Functional Analysis of Four Bile Salt Hydrolase and Penicillin Acylase Family Members in *Lactobacillus plantarum* WCFS1

Jolanda M. Lambert, ¹ Roger S. Bongers, ¹,² Willem M. de Vos, ¹,³ and Michiel Kleerebezem ¹,²,³,*

¹ TI Food & Nutrition, P.O. Box 557, 6700 AN Wageningen, The Netherlands; ² NIZO food research, P.O. Box 20, 6710 BA Ede, The Netherlands; ³ Wageningen University, Laboratory of Microbiology, P.O. Box 8033, 6700 EJ Wageningen, The Netherlands

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Bile salts play an important role in the digestion of lipids in vertebrates and are synthesized and conjugated to either glycine or taurine in the liver. Following secretion of bile salts into the small intestine, intestinal microbes are capable of deconjugating the glycine or taurine from the bile salts, using an enzyme called bile salt hydrolase (Bsh). Intestinal lactobacilli are regarded as major contributors to bile salt hydrolysis in vivo. Since the bile salt-hydrolyzing strain *Lactobacillus plantarum* WCFS1 was predicted to carry four *bsh* genes (*bsh1, bsh2, bsh3, and bsh4*), the functionality of these *bsh* genes was explored using *Lactococcus lactis* heterologous overexpression and multiple *bsh* deletion strains. Thus, Bsh1 was shown to be responsible for the majority of Bsh activity in *L. plantarum* WCFS1. In addition, *bsh1* of *L. plantarum* WCFS1 was shown to be involved in conferring tolerance to specific bile salts (i.e., glycocholic acid). Northern blot analysis established that *bsh1, bsh2, bsh3*, and *bsh4* are all expressed in *L. plantarum* WCFS1 during the exponential growth phase. Following biodiversity analysis, *bsh1* appeared to be the only *bsh* homologue that was variable among *L. plantarum* strains; furthermore, the presence of *bsh1* correlated with the presence of Bsh activity, suggesting that Bsh1 is commonly responsible for Bsh activity in *L. plantarum* strains. The fact that *bsh2, bsh3*, and *bsh4* genes appeared to be conserved among *L. plantarum* strains suggests an important role of these genes in the physiology and lifestyle of the species *L. plantarum*. Analysis of these additional *bsh*-like genes in *L. plantarum* WCFS1 suggests that they might encode penicillin acylase rather than Bsh activity, indicating their implication in the conversion of substrates other than bile acids in the natural habitat.

*Corresponding author. Mailing address: NIZO food research, P.O. Box 20, 6710 BA Ede, The Netherlands. Phone: 31 (0) 318 659 629, Fax: 31 (0) 318 650 400. E-mail: michiel.kleerebezem@nizo.nl.*

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and/or tolerance to various substrates, including bile salts, penicillin V, and acyl-homoserine lactones. Furthermore, the evolutionary conservation of bsh homologs was investigated in several strains of the species L. plantarum, using complete genome hybridization (CGH) (38). These results indicated that bsh2, bsh3, and bsh4 appear to be conserved among L. plantarum strains, suggesting an important physiological role. In addition, the presence of bsh1 appeared to be correlated with the Bsh activity of L. plantarum strains.

MATERIALS AND METHODS

Bacterial strains, plasmids, and primers. The bacterial strains, plasmids, and primers used in this study and their relevant features are listed in Table S1 in the supplemental material. L. plantarum WCFS1 (29) and bsh mutant derivatives were grown at 37°C in MRS broth (Difco, West Molesey, United Kingdom), without aeration. The heterologous nisin-controlled expression (NICE) host L. lactis NZ9000 and its parental strain, MG1363 (21), which was used as an intermediate cloning host for NICE overexpression constructs (31, 36), were grown at 30°C in M17 broth (Oxoid, United Kingdom) supplemented with 0.5% glucose (wt/vol; G-M17), without aeration. Escherichia coli strains DH5α (55) and MC1061 (9, 54) were used as intermediate cloning hosts for L. plantarum mutagenesis constructs and pCR-Blunt constructs, respectively, and were grown at 37°C on TY broth (25), with aeration. When appropriate, antibiotics were added to the media. For L. plantarum, 10 μg/ml chloramphenicol and 10 μg/ml (in liquid medium) or 30 μg/ml (on solid medium) erythromycin were used. For L. lactis, 10 μg/ml chloramphenicol was used. For E. coli, 10 μg/ml chloramphenicol and 250 μg/ml erythromycin were used.

DNA and protein manipulations. Plasmid DNA was isolated from E. coli on a small scale, using the alkaline lysis method (5). Large-scale plasmid DNA isolations were performed using Jetstar columns as recommended by the manufacturer (Genomed GmbH, Bad Oberhausen, Germany). Purification of DNA fragments from agarose gels was performed using the Wizard SV gel and PCR cleanup system (Promega, Leiden, The Netherlands). DNA isolation and transformation of L. plantarum and L. lactis were performed as described previously (16, 33). For DNA manipulations in E. coli, protein extraction, and sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), standard procedures were employed (46). Restriction endonucleases, Taq, Pfu, and Pwo DNA polymerases, T4 DNA ligase, and Klenow enzyme were used as prescribed by the manufacturers (Promega, Leiden, The Netherlands, and Boehringer, Mannheim, Germany). Primers were obtained from Genset Oligos (Paris, France).

RNA isolation and Northern blotting. For RNA isolation, an overnight culture of L. plantarum WCFS1 was diluted 50-fold in 50 ml of fresh MRS medium, with or without the addition of 0.05% (wt/vol) porcine bile (Sigma, Zwijndrecht, The Netherlands), and grown to an optical density at 600 nm (OD600) of 1. Subsequently, 3 volumes of quench buffer (66% methanol, 66.7 mM HEPEs, pH 6.5 [−40°C]) were added (44). The cells were immediately pelleted by centrifugation at 3,500 × g for 10 min (Megafuge 1.0R; Heraeus, Hanau, Germany), resuspended in 750 μl of ice-cold TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH 7.5), and mechanically disrupted (FastPrep FP120; Oobiogene, Illkirch, France) in the presence of 0.8 g of zirconium beads (Biospiec Products, Bartlesville, OK), 0.18 g of Maculoid (Kronos Titan GmbH, Leverkusen, Germany), 50 μl of 10% SDS, and 500 μl of phenol. Subsequently, the RNA was purified from the upper, aqueous phase of the cell extract by phenol-chloroform extraction, precipitated with absolute ethanol, washed with 70% ethanol (46), and resuspended in 50 μl of MQ water. Northern blot analysis was performed as described earlier (46), using total RNA. As probes for bsh1, bsh2, bsh3, and bsh4, PCR amplification products of a large part of the genes (0.7 to 0.8 kb) were used after being amplified with Taq polymerase, using L. plantarum WCFS1 total DNA as a template and primers bsh1F and bsh1R. The resulting amplicon was cloned into pCR-Blunt II-TOPO vector (Invitrogen, Breda, The Netherlands). Subsequently, the bsh1 gene was recovered from the resulting plasmid as a 1.1-kb AflII-HindIII fragment and cloned downstream of and translationally fused to the nisA promoter in NcoI-HindIII-digested pNZ8048 (31, 36). The resulting nisin-controlled bsh1 expression plasmid was designated pNZ5306.

A DNA fragment containing bsh2 was amplified by PCR, using Pfu polymerase, with L. plantarum WCFS1 genomic DNA as a template and primers bsh2F and bsh2R. The amplicon obtained was digested with HindIII, and the resulting 1.1-kb fragment was cloned downstream of and translationally fused to the nisA promoter in NcoI-KpnI-digested pNZ8048, yielding the bsh2 overexpression construct pNZ5307.

The overexpression plasmid for bsh3 was constructed analogously to the bsh1 plasmid pNZ5306. A bsh3-containing PCR ampiclon (using Pfu polymerase, L. plantarum WCFS1 genomic DNA as a template, and primers bsh3F and bsh3R) was initially cloned into pCR-Blunt (Invitrogen, Breda, The Netherlands), and a 1.1-kb fragment containing bsh3 was subcloned into pNZ8048 following the same cloning strategy as that employed with bsh1, yielding pNZ5308, which contains the bsh3 gene under control of the nisC transcription and translation signals.

Finally, a bsh4-containing DNA fragment was amplified by PCR, using Pfu polymerase, with L. plantarum WCFS1 genomic DNA as a template and primers bsh4F2 and bsh4R. The amplicon obtained was cloned into the pCR-Blunt II-TOPO vector. The bsh4 gene was recovered from the resulting plasmid by digestion with KpnI and ApaI, followed by partial digestion with AflII and cloning of the 1.1-kb, bsh4-containing fragment downstream of and translationally fused to the nisA promoter in NcoI-KpnI-digested pNZ8048, yielding the bsh4 overexpression construct pNZ5309.

For all overexpression constructs, the sequence of the cloned bsh gene was confirmed to be correct by sequencing. For overexpression studies of the bsh genes in L. lactis by use of the NICE system, pNZ5306, pNZ5307, pNZ5308, and pNZ5309 were transformed into L. lactis NZ9000.

Construction of bsh deletion mutant strains. For the construction of deletion derivatives of L. plantarum WCFS1 that lack one or more of the bsh genes, the previously reported Cre-lox-based system for multiple gene deletions was used (33). The bsh1 deletion vector pNZ325 and bsh1 deletion strain NZ5305 were constructed as described previously (33). For all overexpression constructs, the sequence of the cloned bsh gene was confirmed to be correct by sequencing. For overexpression studies of the bsh genes in L. lactis by use of the NICE system, pNZ5306, pNZ5307, pNZ5308, and pNZ5309 were transformed into L. lactis NZ9000.

Construction of bsh deletion mutant strains. For the construction of deletion constructs of L. plantarum WCFS1 that lack one or more of the bsh genes, the previously reported Cre-lox-based system for multiple gene deletions was used (33). The bsh1 deletion vector pNZ325 and bsh1 deletion strain NZ5305 were constructed as described previously (33). The bsh2 mutation vector pNZ5329 (see Table S1 in the supplemental material) was constructed by successive cloning of the PCR-amplified 1.0-kb 5′ and 3′ chromosomal flanking regions of bsh2 (lp_0067) (using Pfu polymerase, L. plantarum WCFS1 genomic DNA as a template, and the primer sets bsh2koF1R/bsh2koF1R and bsh2koF2/bsh2koF2R, respectively [see Table S2 in the supplemental material]) into the SwaI and Eco130I restriction sites of pNZ319 (see Table S1 in the supplemental material), respectively (33). Analogously, the bsh2 mutation vector pNZ3332 (see Table S1 in the supplemental material) was constructed by successive cloning of the PCR-amplified 1.0-kb 5′ and 3′ chromosomal flanking regions of bsh3 (lp_3362) (using Pfu polymerase, L. plantarum WCFS1 genomic DNA as a template, and the primer sets bsh3koF1R/bsh3koF1R and bsh3koF2/bsh3koF2R, respectively [see Table S2 in the supplemental material]) into the SwaI and Eco130I restriction sites of pNZ3319 (see Table S1 in the supplemental material), respectively (33). For all bsh deletion constructs, the sequences of the cloned PCR-amplified regions were verified by double-strand sequence analysis (Basclear, Roosendaal, The Netherlands).

The bsh deletion strains NZ5305 (Δbsh1), NZ5307 (Δbsh2), NZ5309 (Δbsh3), NZ5311 (Δbsh4), NZ5313 (Δbsh1Δbsh2), NZ5315 (Δbsh3Δbsh4), NZ5324 (Δbsh1Δbsh2Δbsh3), NZ5326 (Δbsh1Δbsh2Δbsh4), NZ5328 (Δbsh1Δbsh3Δbsh4), NZ5330 (Δbsh2Δbsh3Δbsh4), and NZ5332 (Δbsh1Δbsh2Δbsh3Δbsh4) were constructed as described previously (33) (see Table S1 and Fig. S1 in the supplemental material). Briefly, the desired bsh deletion vector was transformed into L. plantarum WCFS1 or one of its mutant derivatives by electroporation (33), and double-crossover gene replacement mutants in which the target gene was replaced by a loxP-cre-loxP-loxP cassette were selected based on their Cm r phenotype. Correct integration of the loxP-cre-loxP-loxP cassette into the chromosomal flanking regions of the integrated loxP-cre-loxP-loxP cassette, using primers annealing uniquely to genomic sequences combined with the mutagenesis vector-specific primers 85 and 87 that annealed to the P26 cre region (i.e., primers 106a/85 and 87/107a for bsh1 replacement, bsh2ko-up/85 and 87/bsh2ko-down for bsh2 replacement,
determine the Bsh activities of above; see Table S1 in the supplemental material). Stable deletion mutations in one, two, three, or all four of the bsh loci, respectively (Fig. 1), were resuspended to a final OD 600 of 200 in 55 mM sodium acetate buffer, pH 5.5, containing 1 mM DTT and 10% (vol/vol) phenylmethylsulfonyl fluoride (PMSF). Cells were centrifuged at 3,000 × g for 10 min and precleared by 250 μl of 100 mM sodium acetate buffer, pH 5, containing 1 mM PMSF.

To determine penicillin, ampicillin, cephalosporin, acyl-homoserine lactone, and phenylacetyleglycine acylase activities, 5 volumes of the cell extract was mixed with 5 volumes of 100 mM sodium acetate buffer, pH 5, containing 1 mM PMSF, and 5 volumes of the cell extract was added. DTT and 1 volume of 100 mM of penicillin V, penicillin G, ampicillin, cephalosporin C, ketocaproyl-homoserine lactone, oxoocetyl-homoserine lactone, or phenylacetyleglycine and incubated overnight at 37°C. To test the reaction, 35 volumes of 285 mM sodium acetate buffer, pH 4, was added. Free amino groups resulting from enzymatic conversion of the substrate were detected by the addition of 5 volumes of 0.1 mg/ml fluorescein in acetone, centrifugation for 10 min at 3,000 × g at room temperature (TechnoSpin R; Sorval Instruments), and measurement of fluorescence in the supernatant (excitation at 360 nm and emission at 465 nm) (GENios F129004; TECAN Benelux, Giessen, The Netherlands). Acylation of NIPAB at 25°C was determined by the increase of absorption at 405 nm, which was followed for 30 min and measured overnight. As a positive control, purified penicillin acylase and purified end products of the acylase reactions (6-amino-penicillanic acid, 7-aminocephalosporanic acid, homoserine lactone, and glycine) (Sigma, Zwijndrecht, The Netherlands) were used. Furthermore, acylase activity for 6-nitro-3-(phenylacetamido) benzoic acid (NIPAB; Sigma, Zwijndrecht, The Netherlands), which is a commonly used chromogenic substrate for assaying penicillin G acylase activity, was determined as described earlier (1, 32). Brieﬂy, 1 volume of cell extract was mixed with 9 volumes of 100 mM sodium acetate buffer containing 1 mM DTT and 2.5 mM NIPAB. Acylation of NIPAB at 25°C was determined by the increase of absorption at 405 nm, which was followed for 30 min and measured overnight. As a positive control, purified penicillin amidase of E. coli was used (Sigma, Zwijndrecht, The Netherlands).

Bile salt and penicillin V tolerance. To evaluate the tolerance of L. plantarum WCFS1 and its bsh mutant derivatives to bile salts and penicillins, overnight cultures were inoculated 1:20 into fresh MRS medium containing 0 to 30% ox gall (wt/vol), 0 to 0.4% (wt/vol) GDC, 0 to 10% (wt/vol) TDC, and 0 to 14 μg/ml penicillin V. Growth was followed for 16 h by measurement of the OD 600 at 37°C at intervals of 15 min (Spectra Max Plus 384; Molecular Devices, Sunnyvale, CA).

bsh diversity in L. plantarum strains. The genomic diversity of L. plantarum strains (299, 299v, CIP102359, CIP104440, CIP104441, CIP104448, CIP104450, CIP104451, CIP104452, LPS-2, NCIMB12120, and SF2A35B) was previously investigated by use of strain WCFS1-derived DNA microarrays (38). This genomic genotyping database allowed the evaluation of the presence and/or absence of homologues of the four genes that were initially annotated in the L. plantarum WCFS1 genome as bsh genes (bsh1, bsh2, bsh3, and bsh4). Furthermore, a positive cutoff P value of 1e-4 was used for presence calling of bsh homologs.

Detection of Bsh activity in L. plantarum strains. The presence of Bsh activity in different L. plantarum strains (299, 299v, CIP102359, CIP104440, CIP104441, CIP104448, CIP104450, CIP104451, LPS-2, NCIMB12120, and SF2A35B) was detected using a bile salt plate assay, as described earlier (12). Briefly, overnight cultures of L. plantarum strains were transferred to solid MRS medium, with or without 0.5% (wt/vol) of the bile salt TDC, and incubated anaerobically for 48 h at 37°C. Bsh-active strains were recognized by the formation of opaque white colonies in the presence of TDC, which is due to the precipitation of deconjugated bile salt forms.

RESULTS

Expression of bsh genes. L. plantarum WCFS1 contains four related bsh genes (29). The expression of these genes was studied by Northern blotting during exponential growth phase in the presence or absence of porcine bile (data not shown). The probes used for Northern blotting were specific for each of cells of L. plantarum or cell extracts of L. lactis bsh overexpression strains was determined at 37°C by HPLC as described previously (13). Separations were carried out with a reversed-phase resin-based column (PLRP-S; 5 μm by 300 Å by 250 mm by 4.6-mm inner diameter [Polymer Laboratories, Shropshire, United Kingdom]) and matching precolumn. Bile salts were detected using a pulsed amperometric detector (E&G Princeton Applied Research, Princeton, NJ) equipped with a gold working electrode and a reference electrode (Ag/AgCl). Chromatograms were analyzed and integrated using the Chromelon program (Dionex, Sunnyvale, CA), and Bsh activity was determined based on the disappearance of the conjugated bile salts used as a substrate.


determination of Bsh activities of above; see Table S1 in the supplemental material). Stable deletion mutations in one, two, three, or all four of the bsh loci, respectively (Fig. 1), were resuspended to a final OD 600 of 200 in 55 mM sodium acetate buffer, pH 5.5, containing 1 mM DTT and 10% (vol/vol) phenylmethylsulfonyl fluoride (PMSF). Cells were centrifuged at 3,000 × g for 10 min and precleared by 250 μl of 100 mM sodium acetate buffer, pH 5, containing 1 mM PMSF.

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the individual \textit{bsh} genes and did not show cross-hybridization. The Northern blot analysis established that \textit{bsh1}, \textit{bsh2}, \textit{bsh3}, and \textit{bsh4} are all expressed in \textit{L. plantarum} WCFS1 during the exponential growth phase. The estimated sizes of the transcripts corresponded with those predicted for monocistronic transcription of \textit{bsh1}, \textit{bsh2}, \textit{bsh3}, and \textit{bsh4} (1.2 kb for \textit{bsh1}, \textit{bsh2}, and \textit{bsh3} and 1.1 kb for \textit{bsh4}). Incubation of \textit{L. plantarum} WCFS1 grown in liquid medium with 0.5% (wt/vol) porcine bile did not induce significant expression of any of the \textit{bsh} genes.

**Divergence of Bsh in \textit{L. plantarum}**. In order to explore the functionality of the \textit{bsh} genes of \textit{L. plantarum} further, the presence of homologues of the four \textit{bsh} genes of \textit{L. plantarum} WCFS1 in other \textit{L. plantarum} strains was determined by analysis of \textit{L. plantarum} genomic DNA samples on \textit{L. plantarum} WCFS1-specific microarrays, as described earlier (38), using a positive cutoff \textit{P} value of 1e\textsuperscript{-5} for \textit{bsh} gene presence calls. The presence and absence calls were correlated with the experimentally determined capability of these strains to hydrolyze bile salts, using a previously described Bsh plate assay (12; data not shown). Remarkably, in all \textit{L. plantarum} strains, \textit{bsh2}, \textit{bsh3}, and \textit{bsh4} generated very good probability scores for gene presence, suggesting that these genes are highly conserved among \textit{L. plantarum} strains. In contrast, the presence of a gene homologous to \textit{bsh1} appeared to vary among the strains analyzed. The \textit{bsh1} gene appeared to be absent in 4 of the 13 strains analyzed, which appeared to correlate well with the absence of Bsh activity in these four strains. The remaining nine strains appeared to contain a \textit{bsh1} homologue, as concluded from CGH analysis. Seven of these nine strains also displayed clearly detectable Bsh activity in the plate assay employed, while the other two strains did not display activity in this assay. Therefore, it is likely that Bsh activity is related to the presence of a \textit{bsh1} homologue in the species \textit{L. plantarum}.

**Bsh activities of individual Bsh proteins**. Since all \textit{bsh} genes of \textit{L. plantarum} WCFS1 appeared to be expressed during exponential growth, the contributions of the individual \textit{bsh} genes to the total Bsh activity were determined. For this purpose, heterologous \textit{bsh} overexpression strains of \textit{L. lactis} were established. Furthermore, a set of single and multiple \textit{bsh} deletion derivatives of \textit{L. plantarum} WCFS1 was constructed (see Table S1 in the supplemental material).

**Heterologous overexpression strains**. For heterologous overexpression using the NICE system (36), vectors pNZ5306, pNZ5307, pNZ5308, and pNZ5309, containing the \textit{bsh1}, \textit{bsh2}, \textit{bsh3}, and \textit{bsh4} genes, respectively, translationally coupled to the nisin-inducible \textit{nis4} promoter, were constructed and transformed into the Bsh-deficient bacterium \textit{L. lactis} NZ9000. Only following nisin induction could overexpression protein products be detected by SDS-PAGE for \textit{Bsh2} and \textit{Bsh3} (Fig. 2A), at their expected molecular masses of 38 and 36 kDa, respectively. HPLC-based Bsh activity assays (13) using taurine- and glycine-conjugated bile salts of cholic acid, deoxycholic acid, and chenodeoxycholic acid as substrates (see Fig. S1 in the supplemental material) revealed that \textit{Bsh1}, \textit{Bsh3}, and \textit{Bsh4}, but not \textit{Bsh2}, are capable of bile salt deconjugation (Fig. 2A). These results confirmed the functional heterologous expression of both \textit{Bsh1} and \textit{Bsh4}, despite the lack of detection of the corresponding protein products by SDS-PAGE; in heterologous expression, the amount of protein produced is not necessarily correlated to specific activity. \textit{Bsh1} and \textit{Bsh3} displayed a strong preference for glycine-conjugated bile salts compared to taurine-conjugated bile salts, while \textit{Bsh4} appeared to exclusively convert glycine-conjugated bile salts. Moreover, consistent differential substrate specificity of \textit{Bsh1}, \textit{Bsh3}, and \textit{Bsh4} was observed, with a substrate preference diminishing from deoxycholic acid to chenodeoxycholic acid and to cholic acid-conjugated bile salts (Fig. 2B).

**Combinatorial \textit{bsh} deletion strains**. In parallel, a set of combinatorial \textit{bsh} mutant derivatives (Fig. 1; see Table S1 in the supplemental material) of \textit{L. plantarum} WCFS1 was con-
structured using a Cre-lox-based mutation system that allows the effective deletion of multiple genes in a single genetic background (33). The bsh mutant derivatives were checked by PCR, amplifying each of the four bsh loci (Fig. 1). Growth appeared to be unaffected in these mutants compared to that of the parental strain, indicating that the deleted genes were not necessary for growth under normal laboratory conditions (data not shown).

Bsh activity was analyzed using an HPLC-based assay (13), with the purified bile salt GDC (Fig. 2C) and male Fischer rat bile (data not shown) as substrates for the triple and quadruple bsh mutant strains (NZ5324, NZ5326, NZ5328, NZ5330, and NZ5332). These experiments revealed that cells harboring an intact copy of the bsh1 gene (NZ5330) displayed Bsh activity levels that were comparable to those of the wild-type strain, confirming previous studies identifying this gene as the major Bsh-encoding gene in L. plantarum (33). Moreover, Bsh1-dependent Bsh activity declined drastically (about 40 times) in cells harvested during stationary growth phase compared with the activity in cells obtained from the logarithmic phase of growth (Fig. 2C), indicating growth phase-dependent expression of the bsh1 gene. In all triple mutant, bsh1-deficient strains (NZ5324, NZ5326, and NZ5328) and the quadruple bsh mutant strain (NZ5332), a small but detectable amount of bile salt hydrolysis was found (Fig. 2C), which appeared to be bsh2, bsh3, or bsh4 independent.

Alternative functionality of individual Bsh proteins. Since in L. plantarum WCFS1 the bile salt hydrolase activity appeared to be independent of bsh2, bsh3, and bsh4, the functionality of these genes was investigated by determination of the activities of the Bsh proteins on a variety of putative alternative (non bile salt) substrates (see Table S3 in the supplemental material). The Bsh proteins of L. plantarum WCFS1 share significant sequence homology with penicillin V acylase enzyme family members (including the experimentally verified penicillin V acylase of Listeria monocytogenes EGDe [4], with amino acid identity levels ranging from 30 to 26%), which are in turn related to β-lactam acylases and to acyl-homoserine lactone acylases, which play a key role in quorum sensing-dependent gene regulation in gram-negative bacteria. Bsh, penicillin acylase, β-lactam acylase, and acyl-homoserine lactone acylase all act on the same type of chemical bond, although the structures of their substrates differ considerably (see Fig. S2 in the supplemental material). Thus, the activities of the overexpression products for the individual Bsh proteins, using the NICE system (36) in L. lactis NZ9000 (using pNZ5306, pNZ5307, and pNZ5309), were determined for the substrates penicillin V, penicillin G, NIPAB (which is a commonly used substrate for spectrophotometric detection of penicillin acylase activity), the β-lactams ampicillin and cephalosporin C, ketocaproyl-homoserine lactone, oxooctanoyl-homoserine lactone, and phenylacetylgytone, which is a molecule that is involved in phenylalanine metabolism and thereby readily available to L. plantarum and is cleaved by penicillin acylase of E. coli (51) (see Table S3 and Fig. S2 in the supplemental material).

For NIPAB, ampicillin, cephalosporin C, and phenylacetylethyl-glycone, no activity could be detected in any of the Bsh overexpression strains. However, bsh3 (and, to a lesser extent, bsh2 and bsh4) overexpression resulted in an increase in acylase activity of penicillin V (on average, 4.3, 2.2, and 1.8 times, respectively) relative to that of the control strain NZ9000 in several independent experiments (Fig. 3). These findings clearly suggest a role as a penicillin acylase for Bsh3, and possibly Bsh2 and Bsh4. In addition, bsh2, bsh3, and bsh4 overexpression strains all showed increases in activities (on average, 1.9 times) toward penicillin G, ketocaproyl-homoserine lactone, and oxooctanoyl-homoserine lactone relative to those of NZ9000, suggesting a broad range of enzyme specificities. Notably, the bsh1 overexpression strain showed no significant activity toward penicillin V, penicillin G, ketocaproyl-homoserine lactone, or oxooctanoyl-homoserine lactone compared to those of NZ9000, confirming the role of bsh1 as a bona fide Bsh.

Bile salt and penicillin V tolerance. The contributions of the individual bsh genes to the tolerance of the bile salts TDC and GDC, ox gall, and penicillin V (see Fig. S1 and S2 in the supplemental material) were determined using the triple and quadruple L. plantarum bsh mutant strains (NZ5324, NZ5326, NZ5328, NZ5330, and NZ5332). To this end, growth in the presence of a range of bile (salt) and penicillin concentrations was monitored spectrophotometrically.

L. plantarum was able to grow in the presence of up to 30% (wt/vol) of ox gall, with no significant differences in growth between the strains (data not shown). However, due to the high concentrations of ox gall used, measurements were severely hampered. In addition, L. plantarum was able to grow in the presence of >14% (wt/vol) of TDC, with no significant differences found for the strains used (data not shown). However, L. plantarum appeared to be remarkably more sensitive to GDC, with obvious differences between the strains (data not shown). The results clearly established that the presence of bsh1 in L. plantarum enhances GDC bile salt tolerance. Each of the bsh1-deficient derivatives displayed GDC-mediated growth inhibition at concentrations as low as 0.1% (wt/vol) GDC. In contrast, strains containing an intact bsh1 gene were capable of sustaining normal growth characteristics at up to 0.5 to 0.7% (wt/vol) GDC. Analogous with the limited level of hydrolytic activity toward bile salts, bsh2, bsh3, and bsh4 did not appear to contribute significantly to tolerance of GDC.

Furthermore, growth of L. plantarum was inhibited at the lowest concentration of penicillin V tested (0.3 μg/ml), with complete inhibition of growth in the presence of 8 μg/ml penicillin V. However, no difference was found between WCFS1 and...
and its bsh mutant derivatives, indicating that none of the bsh
genes appeared to influence penicillin V tolerance in L. plantarum under the conditions analyzed here.

DISCUSSION

Bile salt hydrolysis is a biologically important reaction in the intestinal tract, since it is the first step in bile salt biotransformations carried out by intestinal bacteria. The formation of secondary bile salts has a significant impact on the physiology of the host, as exemplified by suggestions regarding their implication in lowering of blood cholesterol levels (42) and in various intestinal diseases, such as the formation of gallstones and colon cancer (45). Furthermore, deconjugated bile salts were shown to induce mucin production by intestinal epithelial cells (30), possibly indicating irritation of the epithelial cells by the strong surface-active properties of deconjugated bile salts.

Remarkably, for several strains (e.g., Lactobacillus johnsonii
100-100 and NCC533, Lactobacillus acidophilus NCFM, and L. plantarum WCFS1), the presence of more than one gene encoding a Bsh homologue has been predicted. In L. plantarum WCFS1, the sequences of the bsh genes share higher levels of similarity with bsh genes from other strains or species than with each other. For example, bsh1 shares highest sequence similarity with the bsh genes of other L. plantarum strains and Enterococcus faecalis, whereas bsh2 and bsh4 share sequence similarity with the sequence annotated as bsh of Lactobacillus brevis and share no significant sequence similarity with any other organism whose sequence is publicly available; by analogy, bsh3 shares significant sequence similarity only with the sequence annotated as bsh of Lactobacillus sakei and L. brevis. The bsh genes of L. plantarum WCFS1 may have been acquired via horizontal gene transfer, as suggested earlier for L. johnsonii (18). However, the overall conservation among L. plantarum strains of the bsh2 to bsh4 genes would indicate that this acquisition occurred very early in the evolution of this species. Moreover, this high level of conservation also supports an important role of these genes in the physiology and lifestyle of the species L. plantarum.

In line with the previous finding that bsh1 is the major Bsh in L. plantarum WCFS1 (33), the presence of bsh1 appeared to correlate with the capability to hydrolyze bile salts in 11 of 13 L. plantarum strains. For 2 of 13 strains, however, a Bsh1 homolog was concluded to be present, while the capability to hydrolyze TDC was not detected in these strains. However, this apparent inconsistency can be explained by the fact that the detection of the presence of bsh1 in a particular strain does not necessarily correlate with expression of functional Bsh1 that is identical to the protein expressed by L. plantarum WCFS1. Firstly, the bsh1-like gene found by CGH analysis may contain disruptive (point) mutations, leading to detection of the gene but a lack of functionality. Secondly, the presence of a bsh1-like gene in a particular strain does not give any information on the expression of this gene under the conditions applied in the enzyme assay. Thus, the presence of a gene does not necessarily correlate with expression of this gene. Thirdly, the bsh1 homologues found may display a more stringent substrate preference than that of Bsh1 of L. plantarum WCFS1, thereby failing to convert the specific bile salt TDC used in this experiment. Thus, the Bsh1 homologues may display activity for bile salts other than TDC.

In L. plantarum WCFS1, the presence of bsh1 correlated with GDC tolerance but not with TDC tolerance. The capacity to hydrolyze bile salts has been found to be linked to bile salt tolerance in L. plantarum (15) and several other bacteria, including Lactococcus amylovorus (24), Listeria monocytogenes (4), and Bifidobacterium (40). Thus, the preference of L. plantarum WCFS1 for deconjugation of glycine-over tauroine-conjugated bile salts appears to be reflected in the differential tolerance toward GDC and TDC. This could be related to the higher toxicity of glycine-conjugated bile salts than of taurine-conjugated bile salts, leading to evolution of a preference of Bsh for glycine-conjugated bile salts. Indeed, most Bsh proteins show a preference for glycine-conjugated bile salts (3). Although the precise mechanism is unknown, bile salt hydrolysis could be of great importance for survival in vivo. Since deconjugated bile acids display reduced solubility compared to their conjugated counterparts, especially at lower pH values, bile salt hydrolysis may lead to precipitation of the bile salts and thereby relieve stress levels caused by these surface-active chemicals. Thereby, the capability to hydrolyze bile salts may contribute to the survival and persistence of bacterial strains in the intestinal tract, as previously shown for Listeria monocytogenes (17). Analogously, Bsh activity appears to be present in all lactobacilli isolated from the gastrointestinal environment (48). Notably, L. plantarum WCFS1 is capable of persisting in the mouse gastrointestinal tract for 10 days (41) and has been shown to display relatively high survival and activity during transit of the human gastrointestinal tract (53). Additional animal experiments using the set of L. plantarum WCFS1 bsh mutant derivatives constructed in this work may clarify the role of the bsh genes in persistence and survival of the organism in the gastrointestinal tract.

All four of the bsh genes of L. plantarum WCFS1 are expressed during the exponential phase of growth as monocistrionic transcripts. The expression of the bsh genes did not appear to be induced as a consequence of exposure to porcine bile during growth in liquid media. In contrast, previous whole-genome transcriptome studies suggested that expression of bsh1 was induced by porcine bile when cells were grown on solid media, while the expression of bsh3 appeared to be repressed under these conditions (7). In the same study, no regulation of bsh2 or bsh4 by porcine bile was detected. Notably, these findings are in apparent agreement with the finding that bsh1 and, to a lesser extent, bsh3 are capable of bile salt hydrolysis. Nevertheless, the discrepancy in regulation of bsh expression may be due to the difference in growth conditions used (liquid versus solid medium). Explaining these different findings would require further investigation of the regulation of expression of bsh as a function of culture conditions.

Heterologous overexpression of the bsh genes by use of the NICE system in the Bsh-deficient host L. lactis NZ9000 confirmed that Bsh1 is a functional and bona fide Bsh, as described earlier (33). In addition, Bsh3 and Bsh4 displayed bile salt hydrolysis activity, although with lower enzymatic efficiencies than that of Bsh1, which suggests that these enzymes could contribute to the overall Bsh activity displayed by L. plantarum WCFS1. Analogous to what has been described for most of the Bsh enzymes studied to date (27, 37, 49, 50), the Bsh1, Bsh3,
and Bsh4 enzymes displayed a clear preference for glycoconjugated over tauroconjugated bile salts. Furthermore, the substrate preference of bsh1, bsh3, and bsh4 decreased from deoxycholic to chenodeoxycholic and cholic acid as the steroid moiety of the conjugated bile salts (see Fig. S1 and S2B in the supplemental material), which has previously also been found for the Bsh activity of Lactobacillus buchneri (39). This substrate preference is probably related to the positioning and presence or absence of hydroxyl groups on the steroid moiety of the bile salts (see Fig. S1 in the supplemental material) and, concomitantly, the binding pocket properties of the enzyme.

Mutation analysis with L. plantarum WCFS1 confirmed that Bsh1 is the major Bsh. Nevertheless, differential activity of Bsh2, Bsh3, or Bsh4 under specific conditions or selective activity toward specific bile salts cannot be excluded. Screening for involvement of the bsh genes of L. plantarum WCFS1 in the conversion of various putative alternative substrates showed that Bsh3 and, to a lesser extent, Bsh2 and Bsh4 were able to hydrolyze penicillin V and penicillin G. To date, the in vivo role of penicillin acylase remains unknown. Notably, the enzyme name appears to reflect primarily its industrial application rather than its natural substrate. Therefore, Bsh2, Bsh3, and Bsh4 may play a role in acylation of compounds other than the substrates tested here, such as additional phenylacetic acid derivatives, as suggested by the capability of penicillin acylases to cleave these substrates in addition to penicillins (51). Indeed, phenylacetic acid derivatives would be available to L. plantarum in vivo, since they are formed by microbial activity on plant constituents, where L. plantarum was found to occur naturally (20). However, these substrates are not commercially available. In addition, low-level acylase activity by Bsh2, Bsh3, and Bsh4 toward two types of acyl-homoserine lactones was found, indicating the broad substrate specificity of these enzymes. Acyl-homoserine lactones could have an important function in adhesion of bacteria to the epithelium of the intestinal tract. For example, the pathogen Pseudomonas aeruginosa was found to upregulate PA-1 lectin/adhesin, an important virulence factor in this strain, in response to butanoyl-homoserine lactone in the environment (56). Therefore, bacteria that are available. In addition, low-level acylase activity by Bsh2, Bsh3, and Bsh4 suggests an important but so far unknown role of these genes in the physiology and lifestyle of the species L. plantarum.

REFERENCES
8. Reference deleted.
conjugated bile acids inhibit bacterial growth of *Clostridium perfringens* and induce its extracellular cholyglycine hydrolase. Steroids 59:485–489.


