Conversion of Methionine to Cysteine in *Bacillus subtilis* and Its Regulation

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*Bacillus subtilis* can use methionine as the sole sulfur source, indicating an efficient conversion of methionine to cysteine. To characterize this pathway, the enzymatic activities of CysK, YrhA and YrhB purified in *Escherichia coli* were tested. Both CysK and YrhA have an O-acetylserine-thiol-lyase activity, but YrhA was 75-fold less active than CysK. An atypical cystathionine β-synthase activity using O-acetylserylserine and homocysteine as substrates was observed for YrhA but not for CysK. The YrhB protein had both cystathionine lyase and homocysteine γ-lyase activities in vitro. Due to their activity, we propose that YrhA and YrhB should be renamed MccA and MccB for methionine-to-cysteine conversion. Mutants inactivated for *cysK* or *yrhB* grew similarly to the wild-type strain in the presence of methionine. In contrast, the growth of an ΔyrhA mutant or a luxS mutant, inactivated for the S-ribosyl-homocysteine step of the S-adenosylmethionine recycling pathway, was strongly reduced with methionine, whereas a ΔcysK or ΔcysE mutant did not grow at all under the same conditions. The *yrhB* and *yrhA* genes form an operon together with *yrrT*, *mtnN*, and *yrrC*. The expression of the *yrr* operon was repressed in the presence of sulfate or cysteine. Both purified CysK and CymR, the global repressor of cysteine metabolism, were required to observe the formation of a protein-DNA complex with the *yrr* promoter region in gel-shift experiments. The addition of O-acetyl-serine prevented the formation of this protein-DNA complex.

Methionine plays a central role in a variety of cellular functions. This amino acid is the universal initiator of protein synthesis, and its derivative, S-adenosylmethionine (AdoMet), is involved in several cellular processes, including methylation and polyamine biosynthesis. Methionine synthesis and degradation are, therefore, tightly regulated.

*Bacillus subtilis*, like several microorganisms (21, 28, 55), can use sulfate for the synthesis of organic sulfur metabolites, mostly cysteine, methionine, and AdoMet. As in enterobacteria (28), the sulfate assimilation pathway of *B. subtilis* involves uptake and activation of inorganic sulfate, followed by stepwise reduction to sulfide (Fig. 1) (9, 23, 34, 59). An O-acetylserine-thiol-lyase, the *cysK* gene product, catalyzes the reaction of sulfide and O-acetylserine (OAS) to give cysteine (59). Cysteine is then converted into homocysteine by the transsulfuration pathway, which requires the sequential action of a cystathionine β-synthase, MetI, and two cystathionine β-lyases, MetC and PatB (6, 7). Homocysteine is subsequently methylated to methionine by a sole B$_2$-independent methionine synthase encoded by the *metE* gene (5) (Fig. 1).

AdoMet is synthesized from methionine and ATP by an AdoMet synthase encoded by the *metK* gene (21, 66). Utilization of AdoMet as a methyl donor results in formation of S-adenosylhomocysteine (AdoHcy). AdoHcy is converted to homocysteine in two steps catalyzed by the *mtnN* and *luxS* gene products, respectively (44, 49) (Fig. 1). AdoMet is also the precursor of polyamines leading to the production of methylthioadenosine (MTA). MTA is degraded in adenine and methylthioribose (MTR) by MTA nucleosidase, MtnN. The major subsequent steps for MTR recycling in *B. subtilis* have been recently characterized (4, 50).

*B. subtilis* can also use methionine as sole sulfur source. Microorganisms and mammals catabolize L-methionine through three main pathways (51). Methionine can be directly converted to methanethiol, α-ketobutyrate, and ammonium by a methionine γ-lyase (EC 4.4.1.11), an enzyme that is found in many bacteria (2, 25, 31, 54). Methionine utilization can also occur via a two-step degradation pathway initiated by an ammonia-lyase (67, 68). This enzyme requires the presence of an acceptor, e.g., α-ketoglutarate, to give α-keto-γ-methylthio-butyric acid (KMBA), which is subsequently degraded to methanethiol (10). In *Klebsiella aerogenes* and *Pseudomonas putida*, methanethiol may be oxidized to sulfite via a methane-sulfonate intermediate (48, 61). The third pathway involves the conversion of methionine to homocysteine through AdoMet and then the reverse transsulfuration pathway, which converts homocysteine to cysteine via the intermediary formation of cystathionine (Fig. 1). The reverse transsulfuration pathway, present in mammals, *Saccharomyces cerevisiae*, and *Pseudomonas aeruginosa*, requires the sequential action of cystathionine β-synthase and cystathionine γ-lyase (55, 61).

A number of enzymes involved in the metabolism of cysteine, homocysteine, and methionine are evolutionarily related. Cystathionine γ-synthase, cystathionine β-lyase, cystathionine γ-lyase, O-acetylhomoserine thiol-lyase, and methionine γ-lyase constitute a protein family, whereas OAS thiol-lyase and cat-

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tathionine β-synthase form a second family (35). The YrhB and YrhA proteins of B. subtilis belong to the cystathionine γ-synthase and to the OAS thiol-lyase family of proteins, respectively. They are good candidates to participate in the conversion of methionine to cysteine via the reverse transsulfuration pathway. We have previously shown that the expression of yrhA and yrhB genes is increased in the presence of methionine compared to the presence of sulfate (5). In addition, the expression of these genes is controlled by CymR, the central repressor of cysteine metabolism, and by CysK, the OAS thiol-lyase (1, 18). A CymR-dependent binding to the yrhT promoter region has been observed using crude extracts of Escherichia coli DH5α overproducing CymR, and OAS prevents the binding of this repressor (18). In the present study, the possible involvement of YrhA and YrhB in the utilization of methionine as the sole sulfur source in B. subtilis was analyzed. The signaling pathway, which modulates the CymR-dependent repression of yrhA and yrhB, was also studied.

MATERIALS AND METHODS

Bacterial strains and culture conditions. The B. subtilis strains used in the present study are listed in Table 1. A wild-type E. coli FBS strain was used in the present study (11), as well as an NK3 mutant (ΔpfpE5 leu-6 thi hldR hsdM− cysK− cysM) which is a cysteine auxotroph (24). E. coli cells were grown in LB broth (45) in M63 medium (100 mM KH2PO4, 40 mM (NH4)2SO4, 1 mM MgSO4, 0.1 mM FeCl3, 0.5% glucose, and 15 μM vitamin B12) or in a sulfate-free M63 medium (100 mM KH2PO4, 40 mM NH4Cl, 1 mM MgCl2, 0.1 mM FeCl3, 0.5% glucose, and 15 μM vitamin B12). B. subtilis was grown in SP medium or in minimal medium (6 mM KH2PO4, 4.4 mM KH2PO4, 0.3 mM trisodium citrate, 5 mM MgCl2, 50 μM CaCl2, 5 μM MnCl2, 0.5% glucose, 50 mg of t-tryptophan liter−1, 22 mg of ferric ammonium citrate liter−1, 0.1% l-glutamine, 20 mM asparagine) supplemented with a sulfur source as stated: 0.2 or 1 mM K2SO4, 0.2 or 1 mM L-methionine, 0.1 or 0.25 mM L-cystine, 0.05 or 0.5 mM L-cysteine, 0.2 or 1 mM β-D-thiogalactopyranoside) was added, allowing the expression of genes downstream from yrhA under the control of the Ppuc promoter. When required, antibiotics were added at the following concentrations: ampicillin, 100 μg ml−1, chloramphenicol, 5 μg ml−1, spectinomycin, 100 μg ml−1; and kanamycin 5 μg ml−1. Solid media were prepared by the addition of 20 g of Noble agar (Difco) liter−1. The loss of amylase activity was detected compared to the presence of sulfate (5). In addition, the expression of these genes is controlled by CymR, the central repressor of cysteine metabolism, and by CysK, the OAS thiol-lyase (1, 18). A CymR-dependent binding to the yrhT promoter region has been observed using crude extracts of Escherichia coli DH5α overproducing CymR, and OAS prevents the binding of this repressor (18). In the present study, the possible involvement of YrhA and YrhB in the utilization of methionine as the sole sulfur source in B. subtilis was analyzed. The signaling pathway, which modulates the CymR-dependent repression of yrhA and yrhB, was also studied.

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Plasmid and strain construction. Plasmids from E. coli and chromosomal DNA from B. subtilis were prepared according to standard procedures. Restriction enzymes, DNA polymerase, and phage T4 DNA ligase were used as rec-

FIG. 1. Biosynthesis and recycling pathways of sulfur containing amino acids. The enzymes present in B. subtilis are indicated by the corresponding genes: cysP, sulfate permease; sat, ATP sulfurylase; cystE, APS kinase; cysH, APS-PAPS reductase; sulfA, sulfite reductase; cysF, serine O-acetyltransferase; cysK, OAS thiol-lyase; YrhB, MetC, PatB, and CysK have cysteine desulfhydrase activity in vitro; metI, cystathionine γ-synthase; metC and PatB, cystathionine β-lyases; metE, methionine synthase; metK, AdoMet synthetase; speE, AdoMet decarboxylase; speF, sulfamide synthase; mtnN, AdoHey/MTA nucleosidase; mtnK, methylthioribose kinase, mtnA and mtnWXBD genes products are involved in the MTR-to-KMBA recycling pathway; mtnE, aminotransferase; lacS, S-rabiosylhomocysteine hydrolase; yrhA (mccA), cystathionine β-synthase; yrhB (mccB), cystathionine γ-lyase and homocysteine γ-lyase. The presence of underlined yrhA indicates low OAS thiol-lyase activity in vitro for YrhA. APS, adenosine 5′-phosphosulfate; KMBA, α-keto-γ-methyl-thiobutyric acid; MTA, methylthioribose; MTR, methylthioribose kinase; PAPS, 3′-phosphoadenosine 5′-phosphosulfate; SRH, S-rabiosylhomocysteine.

TABLE 1. Strains used in this study*

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<th>Strain</th>
<th>Genotype</th>
<th>Source or reference</th>
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<td>trpC2</td>
<td>Laboratory stock</td>
</tr>
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<td>BSIP1144</td>
<td>trpC2 amyE::pΔA(-108, +126) yrhT-lacZ cat</td>
<td>BSIP1304 → BSIP1303</td>
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<tr>
<td>BSIP1165</td>
<td>trpC2 yrhAB::aphA3</td>
<td>pDIA5533 → 168</td>
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<tr>
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<td>BSIP1304</td>
<td>trpC2 ΔcysK::spa</td>
<td>6</td>
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<tr>
<td>BSIP1305</td>
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<td>BSIP1304 → BSIP1303</td>
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<td>pDIA5512 → QB944</td>
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* Arrows indicate construction by transformation. cat is the pC194 chloramphenicol acetyltransferase gene, aphA3 is a kanamycin resistance gene, and erm is the PMutin tetracycline resistance gene. BFS2063 was constructed during the Bacillus Functional Analysis Program using the pMutin4 system (27).

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encoding the LacI repressor. The resulting strains were grown at room temperature in LB medium to an OD_{600} of 1.5, and IPTG (1 mM) was added to induce the expression of yrhB, cysK, or cymR, followed by incubation for 3 h. In the case of yrhA overexpression, 0.1 mM IPTG was used for induction. Cells were centrifuged and resuspended in 50 mM sodium phosphate (pH 8) and 300 mM NaCl. E. coli crude extracts were loaded on a Ni-nitrilotriacetic acid agarose column. The YrhB$_{NcoI}$, YrhB$_{NspI}$, and CysK$_{NcoI}$ proteins were eluted in the presence of 300 mM imidazole. E. coli crude extracts containing the CymR-intein-CBD protein fusion were loaded on a chitin column. The CymR protein was eluted after the cleavage of intein in the presence of 50 mM dithiothreitol (DTT). The purified CymR was concentrated, and DTT was eliminated by using an Amicon Ultra column (UFCB801024). The purified YrhA$_{NcoI}$, YrhB$_{NspI}$, CysK$_{NcoI}$, and CymR was confirmed by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (data not shown).

Enzyme assays. β-Galactosidase specific activity was measured as described by Miller (36) with cell extracts obtained by lysosome treatment. Protein concentrations were determined by the method of Bradford. One unit of β-galactosi-dase is defined as the amount of enzyme that produces 1 nmol of O-nitrophenol (ONP) min$^{-1}$ at 26°C. The mean value of at least three independent experiments is presented. The standard deviations are <15%.

OAS thiol-lyase catalyzes the reaction of O-acetylcysteine and sulfide to give cysteine. The production of cysteine was tested with the nihydrid method described by Gaitonde (20) and adapted by Ravanel et al. (42). The reaction mixture contained 100 mM phosphate (pH 7.5), 20 μM pyridoxal phosphate (PLP), 10 mM OAS, 10 mM Na$_2$S, and various amounts of purified CysK$_{NcoI}$ or YrhA$_{NcoI}$ in a final volume of 1 ml. The reaction mixture was incubated for 2 to 10 min at 30°C. The reaction was stopped by the addition of 300 μl of the acridine acid reagent. The samples were heated for 10 min at 100°C cooled for 2 min on ice, and kept at room temperature. Then, 650 μl of 95% ethanol was added, and the absorbance at 560 nm of the sample was determined. The amount of cysteine produced is calculated from a calibration curve, which shows a linear relation between absorbance and concentration up to 2 mM cysteine.

Cystathionine β-synthase catalyzes the reaction of 1-homocysteine and serine or serine derivatives to give cystathionine. This compound displays a peak of absorption at 455 nm in acidic solution as described by Kashiwamata and Greenberg (26). The reaction mixture contained 100 mM phosphate (pH 7.5), 20 μM PLP, 10 mM homocysteine, 10 mM OAS, or 10 mM serine and various amounts of enzyme. The reaction mixture was incubated at 37°C. Samples were taken at regular intervals, and the reaction was stopped by the addition of the acid nihydrid solution. After 5 min at 100°C, samples were cooled at room temperature, and the absorbance at 455 nm was determined. The amount of cystathionine produced was calculated from a calibration curve, which shows a linear relation between absorbance and concentration up to 20 mM cystathionine.

The OAS thiol-lyase and cystathionine β-synthase activity was also estimated spectrophotometrically at 340 nm by following the amount of acid released from OAS, in a coupling system that involved acetate kinase, pyruvate kinase, lactate dehydrogenase, and phosphoenolpyruvate carboxykinase. The reaction mixture (0.5 ml final volume) contained the following reagents: 50 mM Tris-HCl (pH 7.5), 50 mM KC1, 5 mM MgC$_2$, 10 μM PLP, 1 mM ATP, 1 mM fructose-1,6-biphosphate, 0.5 mM phosphoenolpyruvate, 0.15 mM NADH, 1 mM OAS, 20 μl of acetate kinase, 3 U of each pyruvate kinase, lactate dehydrogenase. The samples were equilibrated for several minutes at 30°C, and then the reaction was started by adding appropriate amounts of enzyme and either 1 mM Na$_2$S or 2 mM homocysteine. The decrease in absorption at 340 nm was monitored for 5 min. The linear segment being used to calculate the specific activity.

Cystathionine γ-lyase catalyzes the conversion of cystathionine to ammonia, α-ketobutyrate, and cysteine. The production of free thiol groups was measured by spontaneous disulfide interchange with 5,5'-dithiobis-(2-nitrobenzoic acid) (DTNB) (57). The reaction mixture contained 100 mM Tris-HCl (pH 9.0), 0.2 mM DTNB, 20 μM PLP, various cystathionine concentrations (200 μM to 10 mM), and 10 μg of purified YrhB. The increase in absorbance at 412 nm was measured at intervals of 0.5 min for 20 min. A molar absorption coefficient for the aryl mercaptide of 13,400 M$^{-1}$ cm$^{-1}$ was used to calculate the enzyme activity. One enzyme unit represents the formation of 1 μmol of arylmercaptan min$^{-1}$ at 30°C. Methionine γ-lyase activity leading to the production of meth-anethiol was also measured with DTNB using methionine instead of cystathio nine as substrate.

Homocysteine γ-lyase was assayed as described by Thong and Coombs (56). The reaction mixture contained 50 mM Tris-HCl (pH 7.5), 20 μM PLP, 3 mM l-homocysteine, 0.33 mM lead acetate, and 10 μg of purified YrhB in a final volume of 1 ml. The release of H$_2$S at 37°C was detected by using lead acetate as a trapping agent, and P$_5$S production was monitored at 360 nm. The amount
Sulfur containing products were identified by using high-pressure liquid chromatography (HPLC) as follows. The enzymatic activity was assayed as indicated for each enzyme in the presence of 10 or 50 \( \mu \text{M} \) of the corresponding purified protein. The reaction was stopped after 15 or 30 min of incubation at 30°C by the addition of sulfosalicylic acid (final concentration, 3%). Samples of the supernatant obtained after centrifugation were analyzed by ion-exchange chromatography, followed by ninhydrin post-column derivatization (22). A PCX5200 post-column derivatizer (Pickering Laboratories, Mountain View, CA) was connected to a Waters 626 HPLC system (Waters, Milford, MA). Amino acids were separated on a lithium ion-exchange column (3 mm [inner diameter] by 150 mm; Pickering) and detected at 570 and 440 nm on a Waters 2487 dual absorbance detector (22).

Volatile molecules were extracted by solid-phase microextraction (SPME) and analyzed by using a system composed of a gas chromatograph (GC6890+; Agilent Technologies, Inc., Palo Alto, CA) connected to a mass spectrometer detector (MS5973N; Agilent Technologies). Briefly, volatile molecules were extracted by plunging a 75-μm Carbonex PDMS fiber (Supelco, Bellefonte, PA) into the headspace of the reacting vial at 37°C for 15 min and were therapeutically desorbed from the fiber at 240°C in the splitless injector. Molecules were separated by using a HP Innowax capillary column (60 m by 0.25 mm, 0.50-μm film thickness) (Agilent Technologies) flushed with high-purity helium at a constant flow of 1.4 ml min \(^{-1}\). The temperature was increased from 40 to 240°C with a linear gradient of 10°C min \(^{-1}\). The column was directly connected to the mass detector in scan mode (electronic impact energy = 70 eV, m/z range = 20 to 200). HS was identified by its retention time and comparison of its spectrum with those of NIST 98 library (National Institute of Standards and Technology) and quantified by external standard calibration (12).

The bioluminescence assay for the detection of autoinducer 2 (AI-2) was performed as follows. \( B. \ subtilis \) strains were grown at 37°C in minimal medium with 1 mM homocysteine until the late exponential phase. Cells were centrifuged at 5,000 rpm, and the supernatants were filtered through a 0.22-μm pore-size filter. This cell-free conditioned culture medium was kept at -20°C. \( V. \ harveyi \) indicator strain BB 170 (8) was grown overnight at 30°C in autoinducer bioassay medium and then diluted 1:10,000 in fresh autoinducer bioassay broth, with 10% cell-free conditioned culture medium. Aliquots (1 ml) were taken hourly during 4 h to measure the bioluminescence using a Lumat 9507 luminometer (Berthold).

**RESULTS**

Role of LuxS in AI-2 production and in the conversion of methionine to homocysteine. Methionine is converted to AdoMet by the AdoMet synthase MetK. The by-product of AdoMet-dependent methylation reactions is AdoHcy, which is degraded to homocysteine by the successive action of the MtnN and LuxS enzymes (Fig. 1). An mtnN mutant grows poorly in the presence of methionine as the sole sulfur source (49). Nevertheless, MtnN is a AdoHcy/MTA nucleosidase participating both in AdoMet recycling and in the methionine salvage pathway via MTR (Fig. 1). It is therefore not possible to conclude whether the production of homocysteine via AdoMet, AdoHcy, and S-ribofylhomocysteine is essential for the conversion of methionine to cysteine. A mutant with an inactivated luxS gene was thus constructed. The LuxS enzyme is involved in the degradation of S-ribofylhomocysteine to homocysteine with the concomitant production of AI-2, a signaling molecule of quorum sensing (46, 60). We first verified that the \( B. \ subtilis \) luxS::cat mutant (BSIP1758) cannot produce AI-2. For this purpose, the ability of the wild-type and BSIP1758 strains to synthesize active AI-2 was determined using a \( V. \ harveyi \) reporter strain BB170 (8). In this system, the addition of cell-free supernatant from the wild-type strain stimulated light emission in \( V. \ harveyi \), whereas bioluminescence was not increased by the addition of the supernatant prepared from the luxS mutant. To determine whether the AdoMet recycling pathway is necessary for the utilization of methionine, we compared the growth of the wild-type strain and the \( \Delta \)luxS::cat mutant in minimal medium with 200 \( \mu \text{M} \) methionine, 200 \( \mu \text{M} \) homocysteine, 200 \( \mu \text{M} \) sulfate, or 100 \( \mu \text{M} \) cysteine as the sole sulfur source. Both strains grew similarly with sulfate, homocysteine, or cysteine (Fig. 2 and data not shown). In contrast, the \( \Delta \)luxS::cat mutant grew poorly in the presence of methionine compared to the wild-type strain (Fig. 2). This strongly suggests that methionine utilization requires first its conversion to homocysteine via the AdoMet recycling pathway. To study the possible involvement of the CysK, YrhA, and YrhB polypeptides in subsequent homocysteine degradation, the enzymatic activity of these proteins was determined.

Comparison of the enzymatic activity of the CysK and YrhA proteins. Cystathionine β-synthases and OAS thiol-lyases be-
long to the same family of PLP-dependent enzymes (35). The *B. subtilis* cysK and yrhA genes encode proteins of this family. CysK and YrhA share similarities to the *E. coli* OAS thiol-lyases, CysK (55 and 39% identity, respectively) and CysM (43 and 35% identity, respectively). A rather low level of identity (35%) is observed between CysK and YrhA. Previous work indicates that CysK plays a major role in the sulfate assimilation pathway (59). The YrhA<sub>Hsio</sub> and CysK<sub>Hsio</sub> polypeptides were overproduced in *E. coli*, purified, and assayed in vitro. OAS thiol-lyase catalyzes the reaction of OAS and sulfide to give l-cysteine and acetate. We first measured the production of cysteine by using the method of Gaitonde. In the presence of 10 mM OAS and 10 mM Na<sub>2</sub>S, purified CysK produced 300 μmol of cysteine min<sup>−1</sup> mg of protein<sup>−1</sup>, whereas purified YrhA merely produced 4 μmol of cysteine min<sup>−1</sup> mg of protein<sup>−1</sup>. The release of acetate from OAS was also assayed by measuring the disappearance of NADH using coupled reactions catalyzed by acetate kinase, pyruvate kinase, and lactate dehydrogenase. YrhA was 100-fold less active than CysK. The OAS thiol-lyase activity was then measured in crude extracts of the wild-type and the ΔcysK strains grown with methionine. The activity was 1.5 μmol of cysteine min<sup>−1</sup> mg of protein<sup>−1</sup> for the wild-type strain and 0.07 μmol of cysteine min<sup>−1</sup> mg of protein<sup>−1</sup> for the cysK mutant in agreement with the data obtained with purified proteins. To study the involvement of YrhA and CysK in cysteine biosynthesis in vivo, we compared the growth of the wild-type strain, a ΔcysK mutant, or a ΔyrhA mutant (BFS2063) in the presence of 200 μM sulfate or 100 μM cysteine. The wild-type and ΔyrhA strains grew similarly, with a doubling time of 60 min with sulfate and 50 min with cysteine. In contrast, the doubling time of the ΔcysK mutant in the presence of sulfate increased up to 500 min. Surprisingly, the growth of a ΔcysK mutant also decreased in the presence of cysteine, with a doubling time of 110 min and a reduced growth yield (data not shown). These results indicate that YrhA plays a minor role in sulfate assimilation and suggest another function for this protein.

We further examined the cystathionine β-synthase activity of YrhA and CysK. Cystathionine β-synthase catalyzes the reaction of homocysteine with serine to give cystathionine. In the presence of 10 mM homocysteine and 10 mM serine, no cystathionine was formed with purified YrhA and CysK using either an HPLC or a ninhydrin colorimetric assay. We then tested the possibility that OAS instead of serine could be a substrate for these enzymes. A specific activity of 4 μmol of cystathionine produced min<sup>−1</sup> mg of protein<sup>−1</sup> was observed with YrhA using the colorimetric assay. Under the same conditions, no cystathionine β-synthase activity was detected with CysK. The sulfur compound resulting from the reaction of OAS and homocysteine was determined by using HPLC. Cystathionine was detected only with purified YrhA.

YrhB, an enzyme with cystathionine-lyase and homocysteine γ-lyase activities in vitro. YrhB shares 50% identity with the cystathionine β-lyase (MetC) from *B. subtilis*, 50% identity with the recently characterized cystathionine γ-lyase from *Mycobacterium tuberculosis* (63), 46% identity with the cystathionine γ-lyases from rat and *Saccharomyces cerevisiae* (15), and 40 to 41% identity with the methionine γ-lyases from *P. putida* and *Trichomonas vaginalis* (25, 31). We had previously shown that YrhB has cysteine desulfhydrase activity in vitro (6). To analyze in more details the YrhB function, the protein was purified. Using an assay based on the rate of free-thiol group production, cystathionine degradation was detected. The YrhB enzyme obeyed Michaelis-Menten kinetics with cystathionine as a substrate (data not shown). The apparent *Kₘ* value, determined from Lineweaver-Burk plots, was 3 mM, and the *Vₘₕₖₐ₅* was about 2 μmol of free thiol groups produced min<sup>−1</sup> mg of protein<sup>−1</sup>. However, cystathionine degradation by cystathionine β-lyase or cystathionine γ-lyase leads to the production of different free-thiol containing compounds, homocysteine or cysteine, respectively. We thus tried to determine the sulfur compound resulting from cystathionine cleavage by YrhB using HPLC. Surprisingly, no significant amount of homocysteine or cysteine was detected after incubation of cystathionine with YrhB, despite the fact that this compound disappeared. Since YrhB has cysteine desulfhydrase activity in vitro (6), we propose that this enzyme could further produce sulfide from cysteine. In the presence of YrhB and cystathionine, sulfide formation was indeed detected with SPME-gas chromatography-mass spectrometry (data not shown). At the same time, pyruvate was produced, which could in turn react with cysteine to give 2-methyl-2,4-thiazolidine-carboxylic acid, as previously observed (16, 64). Finally, addition of the cystathionine degradation product obtained in vitro with YrhB allowed a weak growth of a *cysK cysM* double mutant of *E. coli* (NK3), which is a cysteine auxotroph (data not shown). This mutant cannot grow in M63 medium with cystathionine or homocysteine. This result suggests that YrhB produces cysteine and has cystathionine γ-lyase activity (see Discussion).

We further examined whether YrhB can use homocysteine or methionine as a substrate. Homocysteine γ-lyase activity was monitored by detecting PbS formation after incubation of YrhB<sub>Hsio</sub> in the presence of 3 mM homocysteine and lead acetate (see Materials and Methods). A specific activity of 0.33 μmol of PbS produced min<sup>−1</sup> mg of protein<sup>−1</sup> was detected with purified YrhB<sub>Hsio</sub>. Methionine γ-lyases directly produce methanethiol and α-ketobutyrate from methionine. No methionine γ-lyase activity was found for YrhB using 2 or 10 mM methionine as a substrate. Moreover, this activity could not be detected in crude extracts of the *B. subtilis* wild-type strain grown with methionine. These data indicate that there is no methionine γ-lyase in *B. subtilis* or that we failed to induce its synthesis.

**Growth of *E. coli* in the presence of the *B. subtilis* yrhB gene.** *E. coli* is unable to grow with cystathionine, homocysteine, or methionine as the sole sulfur source (21). We tested the ability of the *B. subtilis* yrhB gene to allow growth of *E. coli* with these sulfur sources. The yrhB gene was first cloned in pHT315 under the control of the lac promoter. *E. coli* FB8 strain containing either pHT315 or pHT315<sub>yrb</sub>B was grown in a sulfur-free M63 medium in the presence of sulfate, methionine, cystathionine, or homocysteine as the sole sulfur source. These two strains grew similarly on sulfate (data not shown). The presence of the yrbB gene in multicopy significantly increased the growth of strain FB8 with 1 mM homocysteine or 1 mM cystathionine compared to the same strain containing pHT315 alone (Fig. 3). In contrast, the presence of the yrhB gene in *E. coli* did not modify the poor growth of this bacterium with methionine (data not shown).
Growth of *B. subtilis* mutants inactivated in the *yrhA*, *yrhB*, *cysK*, or *cysE* gene. To investigate the role of YrhA, YrhB, and CysK in methionine utilization, we compared the growth of a Δ*yrhA* mutant (BFS2063), an *yrhB* mutant (BSIP1165), a Δ*cysK* mutant (BSIP1304), and the wild-type strain in the presence of 200 μM methionine. In the BFS2063 mutant, the genes downstream from *yrhA* are expressed under the control of the IPTG-inducible Pspac promoter to avoid a major polar effect (27). The *yrhB* and Δ*cysK* mutants grew similarly to the wild-type strain in the presence of methionine (Fig. 4). In contrast, the growth of the Δ*yrhA* mutant significantly decreased compared to the wild-type strain even when 1 mM IPTG was added (Fig. 4). The low residual growth observed with strain BFS2063 (Δ*yrhA*) was abolished in the presence of a *cysK* gene disruption (Fig. 4). This indicates that methionine utilization mostly requires YrhA. A Δ*cysK* Δ*yrhA* double mutant cannot grow with homocysteine, but significant growth was still observed in the absence of cystathionine (data not shown). *B. subtilis* therefore synthesizes cysteine from cystathionine without the intermediary production of homocysteine.

Since OAS is a substrate for the OAS thiol-lyase and the cystathionine β-synthase activity of YrhA, we tested the phenotype of a *cysE*14 mutant (QB944) lacking the serine O-acetyltransferase activity (17). The *cysE* mutant did not grow in the presence of 200 μM sulfate, 200 μM methionine and 200 μM homocysteine, whereas it grew with 100 μM cystine (data not shown). This indicates that CysE is necessary for sulfate assimilation and also for methionine and homocysteine conversion to cysteine.

The YrhB protein seems to be important both for the reverse transsulfuration and the sulfide-dependent pathways of the methionine-to-cysteine conversion (Fig. 1). Surprisingly, the *yrhB* mutant grew similarly to the wild-type strain in the presence of homocysteine or methionine (Fig. 4 and data not shown), suggesting the existence of bypass(es) for YrhB activity. We tested the growth of a Δ*yrhAB* Δ*cysK* mutant (BSIP1305) in the presence of cystathionine or homocysteine. A significant growth was observed in the presence of cystathionine, whereas the mutant did not grow in the presence of homocysteine (Fig. 5). Thus, a strain inactivated for CysK, YrhA, and YrhB is still able to synthesize directly cysteine from cystathionine.

**FIG. 3.** Growth of an *E. coli* FB8 strain in the presence of pHT315 or pHT315*yrhB*. Growth curves for strain FB8(pHT315) (squares) or FB8(pHT315*yrhB*) (circles) in the presence of dl-homocysteine (open symbols) or dl-cystathionine (closed symbols).

**FIG. 4.** Growth of the *B. subtilis* wild-type strain and various mutants in the presence of methionine. Growth curves for strains 168 (●), BSIP1165 (yrhB:aphA3) (○), BSIP1304 (ΔcysK::spc) (■), BFS2063 (Δ*yrhA*) (□), BSIP1870 (Δ*yrhA* Δ*cysK::spc*) (▲), or QB944 (cysE14) (△) in the presence of 200 μM methionine. For strains BFS2063 and BSIP1870, 1 mM IPTG was added.

**FIG. 5.** Growth of the *B. subtilis* wild-type strain and Δ*yrhAB*Δ*cysK* mutant in the presence of cystathionine or homocysteine. Growth curves in cystathionine (open symbols) or homocysteine (closed symbols) for strains 168 (triangles) and BSIP1305 (Δ*yrhAB*:aphA3 Δ*cysK::spc*) (circles).
**yrrTmtnNyrhABC operon.** Using transcriptome experiments, the expression of the *yrhA*, *yrhB*, and *mtnN* genes has been shown to decrease in the presence of sulfate (5). These genes, which are adjacent in the chromosome, may belong to the same operon. A Northern blot experiment with total RNA isolated from *B. subtilis* 168 grown with methionine or sulfate and hybridization with an *yrhB*-specific probe revealed a major 3.9-kb transcript (Fig. 6). This transcript was detected in RNA extracted from methionine-grown cells (Fig. 6B, lane 2) and absent from sulfate-grown cells (Fig. 6B, lane 1). The size expected for a *mtnN yrhAB* transcript is 3 kb. The *yrrT* and *yrhC* genes, located upstream and downstream of these genes, are transcribed in the same direction. Using *yrrT*- or *yrhC*-specific probes, we also detected the same 3.9-kb transcript (data not shown). The *yrrT*, *mtnN*, and *yrhABC* genes therefore form an operon (*yrrTmtnNyrhABC*). The transcription start site of the *yrrTmntNyrhABC* operon was mapped by primer extension using RNA isolated from strain 168 grown with methionine or sulfate. A band was observed only in the reaction performed with RNA extracted from methionine-grown cells (Fig. 6C, lane 2). The transcription was initiated at a G residue located 59 bp upstream from the *yrrT* translational start site (Fig. 6A). The −10 region (TATTAT) is similar to the consensus sequence of σA-dependent promoters, whereas the deduced −35 region (TAGTAC) presents only low similarity with known *B. subtilis* promoters.

**Expression of the yrrT operon in response to sulfur availability.** The expression of a pΔA(−108,+126)yr*rrT*-lacZ transcriptional fusion (18) was tested in the presence of different sulfur sources. The level of β-galactosidase activity was high in the presence of methionine (660 nmol of ONP mg of protein⁻¹) and slightly reduced in the presence of homocysteine or cystathionine (360 nmol of ONP mg of protein⁻¹). In the presence of sulfate or cysteine, the expression of this fusion was reduced 10- and 60-fold compared to the level of expression in the presence of methionine. Interestingly, the addition of 0.5 mM cysteine or 1 mM sulfate to methionine also repressed the expression of the *yrrT*-lacZ fusion. In contrast, the addition of 50 μM cysteine did not modify *yrrT* expression. These results showed that the transcription of the *yrrT* operon was repressed by sulfate and cysteine at high concentrations in agreement with the role of MtnN, YrhA, and YrhB in the conversion of methionine to cysteine.

We have previously shown that CymR is involved in the sulfate-dependent repression of *yrrT* transcript and that OAS prevents the CymR-dependent binding to the *yrrT* promoter region (18). Albanesi et al. have shown that CysK, the OAS thiol-lyase, is a global regulator of genes participating in...
cysteine biosynthesis, including yrhA (1). To determine the role of CysE, the serine O-acetyltransferase and CysK in the regulation of the yrrT operon, the expression of the pΔAyrT-lacZ fusion was tested in a wild-type strain and in a cysE14 mutant, a ΔcysK mutant, or a cysE14 ΔcysK double mutant. Since the cysE14 and cysE14 ΔcysK mutants cannot grow with methionine, all of the strains were grown in minimal medium in the presence of methionine and cystine to an OD600 of 2 and then incubated for 2 h in the same medium with either 1 mM methionine or 250 μM cystine. The β-galactosidase activity was subsequently assayed (Table 2). In a wild-type background, the yrrT-lacZ fusion was expressed with methionine and repressed eightfold with cystine, whereas the level of expression was low in a cysE14 mutant in both conditions. Moreover, addition of OAS to the growth medium containing cystine resulted in an increase of pryrrT-lacZ expression (250 nmol of ONP mg of protein−1) compared to the level observed without OAS (40 nmol of ONP mg of protein−1). These results confirm the role of OAS in the signaling pathway controlling yrrT. In contrast, constitutive expression was observed with a ΔcymR, ΔcysK, or cysE14 ΔcysK mutant (Table 2). The yrrT derepression observed in a cysK mutant was probably not due to an accumulation of OAS in the absence of the major OAS thiolase but rather to a more direct role of CysK, as proposed for cysH regulation (1).

Involvement of CysK in the CymR-dependent binding to the yrrT promoter region. A CymR-dependent binding to the yrrT promoter region has been observed using crude extracts of E. coli DH5α overproducing CymR (18). To determine whether CysK or CysM from E. coli participated to this binding, pDIA5735 (pysyA-cymR) or pXT were used to transform an E. coli cysK cysM double mutant (NK3). Crude extracts of NK3 producing or not CymR were used in mobility shift DNA-binding assays with a labeled DNA fragment containing the yrrT promoter region (−108 to +126). No complex was formed in the presence of 7.5 μg of crude extracts of NK3 carrying either pDIA5735 or pXT (Fig. 7A, lanes 2 and 3), whereas a protein-DNA complex was observed with 7.5 μg of crude extracts of DH5α carrying pDIA5735 (data not shown). The addition of 0.1 or 0.3 μg of purified CysK from B. subtilis restored the formation of the protein-DNA complex in the presence of crude extracts of NK3 carrying pDIA5735 (Fig. 7A, lane 5). The same result was not obtained when YrhA replaced CysK (Fig. 7A, lane 6). These results strongly suggest that both CymR and CysK are required to observe a gel shift with the yrrT promoter region.

For further analysis, we purified a native CymR protein by using the CBD system. The ability of purified native CymR protein to interact with the yrrT promoter region in the presence or absence of CysK was then tested in gel shift experiments. No binding was observed in the presence of 1 μg of purified CymR or CysK alone (Fig. 7B, lanes 2 and 3). However, a complex was formed when increasing amounts of both CymR and CysK at a 1:1 molar ratio were added to the yrrT promoter fragment (Fig. 7B, lanes 4 to 9). Indeed, 200 ng of CymR and 400 ng of CysK were sufficient for the full retardation of the DNA fragment (Fig. 7B, lane 8).

OAS prevents the CymR-dependent binding to the yrrT promoter region. (A) Gel mobility shift experiments were performed by incubating crude extracts of a cysK cysM double mutant of E. coli (NK3) (7.5 μg of proteins) carrying either pXT (lanes 2 and 4) or pDIA5735 (pysyA-cymR) (lanes 3, 5, and 6) with 5′-radiolabeled DNA fragments containing the yrrT promoter region. A total of 0.3 μg of purified CysK protein (lanes 4 and 5) or 0.3 μg of purified YrhA protein (lane 6) was added to the reaction mixture. Lane 1, free probe; (B) Binding of the purified CysK and CymR proteins to the yrrT promoter region. Lane 1, free probe; lanes 2, 1 μg of purified CymR protein added; lane 3, 1 μg of purified CysK protein added; lanes 4 to 9, increasing amounts of CymR and CysK proteins added at a 1:1 molar ratio (50:100 ng, 75:150 ng, 100:200 ng, 150:300 ng, 200:400 ng, and 250:500 ng of CymR and CysK proteins, respectively). (C) Negative effect of OAS on the binding of the CymR-CysK complex to the yrrT promoter region. Lane 1, free probe; lanes 2 to 9, 250 ng of CymR and 500 ng of CysK. Lanes 3 to 8 show results with increasing amounts of OAS (0, 0.1, 0.2, 0.5, 1, 2, and 5 mM, respectively). Lane 9, 5 mM NAS.

<table>
<thead>
<tr>
<th>Strain</th>
<th>Genotype</th>
<th>β-Galactosidase activity (nmol of ONP min−1 mg of protein−1)</th>
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<tbody>
<tr>
<td>BSIP 1144</td>
<td>p(−108, +126) yrrT-lacZ</td>
<td>Methionine (1 mM) 55</td>
</tr>
<tr>
<td>BSIP 1534</td>
<td>p(−108, +126) yrrT-lacZ</td>
<td>Cystine (0.25 mM) 40</td>
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<tr>
<td>BSIP 1837</td>
<td>p(−108, +126) yrrT-lacZ</td>
<td>3</td>
</tr>
<tr>
<td>BSIP 1840</td>
<td>p(−108, +126) yrrT-lacZ</td>
<td>310</td>
</tr>
<tr>
<td>BSIP 1794</td>
<td>p(−108, +126) yrrT-lacZ</td>
<td>490</td>
</tr>
</tbody>
</table>

* Cells were grown in minimal medium containing 1 mM methionine and 0.25 mM cystine until they reached an OD600 of 2. They were then centrifuged, washed three times in minimal medium without any added sulfur source, and resuspended in minimal medium containing either 1 mM methionine or 0.25 mM cystine. Cells were harvested 2 h after resuspension.
promoter in mobility shift experiments with crude extracts containing CymR (18). Similarly, the addition of increasing concentrations of OAS (0.1 to 5 mM) to the binding assay resulted in the release of the probe from the CymR-CysK protein-DNA complex (Fig. 7C, lanes 3 to 8). The formation of this complex was strongly reduced in the presence of 2 mM OAS (Fig. 7C, lane 7). Moreover, the negative effect of OAS is specific, since its analogous compound, NAS, at 5 mM is unable to dissociate the DNA-protein complex (Fig. 7C, lane 9).

**DISCUSSION**

As observed for other bacteria such as *K. aerogenes, P. putida, P. aeruginosa*, and *M. tuberculosis* (48, 61, 63), *B. subtilis* is able to grow in the presence of methionine, homocysteine, and cystathionine as a sulfur source, indicating the existence of an efficient conversion of these compounds to cysteine. A luxS mutant inactivated for one step of the AdoMet recycling pathway (Fig. 1) grows poorly in the presence of methionine (Fig. 2). Methionine utilization in *B. subtilis* mostly proceeds via the formation of homocysteine using the AdoMet recycling pathway. In this way, methanethiol and methanesulfonate are not intermediates in the methionine-to-cysteine conversion as proposed in *P. putida* (61). In agreement with these results, neither methionine γ-hyrase activity nor methanethiol is detected in the culture of *B. subtilis* (47; the present study work; S. Auger, unpublished results). Moreover, a mutant inactivated for the ssuD gene encoding the monooxygenase required for aliphatic sulfoxidation (utilization) (Fig. 1) is still able to grow in the presence of methionine as the sole sulfur source (I. Martin-Verstraete, unpublished results). The LuxS enzyme is involved in the production of AI-2, a universal signaling factor for interspecies communication (46, 60). AI-2 is generated by the spontaneous cyclization of 4,5-dihydroxyl-2,3-pentanedi-one, which is produced during the two-step recycling pathway of AdoHcy into homocysteine involving MtnN and LuxS (Fig. 1). As recently shown (32), a *B. subtilis* luxS mutant is unable to produce AI-2. In *B. subtilis*, the luxS gene is monocistronic. The mtnN gene, which encodes the AdoHcy nucleosidase (Fig. 1), forms an operon together with *yrbA, yrbB*, a gene encoding a putative AdoMet-dependent methyltransferase (*yrbT*), and *yrbC* (Fig. 6). Interestingly, either *luxS* or *mtnN* form an operon with *yrbA*, *yrbB*, and *yrbT*-like genes in *Bacillus cereus, Bacillus anthracis, Bacillus stearothermophilus, Oceaconbacillus iheyensis, Clostridium perfringens*, and *Clostridium botulinum* (40, 43). The YrhA and YrbB proteins, which belong to the OAS thiol-lyase and cystathionine γ-synthase family of proteins, respectively, are good candidates for the conversion of homocysteine to cysteine. Both YrhA and CysK proteins have an OAS thiol-lyase activity in vitro in agreement with previous results, indicating that a double Δ*cysK ΔyrbA* mutant is unable to grow with sulfate (59). However, several findings indicate that CysK is the major OAS thiol-lyase required for sulfate assimilation in *B. subtilis*. First, the growth of a Δ*cysK* mutant is strongly reduced in the presence of sulfate, whereas a Δ*yrbA* mutant grows similarly to the wild-type strain. Moreover, the OAS thiol-lyase activity of CysK is 75-fold higher than that of YrhA. In contrast, cystathionine β-synthase activity was only observed in vitro with YrhA. Cystathionine β-synthases are usually 150 amino acids longer in their C-terminal region than OAS thiol-lyases (14, 15, 41). YrhA is more closely related to OAS thiol-lyases, as indicated by its size and its similarities. Nevertheless, YrhA has both OAS thiol-lyase and cystathionine β-synthase activities in vitro, like the enzymes of *Aeropyrum pernix* and *Trypanosoma cruzi* (37, 39). Serine is the substrate of cystathionine β-synthases from mammals and microorganisms, whereas YrhA uses OAS instead of serine in the conditions tested. As proposed by Mino and Ishikawa (37), the enzymes with cystathionine β-synthase and OAS thiol-lyase activities could have retained some properties of an ancestral protein that would have diverged later into individual enzymes with narrower or different substrate specificity.

In *B. subtilis*, a Δ*cysK* mutant grows normally with methionine as the sole source of sulfur. Despite the fact that *yrbA* is constitutively expressed at a high level in the *cysK* mutant (Table 2) (1), CysK still represents 95% of the OAS thiol-lyase activity. In contrast, the Δ*yrbA* mutant grows poorly in the presence of methionine, as observed for an *yrbA*-like mutant of *Lactococcus lactis* and a *cbs* (cystathionine β-synthase probable gene) mutant of *Streptomyces venezuelae* (14, 52). The drastic growth defect of the *B. subtilis* Δ*yrbA* mutant in the presence of methionine is most probably due to the cystathionine β-synthase activity of YrhA rather than to its low OAS thiol-lyase activity in the presence of the major OAS thiol-lyase, CysK. YrhA seems therefore to correspond to the first step of the reverse transsulfuration pathway (Fig. 1). The growth of a Δ*cysK ΔyrbA* double mutant with cystathionine but not with homocysteine strongly suggests that a cystathionine γ-lyase, the second step of the reverse transsulfuration pathway, is also present in *B. subtilis*. YrhB has cystathionine lyase, homocysteine γ-lyase, and cysteine desulfhydrase activities in vitro (6; this study). Surprisingly, we fail to detect significant production of cysteine from cystathionine in the presence of YrhB using HPLC. This could be due to the further degradation of cysteine by the cysteine desulfhydrase activity of YrhB. However, several results strongly suggest that YrhB is a cystathionine γ-lyase rather than a cystathionine β-lyase, an enzyme involved in methionine biosynthesis (Fig. 1). (i) The expression of the *yrbB* gene is high with methionine and reduced with sulfate or cysteine. (ii) Two cystathionine β-lyases, MetC and PatB, have already been characterized in *B. subtilis*. A *patB metC* double mutant did not grow with cystathionine, indicating that there is no third cystathionine β-lyase (6). The normal growth of a *B. subtilis* *yrbB* mutant raises the question of the physiological role of the corresponding enzyme in vivo. The significant growth of *E. coli* with homocysteine in the presence of the *yrbB* gene indicates that YrhB may function as a homocysteine γ-lyase in vivo in *E. coli*, leading to the production of sulfide, which can in turn form cysteine with CysK (28). Physiological evidence for the participation of YrhB in the reverse transsulfuration pathway is lacking. Indeed, a strain inactivated for CysK, YrhA, and YrhB is still able to synthesize cysteine from cystathionine without the intermediary production of homocysteine. An alternative cystathionine γ-lyase different from *yrbB* is probably present in *B. subtilis*. The identification of this second cystathionine γ-lyase and its role in methionine degrada-
named MccA and MccB, respectively. In agreement with its role in this conversion, the expression of the yrrT operon is repressed in the presence of cysteine and sulfate. We unraveled some features of the control of yrrT expression in response to sulfur availability. Recent data indicate that CymR and CysK, the major OAS thiol-lyase of B. subtilis, is a global regulator of cysteine metabolism and that OAS plays a key role in the CysK- and CymR-dependent regulations (1, 18). We have shown that crude extracts from E. coli containing CymR interact with the yrrT promoter region only in the presence of CysM and/or CysK from E. coli (Fig. 7A). Purified B. subtilis CymR or CysK alone cannot bind to DNA, whereas the addition of proteins in a molar ratio of one to one allows the formation of a protein-DNA complex (Fig. 7B). Thus, both CysK and CymR are required for DNA binding. It is worth noting that CysK or CysM from E. coli, which shares 55 and 43% identity with CysK from B. subtilis, seem to assist CymR-dependent binding. This suggests that conserved amino acids in OAS thiol-lyases from these two organisms are involved in CymR and CysK interaction. CysK does not contain any DNA-binding motif, whereas CymR, an Rr2-type repressor, has a typical helix-turn-helix motif. We may then propose that the probable formation of a CysK-CymR complex induces conformational changes to help the CymR-dependent binding to its targets. Interestingly, the glutamine synthase of B. subtilis also directly interacts with the global regulator of nitrogen assimilation TnrA (65). OAS is required for the transcription of the cysH, ssu, and yrrT operons and ytlI, encoding the activator of the ytmI operon (13, 38, 58; the present study). As observed for the cysH operon (1), yrrT is actively transcribed regardless of the absence of OAS in a cysK background (Table 2). OAS modulates the interaction of the CymR-CysK probable complex with DNA. CysK may be a good candidate for sensing the cysteine pool of the cell by the regulatory complex. Indeed, OAS, which is a substrate of CysK, binds to the active site of this enzyme. Whether OAS binding dissociates the CymR-CysK complex or decreases the affinity of this complex for DNA deserves further investigation. For the cysH operon, Albanesi et al. (1) have proposed that CysK may interact with an uncharacterized repressor, CysR, to promote its binding to the cysH promoter. Further experiments are needed to determine the existence of a unique repressor or two distinct regulators, CysR and CymR. Cysteine probably limits the size of the intracellular pool of OAS, leading to CymR-dependent repression. As observed for E. coli, CysE activity may be feedback inhibited by cysteine (28). In B. subtilis, cysE expression is also regulated by transcription antitermination at a cysteine-specific T-box (19). Thus, the level of OAS in microorganisms may be correlated with the level of uncharged cysteinyl-tRNA, which signals the cysteine status in the cell. A cysE mutant inactivated for the serine O-acetyltransferase is unable to grow with methionine or homocysteine. OAS plays a key role in the methionine-to-cysteine conversion both as a substrate for CysK and MccA and as a signaling molecule for the sulfur-dependent control of the yrtT operon. In E. coli and L. lactis, OAS or its derivative, NAS, also plays a crucial role in the signaling pathway controlling the CysB and the CmbR/PhuR regulons (30, 52).

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