Carbon catabolite repression (CCR) of many genes in *Bacillus subtilis* is caused by the availability of glucose or other rapidly metabolized carbon sources during growth. The regulatory process involves several proteins including HPr, Crh, HPr kinase, and CcpA. The HPr protein, encoded by the *ptsH* gene, functions in CCR as well as in the phosphoenolpyruvate-sugar phosphotransferase system (PTS). Crh (catabolite repression HPr) is a protein homologous to HPr that functions in the PTS, glycolytic intermediates such as fructose-1,6-bisphosphate (F1,6BP) but not in the PTS (13, 29). After glucose is taken up via the PTS, glycolytic intermediates such as fructose-1,6-bisphosphate activate HPr kinase (14, 19, 25). Then, HPr kinase phosphorylates HPr (14, 36) and Crh (13, 14) at a serine residue (Ser46) in an ATP-dependent manner. The seryl-phosphorylated proteins act as corepressors (11, 20, 24) by forming a complex (6) with the trans-acting CcpA (catabolite control protein A), a member of the LactoGalR family of regulatory proteins (17). The complex binds to a consensus DNA sequence, the so-called cre site (catabolite-responsive element) (18), located upstream of the target gene, where it may act either as a repressor or as an activator of transcription (18).

The key role of CcpA in carbon regulation in *B. subtilis* has been demonstrated by inactivating the *ccpA* gene, which resulted in relief of glucose catabolite repression (11, 17) or the abolition of gene activation (34, 40). Mutating Ser46 of HPr resulted in relief of glucose catabolite repression (11, 17) or the abolition of gene activation (34, 40). Mutating Ser46 of HPr resulted in relief of glucose catabolite repression (11, 17) or the abolition of gene activation (34, 40). Mutating Ser46 of HPr resulted in relief of glucose catabolite repression (11, 17) or the abolition of gene activation (34, 40). In minimal media, CitM was only expressed when citrate was present. The presence of glucose in the medium resulted in relief from CCR of *B. subtilis* (12), and no relief of glucose catabolite repression was observed in media containing glycerol, inositol, or succinate-glutamate. Studies with *B. subtilis* mutants defective in the catabolite repression components HPr, Crh, and CcpA showed that the repression exerted by all these medium components was mediated via the carbon catabolite repression system. During growth on inositol and succinate, the presence of glutamate strongly potentiated the repression of *citM* expression by glucose. A reasonable correlation between *citM* promoter activity and CitM transport activity was observed in this study, indicating that the Mg$^{2+}$-citrate uptake activity of *B. subtilis* is mainly regulated at the transcriptional level.

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The structural gene coding for CitM is organized in an operon-like structure (Fig. 1), including *citM* and a second gene, *yflN* (26), the function of which is not known. Upstream of *citM* are reading frames *citS* and *citT*, which code for a two-component system (9). The CitS-CitT two-component system is essential for the transcription of the *citM-yflN* operon. The putative CitT target sequence is believed to be located in the region between nucleotides −62 and −113, upstream of the *citM* transcriptional start point (H. Yamamoto, M. Murata, and J. Sekiguchi, Abstr. 10th Int. Conf. Bacilli, p. 71, 1999). In addition, just in front of the *citM* gene lies a sequence that matches the consensus sequence of a cre site (41, 46). The functionality of the cre site has been demonstrated in vivo (32). The location of the *citM* gene on the chromosome of *B. subtilis* suggests that expression of CitM might be under the control of the metabolic state of the cell.

Induction by citrate and inhibition by glucose of citrate uptake in *B. subtilis* have been described already 3 decades ago (42). In this study we report on the regulation of synthesis of the Mg$^{2+}$-citrate transporter of *B. subtilis*, encoded by the *citM* gene. Transcription of *citM* is stringently dependent on the pres-
ence of citrate in the growth medium and is under the control of CCR. Repression was observed in media containing several carbohydrates but also nongrowth. Experiments with wild-type and CCR mutant strains revealed the involvement of the different CCR components.

MATERIALS AND METHODS

Bacterial strains, media, and growth conditions. The B. subtilis strains used in this study are listed in Table 1. B. subtilis was grown in C medium (1) in which ammonium citrate was omitted. The C medium was supplemented with 10 mM trisodium citrate (CC medium) or 25 mM myo-inositol (CI medium) or 25 mM myo-inositol and 5 g of potassium glutamate/liter (CIE medium) or 6 g of sodium succinate and 8 g of potassium glutamate/liter (CSE medium). When strain QB5407 was grown in CI medium, 10 mM potassium glutamate was added as nitrogen source (10). Glucose or trisodium citrate was sometimes added at a concentration of 10 mM. Auxotrophic requirements were added at 20-g/ml final concentration. When appropriate, antibiotics were added at concentrations of 100 µg/ml for spectinomycin (strains QB7907 and QB5407) and 5 µg/ml for kanamycin (QB7102) and chloramphenicol (the PitM-lacZ fusion strains).

Overnight cultures of wild-type and mutant B. subtilis strains were inoculated into 20 ml of medium. The cells were grown in 100-ml flasks at 37°C on a rotary shaker operated at 150 rpm. Growth was monitored by measuring the optical density of the cultures at 660 nm (OD660) using a Hitachi U-1100 spectrophotometer. The cells were harvested by centrifugation in the exponential growth phase at an OD660 between 0.3 and 0.6 and washed once with 50 mM PIPES (piperazine-N,N'-bis[ethanesulfonic acid]), pH 6.5.

Construction of PitM-lacZ fusions. Vector pJM116 (5) contains the promotorless spoVG-lacZ gene between two fragments of the B. subtilis amyE gene and carries the cat gene from pC194 (4). The integration vector pCM160 was constructed by cloning an 819-bp-long PCR fragment of the citM region by homologous recombination, PitM, into the multiple cloning site of pJM116. The PitM promoter region including the cre site was amplified by PCR using a forward primer (5'-GAATTC-3') and a backward primer (5'-AAGCCTAA-TAACACATCC-3') that introduced an EcoRI site (boldface) and a backward primer (5'-AAGGCTTAAGATCGTCTACCAACACATCC-3') that introduced a BamHI site (boldface). Both pJM116 and the PitM PCR fragment were digested with BamHI and EcoRI and ligated to yield pCM160. The vector was constructed in Escherichia coli DH5α grown in Luria-Bertani (LB) medium (30) at 37°C. Transformants were selected by plating 50 µg of ampicillin/ml in LB agar plates. The construct was checked by restriction and DNA sequence analyses.

Wild-type and mutant B. subtilis strains were transformed with pCM160 to yield the CM series of mutants listed in Table 1. Successful integrants into the amyE locus by homologous recombination were selected by resistance against chloramphenicol. Integration into the amyE locus was confirmed by an amylase-negative phenotype of cells plated on LB agar containing soluble starch (16). Integrants contained the lacZ gene under the control of the PitM promoter region. β-Galactosidase activity was assayed qualitatively on LB agar plates containing the chromogenic substrate 5-bromo-4-chloro-3-indolyl-D-galactopyranoside (X-Gal) with or without trisodium citrate (10 mM).

Induction and glucose catabolite repression of citM expression. B. subtilis contains two known transporters for citrate, CitH and CitM. CitM is responsible for citrate-induced citrate uptake activity and transports citrate in complex with Mg2+ and other divalent metal ions (2, 3). To study the expression of CitM, the uptake activity was measured using the Ni2+-citrate complex as the substrate, which is highly specific for CitM because of the higher stability of the Ni2+-citrate complex compared to other metal ions.

RESULTS

Induction and glucose catabolite repression of citM expression. B. subtilis contains two known transporters for citrate, CitH and CitM. CitM is responsible for citrate-induced citrate uptake activity and transports citrate in complex with Mg2+ and other divalent metal ions (2, 3). To study the expression of CitM, the uptake activity was measured using the Ni2+-citrate complex as the substrate, which is highly specific for CitM because of the higher stability of the Ni2+-citrate complex.

Table 1. B. subtilis strains used in this study

<table>
<thead>
<tr>
<th>Strain</th>
<th>Genotype</th>
<th>Source [reference(s)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>168</td>
<td>trpC2</td>
<td>Microbiology, University of Groningen</td>
</tr>
<tr>
<td>SA003</td>
<td>trpC2 sacB::lacZ ptsH1</td>
<td>J. Deutscher (7)</td>
</tr>
<tr>
<td>QB7102</td>
<td>trpC2 crh::spc</td>
<td>I. Martin-Verstraete</td>
</tr>
<tr>
<td>QB5407</td>
<td>trpC2 ccpA::Tn917 spc</td>
<td>I. Martin-Verstraete (14, 34)</td>
</tr>
<tr>
<td>CM002</td>
<td>trpC2 amyE::(PitM-lacZ cat)</td>
<td>I. Martin-Verstraete (10)</td>
</tr>
<tr>
<td>CM004</td>
<td>trpC2 sacB::lacZ trpC2 amyE::(PitM-lacZ cat)</td>
<td>This study</td>
</tr>
<tr>
<td>CM006</td>
<td>trpC2 crh::spc amyE::(PitM-lacZ cat)</td>
<td>This study</td>
</tr>
<tr>
<td>CM008</td>
<td>trpC2 crh::spc amyE::(PitM-lacZ cat)</td>
<td>This study</td>
</tr>
<tr>
<td>CM010</td>
<td>trpC2 ccpA::Tn917 spc amyE::(PitM-lacZ cat)</td>
<td>This study</td>
</tr>
</tbody>
</table>
which assures that all citrate is in the complexed state (log $K_a$ is 5.4 and 3.4 for Ni$^{2+}$ and Mg$^{2+}$, respectively [28]).

*B. subtilis* 168 grown in minimal medium containing succinate and glutamate (CSE medium) showed no uptake of Ni$^{2+}$-citrate, while growth in the same medium with additional citrate resulted in significant uptake (Fig. 2). When, in addition to citrate, glucose was added, the uptake activity dropped dramatically. The experiment suggests that Ni$^{2+}$-citrate uptake activity is induced by citrate and repressed by glucose.

To correlate the Ni$^{2+}$-citrate uptake activity with the expression of the *citM* gene, the gene encoding β-galactosidase was fused behind the promoter region of *citM* (*citM-lacZ* fusion) and the construct was integrated in the *amyE* locus on the genome of *B. subtilis* 168, yielding strain CM002 (Table 1). The β-galactosidase activity of CM002 correlated with the Ni$^{2+}$-citrate uptake activity observed in the wild-type strain. No β-galactosidase activity was seen when cells were grown in the absence of citrate, while a high activity was observed in the presence of citrate. Including glucose in the medium in addition to citrate resulted again in very low β-galactosidase activity (Fig. 2, inset). The correlation between Ni$^{2+}$-citrate uptake activity and *citM* promoter activity indicates that the lack of uptake activity in cells grown in the absence of citrate or in the presence of glucose was due to the lack of expression of *citM*.

The same pattern of induction and glucose repression was observed upon growth of *B. subtilis* in minimal medium containing inositol (CI medium) and inositol plus glutamate (CIE medium) (Table 2). In the absence of citrate neither significant uptake of Ni$^{2+}$-citrate nor promoter activity was observed. In the presence of citrate both uptake and promoter activities were significantly higher in CSE medium than in CI and CIE media, while repression by glucose was most effective in the two media that contained glutamate in addition to succinate or inositol (CSE and CIE media, respectively).

Ni$^{2+}$-citrate uptake and *citM* promoter activity in different growth media. The repressive effects of various growth media on Ni$^{2+}$-citrate uptake and *citM* promoter activity were deter-

<table>
<thead>
<tr>
<th>Medium</th>
<th>Initial uptake rate (nmol/min · mg of protein)</th>
<th>β-Galactosidase activity (Miller units)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CI</td>
<td>0.01</td>
<td>0</td>
</tr>
<tr>
<td>CI + citrate</td>
<td>0.18</td>
<td>35</td>
</tr>
<tr>
<td>CI + citrate + glucose</td>
<td>0.10</td>
<td>10</td>
</tr>
<tr>
<td>CSE</td>
<td>0.01</td>
<td>0</td>
</tr>
<tr>
<td>CSE + citrate</td>
<td>0.38</td>
<td>143</td>
</tr>
<tr>
<td>CSE + citrate + glucose</td>
<td>0.01</td>
<td>11</td>
</tr>
<tr>
<td>CIE</td>
<td>0.01</td>
<td>0</td>
</tr>
<tr>
<td>CIE + citrate</td>
<td>0.17</td>
<td>30</td>
</tr>
<tr>
<td>CIE + citrate + glucose</td>
<td>0.01</td>
<td>2</td>
</tr>
</tbody>
</table>

mined by growing *B. subtilis* in citrate minimal medium supplemented with different carbon sources. Highest uptake and promoter activities were observed when cells were grown in C medium with citrate as the sole carbon and energy source (Fig. 3). Supplementing the growth medium with the carbohydrate glucose, glycercol, or inositol resulted in dramatic loss of both activities. The repressive effect was not restricted to sugars, because, though less prominent, clear decreases in both Ni$^{2+}$-citrate uptake activity and *citM* promoter activity were observed with cells grown in medium containing succinate and glutamate. The similar activities of cells grown in inositol and inositol-glutamate media suggest that the latter repression is caused by succinate. In all the growth media tested, there was a fair correlation between *CitM* transport activity and *citM* promoter activity, with the relatively low promoter activity in the glycerol medium as the exception. The differences between the two activities most likely represent other regulatory factors acting after the transcription of the gene or differences in the metabolic state of the cells (see Discussion).

The highest growth rate was observed in the medium with the lowest Ni$^{2+}$-citrate uptake activity and *citM* promoter activity. The doubling time in the media supplemented with glucose, glycercol, or inositol ranged from 120 to 140 min. The
lowest growth rate was observed in minimal citrate medium (380-min doubling time), while in CSE medium growth rate and $citM$ expression were intermediate (230-min doubling time).

In conclusion, the expression of the Ni$_2^+$-citrate uptake system is under strict control of the components of the growth medium and is repressed by other growth substrates besides glucose. The level of expression of CitM was inversely related to the growth rates in the different media, which is typical for CCR.

**Roles of HPr, Crh, and CcpA in repression of $citM$ expression in CI medium.** *B. subtilis* mutants SA003 (genotype ptsH1), QB7097 (crh), and QB5407 (ccpA) are defective in HPr, Crh, and CcpA, respectively, components involved in CCR in *B. subtilis* (see the introduction) (Table 1). Mutant QB7102 is a double mutant defective in the two homologous proteins HPr and Crh. The $CitM$-lacZ fusion was integrated into the chromosome of each of these mutants to measure the promoter activity in the mutant background under inducing and repressing conditions.

$CitM$ promoter activity in wild-type cells grown in CI medium was approximately sixfold lower than that observed in a medium with only citrate (CC medium) (Fig. 3). The activity was repressed 3.5-fold when glucose was also present in the medium (Fig. 4). Surprisingly, the $ccpA$ mutant strain showed a 17-fold increase in $\beta$-galactosidase activity compared to the wild-type level when grown in the absence of glucose in CI medium. As expected, no significant glucose repression was observed in the mutant. The results show that both inositol and glucose have a repressive effect on $citM$ expression in the wild-type strain and that the repression by both is mediated via CcpA. In agreement, a significantly elevated $\beta$-galactosidase activity was observed in the ptsH1 crh double mutant, but the activity was somewhat lower than that of the $ccpA$ mutant. HPr, Crh, or both are involved in inositol repression in addition to CcpA. Glucose repression was almost completely alleviated in the double mutant. In contrast to the double mutant, the ptsH1 and crh single-mutant strains showed wild-type promoter activity levels when grown in CI medium. Apparently, both proteins are involved in the repression of $citM$ expression by inositol but they can replace one another. Compared to the wild type, repression by glucose was less strong in the ptsH1 single mutant but was stronger in the crh mutant, where repression was very potent (Fig. 4, inset). This suggests an important role for HPr in glucose repression in CI medium.

Initial uptake rates of Ni$_2^+$-citrate in wild-type and mutant cells correlated well with $\beta$-galactosidase activity under the same conditions (Table 3). The results showed similar induction and repression features. Transport activity was 1 order of magnitude higher in the $ccpA$ mutant and the ptsH1 crh double mutant than in the wild-type strain. The repression of transport activity by inositol was equally well relieved in the $ccpA$ mutant and the ptsH1 crh double mutant, but in both mutants there was still significant repression by glucose. The transport activities of the crh and ptsH1 single mutants paralleled the promoter activity. Repression by inositol was not relieved in the single mutants, while glucose repression was most potent in the crh mutant and less so in the ptsH1 mutant.

**Roles of HPr, Crh, and CcpA in repression of $citM$ expression in CSE and CIE media.** Upon growth of cells in the CSE and CIE media, the main features of promoter activity and Ni$_2^+$-citrate uptake activity of the mutant and wild-type strains were similar to those observed in CI medium (Tables 3 and 4). Importantly, in the succinate-glutamate medium (CSE medium) both transport and promoter activities were four- to fivefold higher in the $ccpA$ mutant and ptsH1 crh double mutant than in the wild-type strain, indicating that the medium components (succinate and/or glutamate) repressed $citM$ expression via the CCR system. Noteworthy is that in CSE medium the double mutant and the $ccpA$ mutant gave comparable

![Graph showing $\beta$-galactosidase activity in CI medium](image)

**Table 3. Initial uptake rates of Ni$_2^+$-citrate in wild-type and mutant *B. subtilis* strains**

<table>
<thead>
<tr>
<th>Medium</th>
<th>Initial uptake rate (nmol/min/mg of protein) for strain$^a$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CSE (wild type)</td>
</tr>
<tr>
<td>CI + citrate</td>
<td>0.18</td>
</tr>
<tr>
<td>CI + citrate + glucose</td>
<td>0.10</td>
</tr>
<tr>
<td>CSE + citrate</td>
<td>0.38</td>
</tr>
<tr>
<td>CSE + citrate + glucose</td>
<td>0.01</td>
</tr>
<tr>
<td>CIE + citrate</td>
<td>0.17</td>
</tr>
<tr>
<td>CIE + citrate + glucose</td>
<td>0.01</td>
</tr>
</tbody>
</table>

$^a$ The relevant genotype is in parentheses.

**Table 4. $CitM$ promoter activity measured in wild-type and mutant *B. subtilis* strains**

<table>
<thead>
<tr>
<th>Medium</th>
<th>$\beta$-Galactosidase activity (Miller units) for strain$^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CM002 (wild type)</td>
</tr>
<tr>
<td>CI + citrate</td>
<td>35</td>
</tr>
<tr>
<td>CI + citrate + glucose</td>
<td>10</td>
</tr>
<tr>
<td>CSE + citrate</td>
<td>143</td>
</tr>
<tr>
<td>CSE + citrate + glucose</td>
<td>11</td>
</tr>
<tr>
<td>CIE + citrate</td>
<td>30</td>
</tr>
<tr>
<td>CIE + citrate + glucose</td>
<td>2</td>
</tr>
</tbody>
</table>

$^b$ The relevant genotype is in parentheses.
β-galactosidase and transport activities, indicating that repression by succinate and/or glutamate is completely mediated via HPrl and/or Crh. In the two single mutants, both transport activity and promoter activity were stimulated by a factor of two relative to those of the wild-type strain, indicating that both HPrl and Crh play a role in succinate and/or glutamate repression but that they cannot completely take over each other’s roles as was observed in repression by inositol. Additional repression by glucose in the wild-type and mutant strains was much stronger in CSE medium than in CI medium, except for the ccpA mutant, where repression was completely relieved.

CitM expression is repressed in CSE medium. Unfortunately, B. subtilis did not grow on C medium containing succinate as the sole carbon and energy source and only poorly on glutamate, which did not allow discrimination between succinate and glutamate as the repressive substrate. As an alternative, all experiments were repeated using CI medium to which glutamate was added (CIE medium). The addition of glutamate had no effect on the repression by inositol (Tables 3 and 4). However, there was a marked difference in the repression by glucose in this medium. The repression was much stronger than in the absence of glutamate and comparable to the repression in CSE medium.

**Growth defects of the ptsH1, crh, and ccpA mutant strains.** A (partly) functional CCR system was necessary for normal growth on inositol. Both the ptsH1 crh double mutant and the ccpA mutant grew poorly in CI and CIE minimal media, with growth rates of 0.07 to 0.11 h⁻¹ compared to 0.32 (CI medium) and 0.69 h⁻¹ (CIE medium) for the wild-type strain. Especially long lag phases were observed, and the addition of glucose, citrate, or glutamate (44) had no significant effect on the growth rate.

**DISCUSSION**

In B. subtilis the uptake of citrate is mediated by at least two known homologous secondary transport proteins (3). One of them, termed CitM, is a proton motive force-driven transporter that mediates the transport of citrate complexed to Mg²⁺ and several other divalent metal ions in symport with two protons (3). CitM is believed to be the predominant citrate transporter under physiological conditions (2, 3). The conditions under which CitM is expressed were the topic of our research. The expression of citM was monitored at the transcriptional level by measuring the citM promoter activity and at the protein level by measuring Ni²⁺-citrate uptake activity in whole cells. Our results indicate that citM expression is under strict control of the medium composition. CitM is an inducible protein that is only expressed when citrate is present in the growth medium, which is sensed by the CitS-CitT two-component system, whose coding sequence is located upstream of citM (9; Yamamoto et al., Abstr. 10th Int. Conf. Bacillli). The expression was highest when citrate was the only carbon and energy source in the medium. The carbohydrates glucose, glycerol, and inositol are preferred over citrate, resulting in higher growth rates and a strongly repressed expression of citM. Remarkably, the expression of citM was also repressed during growth on the nonsugars succinate and glutamate, albeit to a lesser extent. In B. subtilis, repression by growth substrates other than glucose has been reported for inositol dehydrogenase, which was repressed by glycerol and mannitol (7), for the hut operon, that was found to be repressed by amino acids (43) and for the first three enzymes of the Krebs cycle were found to be repressed by glutamate and glutamine (8, 33).

Transcription of the structural gene is the first step in the biosynthetic pathway of a protein. The amount of a membrane protein such as CitM that ends up in the cytoplasmic membrane depends on the rate of transcription of the gene (measured by the promoter activity) and on other factors such as messenger stability, efficiency of insertion into the membrane, and protein stability. The activity of the ensemble of protein molecules in the membrane, which is the relevant parameter for the cell, may further depend on regulation of the activity of the individual protein molecules by global cellular factors such as pH and redox potential and by more-specific effectors. Most importantly, for secondary transporters, transport activity depends on the energy status of the cell. Transport activity catalyzed by CitM is driven by the electrochemical proton gradient that is maintained across the cytoplasmic membrane by the cellular energy metabolism (3). In this study, a reasonable correlation between citM promoter activity and Ni²⁺-citrate uptake activity in the different strains and under different growth conditions was observed; this is somewhat surprising considering the above discussion. Apparently, in the media tested, the uptake of citrate complexed to divalent metal ions in B. subtilis is mainly regulated at the level of transcription of the citM gene.

The promoter region of citM contains a cre sequence centered 24 bp upstream of the citM start codon. The 14-bp DNA sequence with dyad symmetry deviates very little from the consensus sequence (41, 46) (Fig. 1), which suggested that CitM is likely to be subject to carbon catabolite repression. It has been suggested that A- and T-rich regions intensify the interactions of catabolite control protein A (CcpA) with the cre site flanking the citM cre site (46). Recently, it was demonstrated that the citM cre site is active in vivo (32). The strong relief of citM repression in all media tested in a defective-CcpA mutant indicates that repression by not only glucose but also inositol and succinate and/or glutamate in the wild-type strain is mediated by the binding of CcpA to the cre site located in the citM promoter region.

In CCR, the binding of CcpA to the cre site is induced by complex formation of CcpA and the phosphorylated forms of HPrl and Crh. Repression of citM expression in CI medium was less relieved in the ptsH1 crh double mutant than in the ccpA mutant. Since in the double mutant the CcpA molecule is present, this may indicate an affinity of uncomplexed CcpA for the cre site or promotion of binding by factors other than HPrl and Crh (15, 23). Complete relief of repression in the double mutant when grown in CSE medium suggests that in CI medium other metabolic intermediates promote CcpA binding to the cre site.

The specific roles of HPrl and Crh in CCR are not clear. HPrl is one of the general proteins in the phosphoenoxyruvate-dependent PTS that transports sugars into the cell with concomitant phosphorylation. HPrl is a phosphocarrier intermediate that is phosphorylated at a histidine residue (His15) by phosphoenoxyruvate in a reaction catalyzed by enzyme I (EI). It donates its phosphoryl group to a sugar-specific enzyme or enzyme domain termed IIA. In CCR, HPrl is phosphorylated by ATP at a serine residue (Ser46), a reaction catalyzed by HPrl kinase. It has been suggested that the functions of HPrl in the PTS and CCR interact when transcription of a gene is repressed by a PTS sugar, for instance, glucose (35). The Crh protein is not operational in the PTS (29), simply because, at the position corresponding to the phosphorylation site (His15) in HPrl, a glutamine residue is found in Crh. The CCR phosphorylation site (Ser46) is present, and it has been shown that Crh is phosphorylated by HPrl kinase in vitro (13, 14). In a number of studies, it has been observed that HPrl and Crh can take over each other’s role in the CCR function (34, 46). Similarly, in this study, repression of expression of citM by
inositol in the ptsH1 and crh single mutants was similar to that observed in the wild-type strain, while repression was alleviated in the ptsH1 crh double mutant (Fig. 4). It should be noted that inositol is believed to be taken up by the cell by a secondary transporter (encoded by ioIF) (45) and, therefore, that there is no turnover of the PTS during growth on inositol. For a PTS sugar, it may be anticipated that the CCR system discriminates between HPr and Crh (35). In agreement, repression of citM by glucose is more effective in the crh mutant than in the HPr mutant, suggesting that repression by HPr is potentiated during turnover of the PTS. Discrimination between HPr and Crh in the repression of citM in medium containing the nonsugars succinate and glutamate (CSE medium) was also observed in this study. Repression was relieved twofold in both single mutants, suggesting that HPr could only partly take over repression exerted by Crh and vice versa. The mechanism by which succinate and/or glutamate metabolism affects the degree of phosphorylation of HPr and Crh is completely unknown.

The addition of glucose to CI medium resulted in an additional threefold repression of citM expression. The addition of glutamate to CI medium had no effect on citM expression. Remarkably, when both glucose and glutamate were added to CI medium, the additional repression was about 15-fold and of the same order of magnitude of the glucose repression in CSE medium, which also contains glutamate. This suggests that the presence of glutamate in the medium makes repression by glucose much stronger. A similar synergistic repression has been observed for the citB gene of B. subtilis, coding for the tricarboxylic acid (TCA) cycle enzyme aconitate. Expression of the citB gene in minimal medium containing glucose or glyceral was fully repressed when a source of 2-ketoglutarate, such as citB gene inactivation, glutamate synthase (10). Addition of TCA cycle intermediates, such as citrate and glutamate, could restore the mutant to utilize ammonium as a single source of nitrogen (44), but not on that containing arabinose (38). In this study, the effect of inositol was as observed with arabinose, i.e., growth in CI and CIE medium was severely slowed down even when citrate or glutamate or both were present. The same was true for the ptsH1 crh double mutant grown in minimal medium in the presence of inositol, whereas the ptsH1 and crh single mutants showed wild-type growth rates under these conditions. Similar growth defects have been described by Zalieckas et al. (47).

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ADDENDUM IN PROOF

An extensive characterization of the citM promoter region was recently published (H. Yamamoto, M. Murata, and J. Sekiguchi, Microbiol. 37:898–912, 2000).

REFERENCES


