



Disruption of a Transcriptional Repressor by an Insertion Sequence Element Integration Leads to Activation of a Novel Silent Cellobiose Transporter in *Lactococcus lactis* MG1363

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ABSTRACT *Lactococcus lactis* subsp. *cremoris* strains typically carry many dairy niche-specific adaptations. During adaptation to the milk environment these former plant strains have acquired various pseudogenes and insertion sequence elements indicative of ongoing genome decay and frequent transposition events in their genomes. Here we describe the reactivation of a silenced plant sugar utilization cluster in an *L. lactis* MG1363 derivative lacking the two main cellobiose transporters, PtcBA-CelB and PtcBAC, upon applying selection pressure to utilize cellobiose. A disruption of the transcriptional repressor gene *llmg_1239* by an insertion sequence (IS) element allows expression of the otherwise silent novel cellobiose transporter *Llmg_1244* and leads to growth of mutant strains on cellobiose. *Llmg_1239* was labeled CcIR, for cellobiose cluster repressor.

IMPORTANCE Insertion sequences (ISs) play an important role in the evolution of lactococci and other bacteria. They facilitate DNA rearrangements and are responsible for creation of new genetic variants with selective advantages under certain environmental conditions. *L. lactis* MG1363 possesses 71 copies in a total of 11 different types of IS elements. This study describes yet another example of an IS-mediated adaptive evolution. An integration of *IS981* or *IS905* into a gene coding for a transcriptional repressor led to activation of the repressed gene cluster coding for a plant sugar utilization pathway. The expression of the gene cluster allowed assembly of a novel cellobiose-specific transporter and led to cell growth on cellobiose.

KEYWORDS *Lactococcus lactis*, cellobiose, plant, dairy, reverse evolution, IS element, sRNA, CcIR, PTS, sugar metabolism

L *Lactococcus lactis* demonstrates an impressive genome plasticity and polymorphism. Various *L. lactis* strains can be found in many different environments, such as plants, milk, animals, and related niches (1, 2). Although a number of *L. lactis* strains have been “domesticated” and cultivated in milk for many years, thorough analyses of *L. lactis* genomes reveals the presence of multiple silent gene clusters and pseudogenes with predicted functions in plant sugar utilization as remnants of their plant niche-related history (2–7). Point mutations and genome rearrangements occasionally lead to activation of silent genes and gene clusters, sometimes even granting them new functions. Adaptive evolution in lactococcal strains occurs mostly by horizontal gene transfer and by intracellular activity of the mobilome (7, 8). Insertion sequences (IS) are the simplest type of mobile genetic elements. A typical IS element usually encodes one or two proteins necessary for its transposition and contains (near) perfect inverted repeats at

Received 9 June 2017 Accepted 21 September 2017

Accepted manuscript posted online 29 September 2017

Citation Solopova A, Kok J, Kuipers OP. 2017. Disruption of a transcriptional repressor by an insertion sequence element integration leads to activation of a novel silent cellobiose transporter in *Lactococcus lactis* MG1363. *Appl Environ Microbiol* 83:e01279-17. <https://doi.org/10.1128/AEM.01279-17>.

Editor Maia Kivisaar, University of Tartu

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its termini (8). The genome of *Lactococcus lactis* subsp. *cremoris* MG1363 contains 11 different types of IS elements and 71 copies in total. Recombination between identical IS elements and their ability to transpose from one location to another in a genome provide great genetic flexibility. IS elements have often been described to be involved in adaptation of *L. lactis* strains to unfavorable conditions during starvation or oxidative stress (8–10).

The metabolism of the plant disaccharide cellobiose by some lactococcal laboratory strains is a good example of an ongoing adaptive evolution process. After constant propagation under laboratory conditions, these strains lost the capability to efficiently take up and utilize cellobiose, and certain changes in their genomes are necessary to restore these deficiencies. *L. lactis* MG1363 can take up and utilize cellobiose when one of two dedicated phosphoenolpyruvate:carbohydrate phosphotransferase systems (PTSs) is functional. Activation of the PtcBA-CelB system in *L. lactis* MG1363 requires a *celB* promoter up-mutation (11), and PtcABC is assembled in NZ9000 (an isogenic derivative of MG1363) when a *cre* site in the promoter region of *ptcC* is disrupted and the gene becomes constitutively expressed (12; see Fig. 4). Besides cellobiose, PtcBA-CelB can transport lactose, while PtcABC usually imports glucose (13, 14).

We found that even after deletion of the membrane-integrated EII subunit C (EIIc) components of these two PTSs, namely, CelB and PtcC, *L. lactis* MG1363 was able to restore growth in a medium supplemented with cellobiose as the sole carbon source. Two types of IS elements were discovered that had integrated at different positions relative to a gene coding for a putative LacI-type transcriptional repressor in the Cel⁺ isolates. This gene is located upstream of a putative cellobiose-, beta-glucoside-, or lichenan-catabolic gene cluster. The aim of this study was to assess the effect of an IS integration in the putative repressor gene and to identify the silent cellobiose transporter responsible for the restored Cel⁺ phenotype of *L. lactis* MG1363 Cel⁻ strains.

RESULTS

Cultivation of Cel⁻ strains in cellobiose-containing medium leads to a Cel⁺ phenotype in *L. lactis* MG1363. An *L. lactis* MG1363 derivative lacking the two main cellobiose transporters (PtcBA-CelB and PtcBAC), strain MGΔ*ptcC*Δ*celB*, does not grow in a chemically defined medium with cellobiose (CDM-Cel) as the sole carbon source for around 24 h. However, after a very long lag phase, an increase in culture optical density can be observed. Recently, van der Meulen et al. described Limg_0963, which is annotated as a cellobiose-specific EIIc PTS component. A cellobiose-inducible small RNA (sRNA), LLMGnc_147, enhances expression of this protein (15). To find out whether expression of Limg_0963 was responsible for the growth of MGΔ*ptcC*Δ*celB* in CDM-Cel, *limg_0963* was deleted in MGΔ*celB*Δ*ptcC* and growth of the triple mutant was monitored in CDM-Cel. After a lag phase of 24 h the strain started growing, demonstrating that Limg_0963 does not play a role in the observed phenomenon (Fig. 1).

When MGΔ*celB*Δ*ptcC*Δ*limg_0963* was streaked out on a rich M17 agar with cellobiose, two types of colonies appeared: small ones which were visible after overnight incubation and presumably grew on residual carbon sources in M17 and large ones (Fig. 1). The latter had (re)gained the ability to metabolize cellobiose and became distinguishable only after 24 h of incubation. When large colonies were restreaked on new M17-Cel agar plates, they retained the large colony Cel⁺ phenotype. When small colonies were restreaked, they again diversified into two phenotypes and formed small and large colonies. The inheritable Cel⁺ phenotype suggested that genetic changes were involved. In order to identify these changes, the whole genomes of colony-purified Cel⁺ strains with different genetic backgrounds (MGΔ*celB* and MGΔ*celB*Δ*ptcC*Δ*limg_0963*) were sequenced and compared to that of the parental strain, MG1363. For the purpose of the current study, a cellobiose-positive variant of MG1363 with a *celB* promoter up-mutation (Pcel⁺) was used (11). Cel⁺ isolates were picked from M17-Cel agar plates with *L. lactis* MGΔ*celB* and MGΔ*celB*Δ*ptcC*Δ*limg_0963*. The number of Cel⁺ colonies suggested that the genetic changes had occurred at very

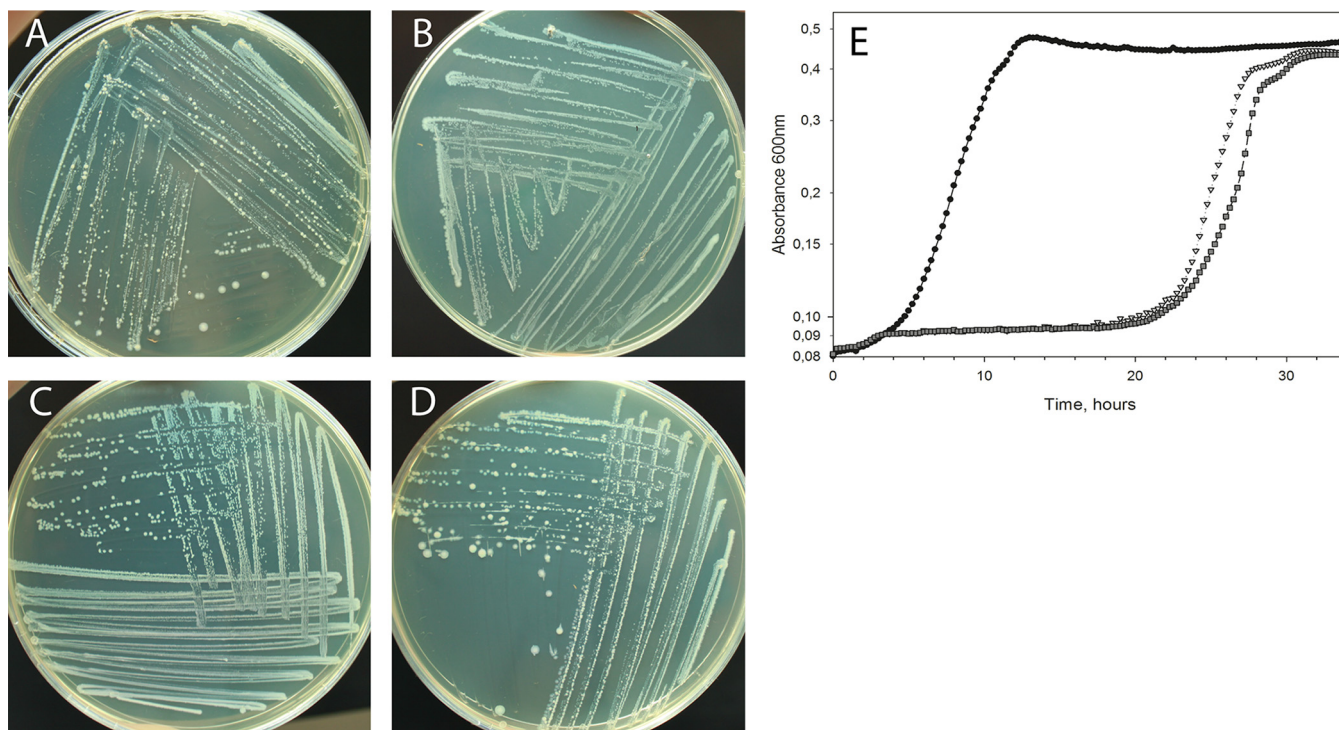


FIG 1 (A to D) Growth of *L. lactis* MGΔ*celB*Δ*ptcC*Δ*llmg_0963* on M17 agar with or without cellobiose for 24 h. (A) Two types of colonies are formed by MGΔ*celB*Δ*ptcC*Δ*llmg_0963* on M17-Cel. (B) Growth of the same strain on solidified M17. (C) Only large colonies appear when a large colony from panel A is restreaked on solidified M17-Cel. (D) Two types of colonies appear when a small colony from panel A is restreaked on solidified M17-Cel. (E) Growth of *L. lactis* strains in CDM-Cel. Circles, MG1363 carrying a P_{cel} promoter up-mutation (growth rate [k] = 0.26 h⁻¹); triangles, MGΔ*celB*Δ*ptcC* (k = 0.28 h⁻¹); squares, MGΔ*celB*Δ*ptcC*Δ*llmg_0963* (k = 0.2 h⁻¹). Growth curves show means for 3 replicates.

high frequencies, which pointed to possible involvement of a mobile genetic element(s).

Analyses of genome sequences of the Cel⁺ isolates revealed that, indeed, some genetic changes had occurred. Two types of IS elements, namely, IS981 and IS905N, had integrated in *llmg_pseudo_39* of MGΔ*celB* and MGΔ*celB*Δ*ptcC*Δ*llmg_0963*, respectively. In fact, the frameshift that was annotated in the sequence of *llmg_pseudo_39* of *L. lactis* MG1363 (4) was not present in any of our sequenced strains, meaning that the gene is intact in these strains. The *L. lactis* MG1363 strain sequenced by Linares et al. did not contain the frameshift either, but the annotation of the gene was not changed (12). Since the gene encodes an intact protein of 348 amino acids in our strains, throughout this paper we label it *llmg_1239*, not *pseudo_39*. Sequence analyses of the PCR-amplified chromosomal regions of many other Cel⁻ and Cel⁺ isolates revealed that the IS elements had integrated in *llmg_1239* of Cel⁺ isolates only while all Cel⁻ strains possessed an intact gene. The gene, which is identical to LLNZ_06380 of *L. lactis* NZ9000, codes for a transcriptional regulator of the LacI family and lies upstream of a putative plant sugar utilization gene cluster (Fig. 2). *L. lactis* NZ9000 is a derivative of MG1363 in which two genes (*nisRK*) were integrated into the chromosome as a part of the nisin-inducible gene expression system (16).

Besides IS integration, one change in the promoter region of the catabolite control protein CcpA-encoding gene was discovered in the chromosomes of two Cel⁺ isolates. A transversion from an A to a T had occurred in the -10 region of the promoter, changing it from TATAAT into TATATT. The mutation might affect transcription of the *ccpA* gene, lowering the number of regulator molecules, since the changed -10 box is less similar to the consensus -10 sequence.

Disruption of *llmg_1239* leads to a Cel⁺ phenotype of *L. lactis* MG Cel⁻. Closer analyses of the MGΔ*celB*Δ*ptcC*Δ*llmg_0963* Cel⁺ genome sequence showed that the integration of IS905N did not result in a premature translation stop but changed the last

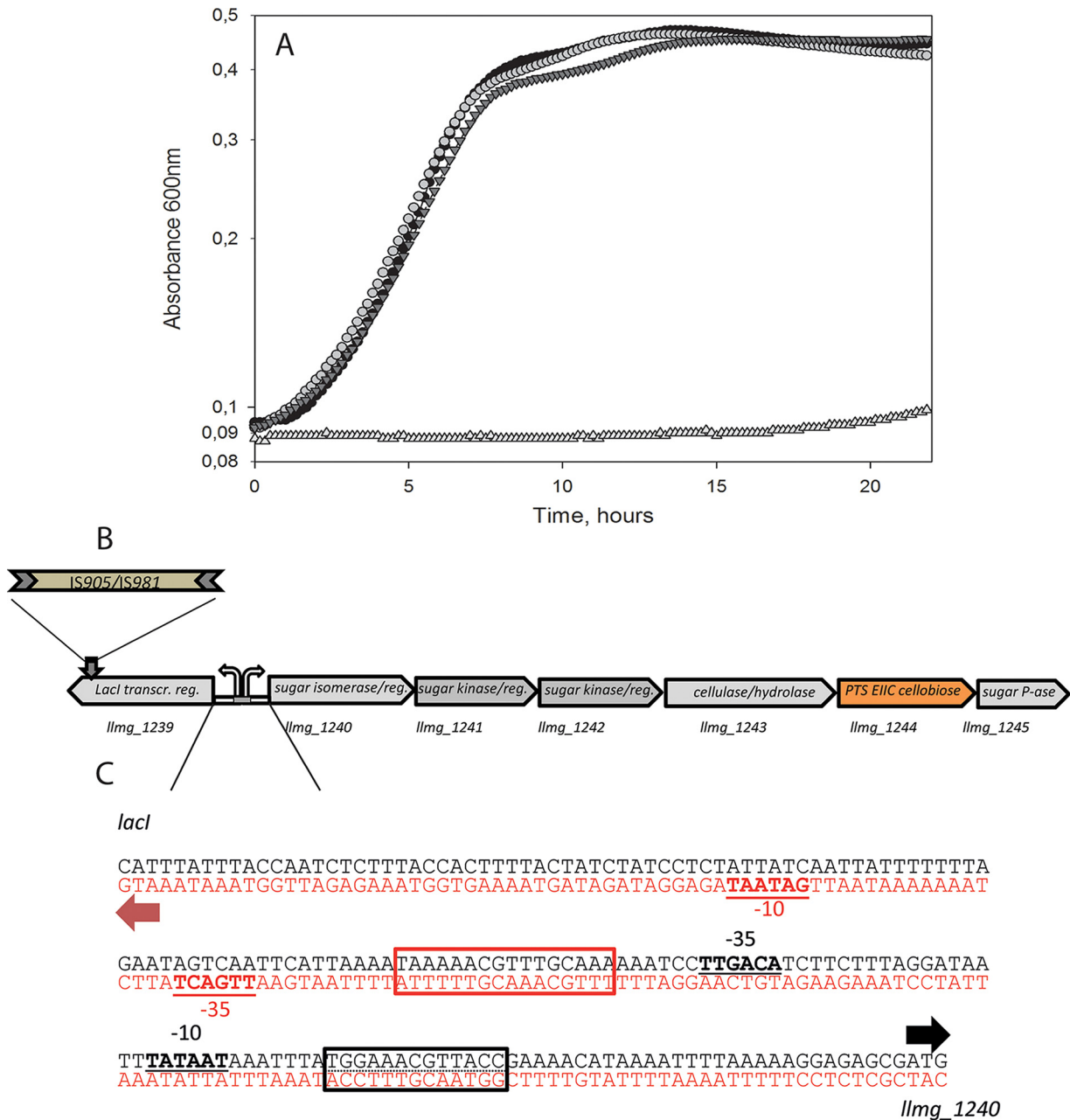


FIG 2 Integration of IS/GFP into *limg_1239* and growth of *L. lactis* derivatives in CDM-Cel. (A) Black circles, MG1363 carrying a *P*_{cel} promoter up-mutation ($k = 0.25 \text{ h}^{-1}$); light gray circles, MGΔ*celB*Δ*ptcC*Δ*limg_0963 limg_1239::IS905* (M-IS) ($k = 0.24 \text{ h}^{-1}$); dark gray inverted triangles, MGΔ*celB*Δ*ptcC*Δ*limg_0963 limg_1239::gfp* (M-GFP) ($k = 0.23 \text{ h}^{-1}$); light gray triangles, MGΔ*celB*Δ*ptcC*Δ*limg_0963* ($k = 0.0039 \text{ h}^{-1}$). Growth curves show means for 3 replicates. (B) Genetic cluster *limg_1240-45* containing genes with predicted functions in sugar utilization and the predicted transcriptional repressor gene *limg_1239*, located upstream. transcr. reg, transcription regulator; P-ase, phosphatase. (C) *limg_1239-limg_1240* intergenic region. Translation start sites (ATG) are indicated by arrows, -10 and -35 sequences are underlined, and predicted operator sequences to which Lacl family regulators can bind are shown in boxes. Black box, CcpA; red box, GalR of *E. coli* (note that *L. lactis* MG1363 does not possess GalR). The prediction was made using <http://genome2d.molgenrug.nl>.

seven amino acids of the C terminus of the putative Lacl family transcriptional regulator. These substitutions occurred in the putative ligand molecule binding domain. The changes in this domain, which most likely binds sugar, might affect the function of the regulator. MGΔ*celB* Cel⁺ colonies possessed a truncated version of the same regulator after the integration of IS981 into *limg_1239*. The IS981 integration introduced a stop codon, leading to a truncated version of the putative regulator of 290 amino acids instead of the full-length 348-amino-acid protein. Again, the insertion only disrupted

TABLE 1 Relevant genes that were differentially expressed in M-GFP and M-IS in comparison with the WT^a

Locus	Gene	Predicted function	Fold difference from WT	
			M-GFP	M-IS
<i>llmg_1242</i>	<i>llmg_1242</i>	Sugar kinase/transcriptional regulator	52.5	20.7
<i>llmg_1240</i>	<i>llmg_1240</i>	Sugar isomerase/transcriptional regulator	22.0	12.8
<i>llmg_1244</i>	<i>llmg_1244</i>	Cellobiose-specific PTS EIIc	16.4	9.3
<i>llmg_1245</i>	<i>llmg_1245</i>	Sugar phosphatase	20.8	8.3
<i>llmg_1241</i>	<i>llmg_1241</i>	Sugar kinase/transcriptional regulator	14.3	7.3
<i>llmg_1243</i>	<i>llmg_1243</i>	Cellulase/hydrolase	8.1	2.9
<i>llmg_0437</i>	<i>ptcB</i>	Cellobiose-specific PTS EIIb	2.2	
<i>llmg_0438</i>	<i>ptcA</i>	Cellobiose-specific PTS EIIa	3.4	
<i>llmg_0439</i>	<i>llmg_0439</i>	LacI family transcriptional regulator	1.9	
<i>llmg_2311</i>	<i>arcD1</i>	Arginine/ornithine antiporter	9.1	2.2
<i>llmg_2310</i>	<i>arcC1</i>	Carbamate kinase	4.9	2.2
<i>llmg_2312</i>	<i>arcB</i>	Ornithine carbamoyltransferase	4.3	
<i>llmg_2309</i>	<i>arcC2</i>	Carbamate kinase	2.7	
<i>llmg_2313</i>	<i>arcA</i>	Arginine deiminase	2.4	
<i>llmg_1559</i>	<i>flpA</i>	Transcriptional regulator, Crp/Fnr family		4.9
<i>llmg_0456</i>	<i>pgmB</i>	Phosphoglucomutase	3.0	
<i>llmg_0452</i>	<i>treR</i>	Trehalose operon transcriptional repressor	2.1	
<i>LLMGnc_172</i>		sRNA	13.3	
<i>LLMGnc_008</i>		sRNA		3.2
<i>LLMGnc_004</i>		6S RNA	2.4	
<i>LLMGnc_131</i>		sRNA	2.4	
<i>LLMGnc_089</i>		sRNA	2.3	
<i>LLMGnc_133</i>		sRNA	2.1	
<i>LLMGnc_080</i>		sRNA	2.1	
<i>LLMGnc_139</i>		sRNA	2.1	
<i>LLMGnc_150</i>		sRNA	2.0	
<i>LLMGnc_002</i>		sRNA	2.0	
<i>LLMGnc_026</i>		sRNA	2.0	
<i>LLMGnc_032</i>		sRNA		2.5
<i>LLMGnc_099</i>		sRNA		2.2
<i>LLMGnc_044</i>		sRNA		2.2
<i>LLMGnc_173</i>		sRNA		2.2
<i>LLMGnc_150</i>		sRNA		2.2
<i>LLMGnc_090</i>		sRNA		2.1
<i>LLMGnc_134</i>		sRNA		2.1
<i>LLMGnc_013</i>		sRNA		2.0
<i>LLMGnc_056</i>		sRNA		2.0
<i>LLMGnc_045</i>		sRNA		2.0
<i>llmg_0190</i>	<i>bgIS</i>	P-β-glucosidase BglS		-2.2
<i>llmg_0188</i>	<i>llmg_0188</i>	Function unknown	-5.1	-5.1
<i>llmg_0189</i>	<i>llmg_0189</i>	Function unknown	-6.6	-9.6
<i>llmg_0186</i>	<i>llmg_0186</i>	Surface protein/glycosylhydrolase	-22.7	-12.4
<i>llmg_0187</i>	<i>celB</i>	Cellobiose-specific PTS EIIc	-14.4	-20.8
<i>tnp904</i>	<i>tnp904</i>	Transposase	-4.0	-4.8
<i>tnp905</i>	<i>tnp905</i>	Transposase	-4.2	-4.1
<i>llmg_0550</i>	<i>tnp891</i>	Transposase	-10.3	-3.2
<i>llmg_0775</i>	<i>ccpA</i>	Catabolite control protein A	-1.4	-2.8
<i>llmg_1165</i>	<i>llmg_1165</i>	ABC-type sugar transport system	-17.0	-7.3
<i>llmg_2539</i>	<i>gapB</i>	Glyceraldehyde 3-phosphate dehydrogenase	-9.5	-10.9
<i>llmg_0099</i>	<i>llmg_0099</i>	Ribosomal protein	-3.5	-16.5
<i>llmg_1208</i>	<i>llmg_1208</i>	Ribosomal protein		-15.0
<i>llmg_2356</i>	<i>llmg_2356</i>	Ribosomal protein	-6.8	-9.1
<i>llmg_2357</i>	<i>llmg_2357</i>	Ribosomal protein	-8.4	-9.2
<i>llmg_2374</i>	<i>llmg_2374</i>	Ribosomal protein	-6.4	-6.6
<i>llmg_2366</i>	<i>llmg_2366</i>	Ribosomal protein	-5.5	
<i>llmg_2365</i>	<i>llmg_2365</i>	Ribosomal protein	-5.3	-5.3
<i>llmg_1493</i>	<i>llmg_1493</i>	Ribosomal protein	-6.0	-9.5
<i>llmg_2354</i>	<i>rpoA</i>	RNA polymerase	-6.0	-8.1
<i>llmg_2353</i>	<i>llmg_2353</i>	Ribosomal protein	-4.1	-10.1

^aOnly significantly expressed genes are included (Bayesian $P < 0.001$).

dehalogenase-hydrolase activity, similar to trehalose/sucrose phosphatase. All these genes were upregulated around 3- to 20-fold in *llmg_1239::IS905*-carrying and 8- to 52-fold in *llmg_1239::gfp*-carrying Cel⁺ strains. Two cellobiose-specific EII cytoplasmic AB components containing *ptcBA* were upregulated in the M-GFP strain compared to their expression in the WT. The most highly upregulated gene cluster in the WT relative to its expression in M-IS and M-GFP was the *celB* operon (up to 20- to 22-fold), confirming its importance in cellobiose uptake in this strain.

Besides sugar utilization genes, the operon encoding the arginine deiminase pathway enzymes was overexpressed in M-GFP and M-IS cells. A small RNA molecule that is the product of the *argR* transcript, LLMGnc_172 (15), was also detected in the transcriptome of the M-GFP strain. sRNA LLMGnc_147, which was previously reported to be activated in the presence of cellobiose (15), was not differentially expressed in the strains compared.

Another regulatory RNA abundant in M-GFP cells was 6S RNA. This regulatory small RNA is known to be upregulated in *L. lactis* MG1363 during the exponential growth phase when galactose or cellobiose is the carbon source (15). It is interesting that although many genes show similar expression patterns in M-IS and M-GFP strains, different sets of sRNAs seem to be present in the two strains. Presumably, changes in sRNA expression are more dynamic and their expression differences are caused by minor variances in RNA sampling time.

Cellobiose utilization in *L. lactis* MG1363. A *cre* site (TAAAAACGTTTGCAAAAAT) is present in the *llmg_1239-1240* intergenic region, suggesting global regulation of the operon by a LacI family protein, carbon catabolite repression (CCR) mediator CcpA (prediction performed with an online tool [<http://genome2d.molgenrug.nl>]) (Fig. 2). Regulation of catabolic genes and operons by CCR is very common in *L. lactis* and ensures dominant utilization of glucose when it is present in the growth medium. On the other hand, a weak basal expression of the repressor-encoding gene *llmg_1239* can be observed in transcriptomics and RNA-seq data even in the presence of glucose (15, 17, 18). Another LacI family operator-like palindrome, TGGAAACGTTACCA, was identified downstream of the *cre* site in the *llmg_1239-1240* intergenic region. The operator is similar to the consensus sequence recognized by GalR of *Escherichia coli* (18; predicted using the tool at <http://genome2d.molgenrug.nl>). Typical LacI family repressors bind their operator sites as dimers or tetramers in the absence of the effector molecules. The binding of the effector molecule (very often a carbohydrate) reduces the DNA binding affinity, releasing the regulator from the operator and allowing transcription (19).

Based on the data generated in this study, we propose that three cellobiose transporters can be activated in *L. lactis* MG1363 (Fig. 4).

DISCUSSION

The *L. lactis* subsp. *cremoris* MG1363 genome contains multiple silent gene clusters and pseudogenes with predicted functions in plant sugar utilization, as a heritage of its plant-related origin (4). Nowadays many of the *L. lactis* subsp. *cremoris* strains are adapted to a milk environment and are used for industrial fermentations. It has been suggested that the high number of pseudogenes could indicate ongoing specialization of the strains to their current niche, with constant degeneration of dispensable genes and their encoded functions (5, 7). Genome decay was reproduced in an experimental evolution study through cultivation of the plant isolate *Lactococcus lactis* subsp. *lactis* KF147 in milk for 1,000 generations (20). The outcome of the experiment was isolation of strains which had lost or had downregulated plant polymer utilization genes and gene clusters (20).

Here we describe a “reverse evolution” experiment: a reactivation of a silenced plant sugar utilization pathway when selection pressure to utilize cellobiose is applied. Disruption of the repressor gene *llmg_1239* by an IS element allowed expression of the otherwise silent cellobiose transporter *llmg_1244* and led to survival of mutant strains on cellobiose. LacI family transcriptional regulators homologous to that encoded by

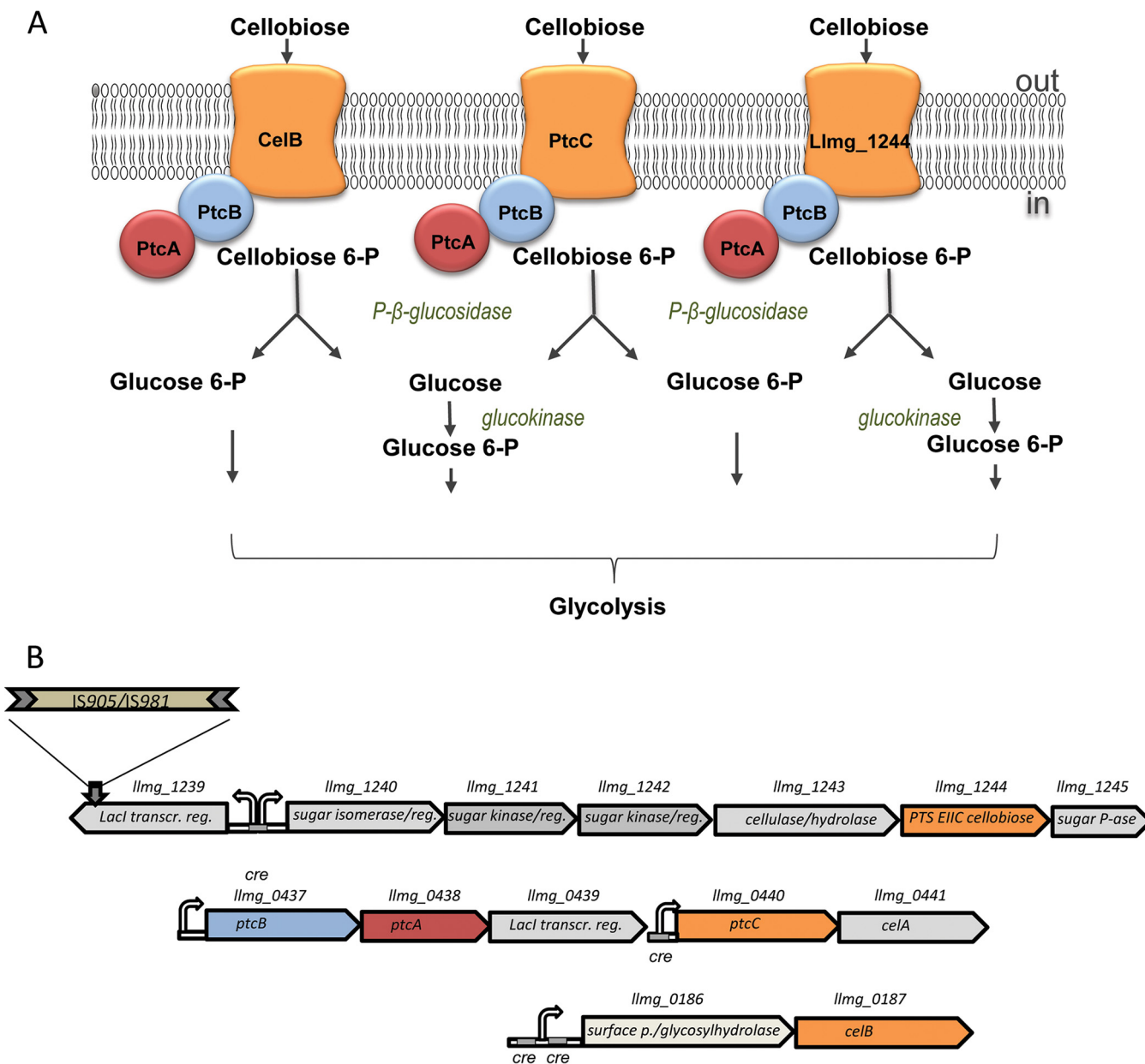


FIG 4 Cellobiose-specific uptake and metabolism in *L. lactis* MG1363. (A) Three cellobiose-specific PTSs and subsequent metabolic routes for degradation of this sugar. P, phosphate. (B) Gene clusters containing cellobiose transporter genes. Cellobiose-specific EIIc component Limg_1244 is expressed only when *limg_1239*, coding for a transcriptional repressor, is disrupted.

limg_1239 are present in the genomes of only a few *L. lactis* strains: NZ9000 and KW2 of *L. lactis* subsp. *cremoris* (gene sequence homology of 99%) and IO-1 of *L. lactis* subsp. *lactis* (gene sequence homology of 88%; KEGG database). While *L. lactis* NZ9000 is of dairy origin, KW2 is isolated from fermented corn and the xylose-utilizing strain IO-1 was first isolated from a drain pit of a kitchen sink (16, 21, 22). The transcriptional regulator is homologous to PurR and GalR from *Cedecea*, *Klebsiella*, and *Listeria*. It does not resemble any of the other cellobiose metabolism regulators, such as Limg_0439 (LacI), CcpA, or ClaR (DexR) of *L. lactis* or CelR of *Streptococcus*. Due to the lack of clear homology to any other transcriptional regulator, we propose to designate Limg_1239 CcIR, for cellobiose cluster repressor.

It remains unclear to which effector molecule CcIR/Limg_1239 responds. Apparently, the absence of glucose and the presence of cellobiose in the medium were

not the right cues for the relief of repression of the catabolic gene cluster *llmg_1240-45*. The reason for such tight repression could be the lack either of the proper effector molecule or of an additional coregulator protein that might have been lost during evolution. Alternatively, the ability of the regulator to bind the inducer may have been lost.

A functional PTS EII consists of three components. Once the sugar molecule is imported via the membrane-spanning EII component C, it needs to be phosphorylated by the cytosolic EIIB subunit. Phosphorylation of sugar lowers the affinity of EIIC for it, and the molecule is released to the cytosol (23). The EIIB component receives its phosphoryl group from an EIIA protein. Both of them are sugar specific and usually interact with only one EIIC. As the *llmg_1240-45* cluster contains no genes potentially encoding EIIAB subunits, *Lmg_1244* presumably works with EII components of another PTS. PtcAB are promiscuous EIIAB proteins and phosphorylate carbohydrates imported via PtcC, CelB, and, presumably, *Lmg_0963*. All of these EIIC components are annotated as cellobiose specific, although they can import other sugars (glucose and galactose) as well (14, 15). The ability of PtcBA to interact with more than one EIIC component is very unusual for bacterial PTSs. Genes coding for PtcBA form a separate cluster and can be transcribed independently from their membrane-integrated partners (Fig. 4). The transcriptome data showed an around 2- to 3-fold upregulation of *ptcBA* in the M-GFP strain compared to their expression in the WT strain. We propose that the full novel cellobiose PTS consists of *Lmg_1244PtcBA*.

Possibly because of the change from the consensus -10 box TATAAT into a less perfect TATATT in its promoter, *ccpA* expression is slightly lower in the M-GFP and M-IS strains than it is in the WT strain. Presumably, related to that, expression of the *arc* and the trehalose (*pgm* and *treR*) operons and that of the 6S RNA are higher in M-GFP and M-IS. Cellobiose has not been shown to cause CCR, but it consists of two glucose molecules and is metabolized by phospho (P)- β -glucosidases rather quickly in Cel⁺ strains (Fig. 2), making it plausible that some level of catabolite repression on cellobiose remains. In accordance with higher *ccpA* expression, glycolysis (*gapB*) and protein synthesis (RNA polymerase RpoA- and ribosomal protein-encoding genes) are more intense in the WT strain. To evaluate the role of the *PccpA* mutation in the strength of CCR, the level of induction of *Parc* and *PackA* in the presence of glucose was studied in *L. lactis* MG1363 (WT) and *MG Δ celB Δ ptcC* using promoter-*gfp* fusion constructs. *Parc* and *PackA* are strong promoters that contain *cre* sites and are repressed by CcpA in *L. lactis* (24). Fluorescence intensities driven from *Parc* and *PackA* differ between the WT and *MG Δ celB Δ ptcC*, especially in glucose-containing media (where CCR is active), although the difference is not very pronounced (data not shown). More studies are needed in order to elucidate the role of the change in the sequence of *PccpA*.

WT cells also strongly express a putative sugar ABC permease, *Lmg_1165*. The gene for this transporter is located in a putative sugar utilization cluster, *llmg_1164-aguA*. The cluster codes for two ABC transporters (*Lmg_1164* and *Lmg_1165*), two putative endoglucanases, polysaccharide (chitin and xylan) deacetylase, and a putative α -glucuronidase, *AguA*. The role of this gene cluster in *L. lactis* MG1363 has not yet been studied. It is not clear whether the expression of *llmg_1165* in WT cells is the result of the *Lmg_1239* regulation or is an indirect effect of possible slight variations in growth rates or CcpA levels among the strains. Changes in expression were observed in a set of sRNAs in the strains studied. Unfortunately, little is known about the exact functions of the regulatory RNAs in *L. lactis*. The housekeeping 6S RNA, which was upregulated in M-GFP, was shown to be expressed during the exponential growth phase in cells utilizing cellobiose (15). The expression of *LLMGnc_172* was associated with that of the *arc* cluster (15), which was also overexpressed in M-GFP. It is possible that the type of disruption of *llmg_1239* (by an IS905 or via *gfp* integration) is the cause of differences in sRNA expression.

The catabolic cluster *llmg_1240-45* and the repressor gene seem to have been acquired by horizontal gene transfer. The *Lmg_1244* protein sequence shows closest similarity to a lichenan-specific EIIC component, LicC, or a cellobiose-specific CelB from

various *Listeria*, *Enterococcus*, and *Lactobacillus* strains. Moreover, comparison of *L. lactis* MG1363 PTS components with *lmg_1244* demonstrates that it is not related to PtcC and CelB of this organism (32% and 29% similarity, respectively). Lactococci are in contact with other bacterial species in both industrial and natural environments. Bacterial genomes can be altered not only by point mutations or genome rearrangements but also by acquisition of foreign DNA (1). Although *L. lactis* strains possess elements of the competence machinery, it is not functional (25). Transduction by phages and transfer mediated by transposons or conjugative plasmids thus serve as the main sources of foreign DNA (3, 26). Integration of the acquired DNA can occur via homologous recombination of common IS elements.

IS elements play an important role in the evolution of lactococci and other bacteria. They facilitate DNA rearrangements and are responsible for creation of new genetic variants with selective advantages under certain environmental conditions. During adaptation of lactococci to the milk environment, IS elements most likely helped to acquire and rapidly disseminate such traits as lactose metabolism, casein protease activity, nisin production and immunity, and bacteriophage resistance (8). Horizontal gene transfer is obviously an essential mechanism for lactic acid bacteria (LAB) to adapt and therefore survive in their respective environments. *L. lactis* MG1363 possesses 71 copies in a total of 11 different types of IS elements, of which *IS981* (27) and *IS905* (28) are the most abundant (14 copies and 16 copies, respectively) (4). *IS981* has been described to be involved in adaptation of *L. lactis* MG1363 and IL1403 to unfavorable conditions during starvation or oxidative- or bacteriocin-induced stress (9, 10, 29). A recombination event between two *IS905* elements has been shown to be responsible for a large chromosomal inversion in *L. lactis* NCDO763, although no resulting phenotypic changes could be observed in comparison to MG1363. The two *IS905* elements between which the inversion had occurred were fixed in these *L. lactis* strains, presumably because of their positive effects for the cell: one of the IS elements provided a -35 sequence for *psp* (pyrrolidone carboxyl peptidase gene), and the other one disrupted a gene for a phage terminase subunit and thus protected the cell from phage-caused lysis (30). The chromosomal region of *L. lactis* MG1363 harboring the cluster *lmg_1240-45* is surrounded by transposases and helper proteins for both *IS981* and *IS905*; e.g., *lmg_1234* and *lmg_1264* code for transposases for *IS981*, while *lmg_1319* specifies a transposase for *IS905*. The abundance of mobile elements in the vicinity of the *lmg_1240-45* cluster could have influenced the high frequency of their integration into *lmg_1239*. Since the integrants acquire a substantial growth advantage in cellobiose-containing medium, they are rapidly selected for.

The fact that *lmg_1239* contains a frameshift and is annotated as a pseudogene in the first sequenced *L. lactis* MG1363 strain (4) may not be coincidental. Silencing of the repressor gene by such a mutation could be another mechanism to activate the *lmg_1240-45* cluster. The frameshift was identified in a stretch of adenines (6 in a row) making the region prone to slipped-strand mispairing during replication. It is also known that this MG1363 derivative did not possess the *PcelB* activating mutation, meaning that it could not use PtcBA-CelB for the import of cellobiose. If this Cel⁻ strain has been exposed to cellobiose before sequencing, it could have acquired the *lmg_1239* frameshift mutation that helped it to survive on this disaccharide. None of the *L. lactis* MG1363 strain stocks sequenced in our laboratory harbored the frameshift in *lmg_1239*. The same was reported for the isolate sequenced by Linares et al. (12). Strains evolve with time, and many processes might have changed in what we call "model strains" because of long laboratory propagation. It is also clear that these model strains vary per laboratory. We show here that it is important to resequence *lmg_1239* when studying the metabolism of cellobiose or other sugars in *L. lactis* MG1363 and its derivatives. While it was determined in this study that the silent sugar utilization cluster can be activated when cellobiose is present in the environment, the real inducer molecule that binds the repressor CclR remains to be elucidated.

TABLE 2 Strains and plasmids used in this study

Strain or plasmid	Description	Source or reference
<i>E. coli</i> DH5 α	F ⁻ ϕ 80 <i>lacZ</i> Δ M15 Δ (<i>lacZYA-argF</i>)U169 <i>rec1A end1A hsdR17 gyrA96 supE44 thi-1 relA1</i>	Invitrogen (Carlsbad, CA)
<i>L. lactis</i> strains		
MG1363 (WT)	<i>L. lactis</i> subsp. <i>cremoris</i> plasmid-free derivative of NCDO712 carrying a <i>PceI</i> up-mutation	11
MG Δ <i>celB</i>	MG1363 carrying a chromosomal deletion of <i>celB</i>	11
MG Δ <i>celB</i> Δ <i>ptcC</i>	MG1363 carrying chromosomal deletions of <i>celB</i> and <i>ptcC</i>	Gift from B. Martinez
MG Δ <i>celB</i> Δ <i>ptcC</i> Δ <i>llmg_0963</i>	MG1363 Δ <i>celB</i> Δ <i>ptcC</i> carrying a chromosomal deletion of <i>llmg_0963</i>	This study
MG Δ <i>celB</i> Δ <i>llmg_1239::IS981</i>	MG1363 Δ <i>celB</i> with <i>IS981</i> integrated in <i>llmg_1239</i>	This study
MG Δ <i>celB</i> Δ <i>ptcC</i> Δ <i>llmg_0963</i>	MG Δ <i>celB</i> Δ <i>ptcC</i> Δ <i>llmg_0963</i> with <i>IS905</i> integrated in <i>llmg_1239</i>	This study
<i>llmg_1239::IS905</i> (M-IS)		
MG Δ <i>celB</i> Δ <i>ptcC</i> Δ <i>llmg_0963</i>	MG Δ <i>celB</i> Δ <i>ptcC</i> Δ <i>llmg_0963</i> with <i>PackA-gfp</i> integrated in <i>llmg_1239</i>	This study
<i>llmg_1239::gfp</i> (M-GFP)		
Plasmids		
pCS1966	Integration vector for <i>L. lactis</i>	29
pCS1966-0963'	pCS1966 containing <i>llmg_0963</i> deletion construct	This study
pSeudo39	Integration vector for <i>L. lactis</i> carrying the flanking regions of <i>llmg_1239</i>	Gift from J. Siebring
pSeudo39 <i>PackA-gfp</i>	pSeudo39 containing <i>PackA-gfp</i>	40

MATERIALS AND METHODS

Microbial strains used and growth conditions. *L. lactis* MG1363 (31) derivatives (Table 2) were grown as standing cultures at 30°C in M17 broth (Difco, Sparks, MD) or in chemically defined medium (CDM PC) (32) supplemented with 1% cellobiose or 0.5% glucose. M17 agar plates were prepared by adding 1.5% (wt/vol) agar to M17 supplemented with either glucose (0.5%) or cellobiose (1%). When appropriate, erythromycin (Sigma-Aldrich, St. Louis, MO) was used at 1 μ g ml⁻¹. *E. coli* DH5 α (Invitrogen, Carlsbad, CA) was used as a cloning host and was grown in LB medium at 37°C or on LB medium solidified with 1.5% (wt/vol) agar. For plasmid selection, 150 μ g ml⁻¹ of erythromycin was added.

To study the growth of various *L. lactis* strains, the strains were grown in 0.2 ml of CDM supplemented with 1% cellobiose in 96-well microtiter plates at 30°C. Growth was monitored by measuring the OD₆₀₀ with an Infinite 200 PRO 16 microplate spectrophotometer (Tecan Group Ltd., Mannedorf, Switzerland).

General DNA techniques. DNA manipulations were executed essentially as described previously (33). Plasmid DNA and PCR products were isolated and purified using the High Pure plasmid isolation kit (Roche Applied Science, Mannheim, Germany) according to the manufacturer's instructions.

Restriction enzymes, T4 DNA ligase, and *Taq* DNA polymerase were obtained from Thermo Fisher Scientific Baltics (Vilnius, Lithuania) and used according to the supplier's instructions. Phusion DNA polymerase was purchased from New England Biolabs (Ipswich, MA). PCR were performed in a Bio-Rad thermal cycler (Hercules, CA) using *L. lactis* MG1363 chromosomal DNA as the template.

Construction of the *L. lactis* deletion strain. The flanking regions of *llmg_0963* were amplified using Ko09631F (AAAATTCTAGAAATTATGGTATGCATTATAG)/Ko09632R (TGCATGGATCCTCTTTAGATTCACTCCTTAAAC) and Ko09633F (TGCATGGATCCGAATTAGAGGTTTCAGAAAC)/Ko09634R (TTTTAGGTACCAGACTTCTACTGACAGATTC). These fragments were ligated into pCS1966 (34, 35) via *Xba*I/*Bam*HI and *Bam*HI/*Kpn*I restriction sites. The resulting vector was designated pCS1966-0963'. The pCS1966 derivative was obtained and maintained in *E. coli* DH5 α .

Vector pCS1966-0963' was introduced in *L. lactis* MG1363 Δ *celB* Δ *ptcC* (a kind gift from B. Martinez) via electroporation (36); cells in which the two-step homologous recombination event had occurred were selected by growing them on selective SA medium plates (37) supplemented with 20 μ g ml⁻¹ of 5-fluoroorotic acid hydrate (29, 32). The obtained strain was labeled MG1363 Δ *celB* Δ *ptcC* Δ 0963. The chromosomal structure of the deletion strain was confirmed by PCR analysis and sequencing of the modified regions.

Construction of the *gfp* integration strain. *L. lactis* plasmid pSeudo39*PackA-gfp* was constructed by introducing *PackA* and *sfgfp*(Bs) (38) into pSeudo39 vector carrying two flanking regions of *llmg_1239* (a kind gift from J. Siebring) via *Sma*I/*Xho*I and *Xho*I/*Bam*HI restriction sites, respectively. The *PackA* promoter fragment was obtained by PCR using primer pair 5'-GCATCCCGGGATCTTTATGGAAGAATTAC-3'/5'-CGATCTCGAGTTTGGTCATGTTAATAAAC-3' (underlined sequences represent the *Sma*I and *Xho*I restriction sites, respectively) and the chromosomal DNA of *L. lactis* MG1363 as a template. The resulting vector was integrated into the chromosome of MG1363 Δ *celB* Δ *ptcC* Δ 0963. The strain obtained after a two-step recombination event via the *llmg_1239*-homologous regions (as described above) was designated MG1363 Δ *celB* Δ *ptcC* Δ 0963 *llmg_1239::gfp* (M-GFP).

Transcriptome analyses. Transcriptome analysis was performed using full-genome *L. lactis* MG1363 DNA microarrays (Agilent Technologies, Santa Clara, CA) as described previously (39). Single colonies of the WT, M-GFP, and M-IS were used to inoculate 10 ml of M17 supplemented with 1% cellobiose. After overnight growth, the cultures were diluted with fresh medium 10 \times and harvested at the mid-exponential growth phase, at an OD₆₀₀ of 0.6. RNA from M-GFP and M-IS was compared to RNA from MG1363 (WT). DNA Microarray slides were scanned with a Genepix 4200 laser scanner (Molecular

Devices, Sunnyvale, CA). Slide images were analyzed using GenePix Pro v.6.0 software. Background subtraction and LOWESS (locally weighted scatterplot smoothing) normalization were done using the standard routines provided by GENOME2D software available at <http://genome2d.molgenrug.nl/index.php/analysis-pipeline>. A gene was considered differentially expressed when the Bayesian *P* value was <0.001, and a fold change cutoff of 2.5 was applied.

Accession number(s). Gene expression data were deposited in the GEO database under accession number [GSE103707](https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE103707).

ACKNOWLEDGMENTS

We thank A. Ravi and M. Tufano for their help with the transcriptome analyses. We thank B. Martinez (IPLA-CSIC, Spain) for kindly providing *L. lactis* MG1363Δ*celB*Δ*ptcC*. We thank J. Siebring for kindly providing plasmid pSeudo39.

This work was supported by a grant from NWO-STW Technology Foundation (project number 10619).

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