana, J. (2000). Selection of conserved blocks from multiple alignments for

L. casei

Bäuerl et al.

Liu, S., Rich, J. O., and Anderson, A. (2015). Antibacterial activity of a cell wall

salicylate from Lactobacillus paracasei NRIIL B-50314 produced by recombinant Bacillus megaterium J. Ind. Microbiol. Biotechnol. 42, 229–235. doi: 10.1007/s10295-014-1557-6

Moxon, R., Bayliss, C., and Hood, D. (2006). Bacterial contingency loci: the role of simple sequence dna repeats in bacterial adaptation. Annu. Rev. Genet. 40, 307–333. doi: 10.1146/annurev.genet.40.110405.09442

Notredame, C., Higgins, D. G., and Heringa, J. (2000). T-coffee: a novel method for fast and accurate multiple sequence alignment. Edited by J. Thornton. J. Mol. Biol. 302, 205–217. doi: 10.1006/jmbi.2000.4042

Oechslin, F., Daraspe, J., Giddey, M., Moreillon, P., and Resch, G. (2013). In vitro characterization of PlySK1249, a novel phage lysin, and assessment of its antibacterial activity in a mouse model of Streptococcus agalactiae bacteremia. Antimicrob. Agents Chemother. 57, 6276–6283. doi: 10.1128/AAC.01701-13

Park, J.-H., Lee, Y.-S., Lim, Y.-K., Kwon, S.-H., Lee, C.-U., and Yoon, B.-S. (2000). Specific binding of recombinant Listeria monocytes p60 protein to Caco-2 cells. FEMS Microbiol. Lett. 186, 35–40. doi: 10.1016/s0168-0561(00)0112-9

Rangan, K. J., Pedicord, V. A., Wang, Y.-C., Kim, B., Lu, Y., Shaham, S., et al. (2016). A secreted bacterial peptidoglycan hydrolase enhances tolerance to enteric pathogens. Science 353, 1434–1437. doi: 10.1126/science.aaf3552

Regulski, K., Courtin, P., Maynard, M., Claes, I. J. J., Lebeer, S., Vanderleyden, J., et al. (2012). Analysis of the peptidoglycan hydrolase complement of Lactobacillus casei and characterization of the major y-d-glutamyl- l-lysyl-endopeptidase. PLoS One 7:e32301. doi: 10.1371/journal.pone.0032301

Rice, P., Longden, I., and Bleasby, A. (2000). EMBOSS: the European molecular biology open software suite. Trends Genet. 16, 276–277. doi: 10.1016/s0168-9525(00)02024-2

Salvetti, E., Harris, H. M. B., Felis, G. E., and O’taole, P. W. (2018). Comparative genomics of the genus lactobacillus reveals robust phylogroups that provide the basis for reclassification. Appl. Environ. Microbiol. 84, e0993-18. doi: 10.1128/AEM.00993-18

Sambrook, J., and Russell, D. W. (2001). Molecular Cloning: A Laboratory Manual. New York, NY: Cold Spring Harbor laboratory Press

Sanz-Gaitero, M., Keary, R., Garcia-Doval, C., Coffer, A., and Van Raaij, M. J. (2014). Crystal structure of the lytic CHAP(K) domain of the endolysin LysK from Streptococcus aureus bacteriophage K. Virol. J. 11:133. doi: 10.1186/1743-422X-11-133

Shen, X., Liu, L., Peek, R. M., Acra, S. A., Moore, D. J., Wilson, K. T., et al. (2018). Supplementation of p40, a Lactobacillus rhamnosus GG-derived protein, in early life promotes epidermal growth factor receptor-dependent intestinal developmental and long-term health outcomes. Microsomal Immunol. 11, 1316–1328. doi: 10.1007/s43185-018-0034-3

Sun, Z., Harris, H. M. B., Mccann, A., Guo, C., Argimón, S., Zhang, W., et al. (2015). Expanding the biotechnology potential of lactobacilli through comparative genomics of 213 strains and associated genera. Nat. Commun. 6:3832. doi: 10.1038/ncomms9322

Teng, F., Kawalec, M., Weinstock, G. M., Hryniwicz, W., and Murray, B. E. (2003). An Enterococcus faecium secreted antigen. SagK, exhibits broad-spectrum binding to extracellular matrix proteins and appears essential for E. faecium Growth. Infect. Immum. 71, 5033–5041. doi: 10.1128/IAI.71.9.5033-5041.2003

Vermassen, A., Leroy, S., Talon, R., Provot, C., Popowska, M., and Desvaux, M. (2019). Cell wall hydrolases in bacteria: insight on the diversity of cell wall amidases, glycosidases and peptidases toward peptidoglycan. Front. Microbiol. 10:331. doi: 10.3389/fmicb.2019.00331

Vissewesaran, G. R. R., Leenhouts, K., Van Roosmalen, M., Kok, J., and Buist, G. (2014). Exploiting the peptidoglycan-binding motif, LysM, for medical and industrial applications. Appl. Microbiol. Biotechnol. 98, 4331–4345. doi: 10.1007/s00253-014-5633-7

Wang, L., Cao, H., Liu, L., Wang, B., Walker, W. A., Acra, S. A., et al. (2014). Activation of epidermal growth factor receptor mediates mucin production stimulated by p40, a Lactobacillus rhamnosus GG-derived protein. J. Biol. Chem. 289, 20234–20244. doi: 10.1074/jbc.M114.11553800

Wang, Y., Liu, L., Moore, D. J., Shen, X., Peek, R. M., Acra, S. A., et al. (2016). A LGG-derived protein promotes IgA production through up-regulation of APRIL expression in intestinal epithelial cells. Microsomal Immunol. 10, 373–384. doi: 10.1007/mi.2016.57

Yamaguchi, H., Furuhata, K., Fukushima, T., Yamamoto, H., and Sekiguchi, J. (2004). Characterization of a new Bacillus subtilis peptidoglycan hydrolase gene, yveC (named cwOd), and the enzymatic properties of its encoded protein. J. Biosci. Bioeng. 98, 174–181. doi: 10.1263/jbb.98.174

Yan, F., Cao, H., Cover, T. L., Washington, M. K., Shi, Y., Liu, L., et al. (2011). Colon-specific delivery of a probiotic-derived soluble protein ameliorates intestinal inflammation in mice through an EGF-R-dependent mechanism. J. Clin. Invest. 121, 2242–2253. doi: 10.1172/JCI40331

Yan, F., Cao, H., Cover, T. L., Whitehead, R., Washington, M. K., and Polk, D. B. (2007). Soluble proteins produced by probiotic bacteria regulate intestinal epithelial cell survival and growth. Gastroenterology 132, 562–575. doi: 10.1053/j.gastro.2006.11.022
Yan, F., Liu, L., Dempsey, P. J., Tsai, Y. H., Raines, E. W., Wilson, C. L., et al. (2013). A Lactobacillus rhamnosus GG-derived soluble protein, p40, stimulates ligand release from intestinal epithelial cells to transactivate epidermal growth factor receptor. J. Biol. Chem. 288, 30742–30751. doi: 10.1074/jbc.M113.492397

Yan, F., and Polk, D. B. (2012). Characterization of a probiotic-derived soluble protein which reveals a mechanism of preventive and treatment effects of probiotics on intestinal inflammatory diseases. Gut Microbes 3, 25–28. doi: 10.4161/gmic.19245

Zhou, K., Aertsen, A., and Michiels, C. W. (2014). The role of variable DNA tandem repeats in bacterial adaptation. FEMS Microbiol. Rev. 38, 119–141. doi: 10.1111/1574-6976.12036

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Bäuerl, Abitayeva, Sosa-Carrillo, Mencher-Beltrán, Navarro-Lleó, Coll-Marqués, Zúñiga-Cabrera, Shaikin and Pérez-Martínez. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.