

Development and Characterization of a Xylose-Dependent System for Expression of Cloned Genes in *Bacillus subtilis*: Conditional Complementation of a Teichoic Acid Mutant

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We have developed a xylose-dependent expression system for tight and modulated expression of cloned genes in *Bacillus subtilis*. The expression system is contained on plasmid pSWEET for integration at the *amyE* locus of *B. subtilis* and incorporates components of the well-characterized, divergently transcribed xylose utilization operon. The system contains the xylose repressor encoded by *xylR*, the promoter and 5' portion of *xylA* containing an optimized catabolite-responsive element, and intergenic *xyl* operator sequences. We have rigorously compared this expression system to the isopropyl- β -D-thiogalactopyranoside-induced *spac* system using a thermostable β -galactosidase reporter (BgaB) and found the *xyl* promoter-operator to have a greater capacity for modulated expression, a higher induction/repression ratio (279-fold for the *xyl* system versus 24-fold with the *spac* promoter), and lower levels of expression in the absence of an inducer. We have used this system to probe an essential function in wall teichoic acid biosynthesis in *B. subtilis*. Expression of the teichoic acid biosynthesis gene *tagD*, encoding glycerol-3-phosphate cytidyltransferase, from the xylose-based expression system integrated at *amyE* exhibited xylose-dependent complementation of the temperature-sensitive mutant *tag-12* when grown at the nonpermissive temperature. Plasmid pSWEET thus provides a robust new expression system for conditional complementation in *B. subtilis*.

Manipulated gene expression in bacteria is of fundamental importance to understanding the effects of expression or depletion of gene products on bacterial physiology. Such investigations ideally require the controlled expression of a cloned gene from a tightly regulated, inducible promoter. This is particularly true in studies of indispensable genes in bacteria where targeted deletions of these genes require expression from complementing copies of the genes. The dispensability of such genes and the phenotypes associated with their loss are most rigorously examined through conditional expression of the complementing genes. Especially challenging in such studies are the need for extremely tight regulation of expression and the tendency for the majority of regulated expression systems to “leak” in the absence of an inducer.

The state of the art in controlled expression in *Bacillus subtilis* is the *spac* expression system, which is based on the application of the *lac* repressor-operator control system from *Escherichia coli*. In *B. subtilis*, the system uses a constitutive penicillinase promoter to express the *lac* repressor. Isopropyl- β -D-thiogalactopyranoside (IPTG)-dependent expression of the target gene is from a hybrid promoter-operator consisting of the SPO1 bacteriophage promoter and *lac* operator sequence (35). Despite its broad use for conditional expression in *B. subtilis*, the *spac* system has been widely recognized for its capacity to allow significant expression in the absence of an

inducer. Indeed, only relatively recently has there been a systematic study of expression from the *spac* promoter, indicating demonstrable uninduced expression of *lacZ* from the *spac* promoter of the pMUTIN vector system (33).

The xylose operon has emerged as a well-characterized *B. subtilis* regulatory system with the potential for particularly tight transcriptional regulation (5, 6, 12, 13, 16–19). Xylose utilization in *B. subtilis* requires the production of xylose isomerase (XylA) and xylulose kinase (XylB) and is regulated at the level of transcription by a xylose-responsive repressor protein encoded by *xylR* and by catabolite repression. Genes *xylR* and *xylAB* are divergently transcribed from a common intergenic region containing *xyl* operator sequences which are bound by XylR in the absence of an inducer. Transcription of the *xyl* operon is also catabolite repressed through the *cis*-acting catabolite-responsive element (CRE) located in the coding sequence of *xylA*.

Cell wall teichoic acids are a diverse group of phosphate-rich polymers which are covalently linked to peptidoglycan and constitute a substantial portion of the cell wall of gram-positive bacteria. Teichoic acids have been implicated as virulence factors in a variety of gram-positive bacterial infections (14, 22, 30), and growing evidence indicates that teichoic acid biosynthesis is indispensable for the growth of *B. subtilis* (2, 21, 23). Conditional lethal mutations in the poly(glycerol phosphate) teichoic acid gene cluster (*tag*) of *B. subtilis* 168 have been mapped to a number of genes, including *tagD* (27), encoding glycerol-3-phosphate cytidyltransferase (25).

In this work, we have taken advantage of a notable depth of knowledge of the xylose utilization operon to develop a xylose-dependent promoter-operator system for tight and modulated expression of cloned genes in *B. subtilis*. As a point of refer-

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TABLE 1. List of strains and plasmids

Strain or plasmid	Description	Reference or source
<i>B. subtilis</i>		
EB161	<i>trpC2 pheA1 amyE::ery</i>	<i>Bacillus</i> Genetic Stock Center 1A717
EB106	<i>trpC2 pheA1 amyE::xylR P_{xylA} cat86</i>	This work
EB107	<i>trpC2 pheA1 amyE::xylR P_{xylA} bgaB cat86</i>	This work
EB105	<i>trpC2 pheA1 amyE::P_{spac} lacI cat86</i>	This work
EB104	<i>trpC2 pheA1 amyE::P_{spac} bgaB cat86</i>	This work
EB103	<i>trpC2 pheA1 amyE::P_{spac} bgaB lacI cat86</i>	This work
EB6	<i>hisA1 argC4 metC3</i>	L5087 (2)
EB4 ^a	<i>hisA1 argC4 metC3 tag-12 ϕ29^f</i>	L6602 (2)
EB123	<i>hisA1 argC4 metC3 tag-12 ϕ29^f amyE::xylR P_{xylA} tagD cat86</i>	This work
EB127	<i>hisA1 argC4 metC3 tag-12 ϕ29^f amyE::P_{spac} tagD lacI cat86</i>	This work
<i>E. coli</i>		
Novablue ^b	<i>endA1 hsdR17 (r_{K12}⁻ m_{K12}⁺) supE44 thi-1 recA1 gyrA96 relA1 lac[F' proA⁺B⁺ lacI^aZΔM15::Tn10(TcR)]</i>	Novagen
Plasmids		
pDG364	<i>B. subtilis</i> vector for integration at <i>amyE</i>	<i>Bacillus</i> Genetic Stock Center (3)
pSWEET	pDG364 with <i>xyl</i> expression system	This work
pSWEET- <i>bgaB</i>	pSWEET with <i>bgaB</i> from <i>B. stearothermophilus</i> and polylinker	This work
pSWEET- <i>tagD</i>	pSWEET with wild-type <i>tagD</i> from <i>B. subtilis</i> 168	This work
pDR67	Plasmid containing <i>spac</i> expression system for integration at <i>amyE</i>	15
pSPAC- <i>bgaB</i> Δ <i>lacI</i>	pDR67 with <i>bgaB</i> from <i>B. stearothermophilus</i> without <i>lacI</i> gene	This work
pSPAC- <i>bgaB</i>	pDR67 with <i>bgaB</i> from <i>B. stearothermophilus</i>	This work
pSPAC- <i>tagD</i>	pDR67 with <i>tagD</i> from <i>B. subtilis</i> 168	This work

^a Temperature-sensitive mutant in teichoic acid biosynthesis ascribed to *tagD*.

^b Cloning strain.

ence, we have rigorously compared this expression system to the IPTG-induced *spac* system for efficiency of regulation and modulation of expression. Finally, we have put expression of *tagD* under control of the xylose-based expression system described here and have for the first time shown *trans* complementation of a teichoic acid mutant.

MATERIALS AND METHODS

General methods. Strains, plasmids, and primers used are listed in Tables 1 and 2. *B. subtilis* strains were grown in rich (Luria-Bertani [LB]) or minimal [15 mM (NH₄)₂SO₄, 80 mM K₂HPO₄, 44 mM KH₂PO₄, 3.4 mM sodium citrate, 1 mM MgSO₄ (pH 7.4)] medium plus arabinose (0.2%). (Arabinose was previously found not to exert a catabolite-repressive effect on the xylose operon [18].) The following concentrations of antibiotics were used for selection: 50 μg of ampicillin per ml, 10 μg of chloramphenicol (CHL) per ml, and 1 μg of erythromycin per ml. Unless otherwise stated, glucose was added to 0.2%, xylose was added to

2%, IPTG was added to 1 mM, and 5-bromo-4-chloro-3-indolyl-β-D-galactopyranoside (X-Gal) was added to 80 μg/ml. Cloning was performed in *E. coli* strain Novablue (Novagen) according to established methods (29). Transformation of these cells was performed according to the manufacturer's instructions. Restriction enzymes were purchased from New England Biolabs (Beverly, Mass.) and used according to the manufacturer's instructions. Transformation of *B. subtilis* organisms was performed according to procedures previously described (4); derivatives of pDG364 were targeted to *amyE* via double recombination as linear DNA, using approximately 1 μg of *Pst*I-digested plasmid. All other chemicals were purchased from Sigma (Mississauga, Ontario, Canada).

Plasmid construction. Plasmid pSWEET-*tagD* was constructed using the following strategy. The *xylR* gene, the *xylA* promoter, and the first 58 nucleotides of the *xylA* gene (including the CRE site) were amplified as a single product from *B. subtilis* W23 chromosomal DNA, using the primers AB01 and AB02. Primer AB02 incorporated two mutations in the CRE site, G→T and A→T at positions 3 and 10, respectively (18, 34). Gene *tagD*, from -24 to stop (including its native ribosome binding site), was amplified from *B. subtilis* 168 using the primers AB03

TABLE 2. List of primers

Oligo-nucleotide	Sequence ^{a,b}	Restriction site(s)
AB01	5'-GGCCGGATCCGAATTACATTTTAACGATATC-3'	<i>Bam</i> HI
AB02 ^c	5'-GCGCTTAATTAACCACCTTTGGTTAACGCTTACAAAATAGTT-3'	<i>Pac</i> I
AB03	5'-GCGCTTAATTAACATTAGGAAGGACGCTTTCTT-3'	<i>Pac</i> I
AB04	5'-CCGGATCCGCTTATAAACCAGCAATTTCCCTC-3'	<i>Bam</i> HI
AB05	5'-GGAAGGATCCGCGCTTAATTAACCACCTTTGGTTAACGCTT-3'	<i>Bam</i> HI/ <i>Pac</i> I
AB06	5'-GGAAAGATCTTTACATTTTAACGATATCTAGAAAAAT-3'	<i>Bgl</i> II
AB07 ^d	5'-CCTGCCCGGG/TTAATTAACAT <u>AGGAAGGAGCGTTTCTTTAAATGAATGTGTTATCCTCAATTTGTTACGG</u> -3'	<i>Sma</i> I/ <i>Pac</i> I
AB08 ^e	5'-GGAAGGATCC/AGCTT/GCGGCCG/CCGGGCTAGCTTAAACCTTCCCGGCTTCATCATGCTCTCT-3'	<i>Bam</i> HI/ <i>Hind</i> III/ <i>Not</i> I/ <i>Srf</i> I/ <i>Nhe</i> I
AB09	5'-GGCGGATCC/AGATCTCTAAACCTTCCCGG-3'	<i>Bam</i> HI/ <i>Bgl</i> II

^a Restriction sites are indicated in italics.

^b Multiple restriction sites are separated by a forward slash (/).

^c Underlined text indicates the optimized CRE site, and bold text indicates mismatches from the *xylA* CRE site.

^d Underlined text indicates the ribosome binding site, and bold text indicates a 9-bp spacer between the ribosome binding site and the initiation codon.

^e Underlined text indicates bases shared by two restriction sites.

and AB04. The above PCR products were digested with *PacI* and ligated together. The ligation product was reamplified with primers AB01 and AB04, digested with *BamHI*, and ligated into *BamHI*-digested pDG364.

To create pSWEET-*bgaB* the *xylR-xylA* fragment was reamplified from pSWEET-*tagD* with primers AB05 and AB06. The amplified fragment was digested with *BamHI* and *BglII* and ligated into *BamHI*-digested pDG364 to create plasmid pSWEET. Plasmid pK14 was used as a template for amplification of *bgaB* (31) using primers AB07 and AB08. Primer AB07 incorporated nucleotides -24 to -1 of *B. subtilis tagD* so that the ribosome binding sites and their contexts were identical in pSWEET-*tagD* and pSWEET-*bgaB*. The *bgaB* PCR product was digested with *PacI* and *BamHI* and ligated into pSWEET digested with the same enzymes.

To facilitate a rigorous comparison of the xylose-based expression system in pSWEET with that of the commonly used *spac* system, we constructed clones of *bgaB* and *tagD* under the control of the *spac* promoter in pDR67 (15) with ribosome binding sites and contexts identical to those used with pSWEET (i.e., nucleotides -24 to -1 of *tagD*, as described above). Those plasmids were named pSPAC-*bgaB* and pSPAC-*tagD*, respectively. In addition, we constructed a *lacI* deletion of pSPAC-*bgaB*, designated pSPAC-*bgaB* Δ *lacI*, in order to test the importance of *lacI* to the regulation of the *spac* promoter. Gene *bgaB* was amplified from pK14 using primers AB07 and AB09. To create pSPAC-*bgaB*, the amplified product was digested with *SmaI* and *BglII* and ligated to *SmaI*- and *BglII*-digested pDR67. For pSPAC-*bgaB* Δ *lacI*, the amplified product was digested with *SmaI* and *BamHI* and ligated to *SmaI*- and *BamHI*-digested pDR67. To place *tagD* under the control of the *spac* promoter, *tagD* was excised from pSWEET-*tagD* with *PacI* and *BamHI* and ligated into *PacI*- and *BamHI*-digested pSPAC-*bgaB*.

β -Galactosidase activity assay. An assay of thermostable β -galactosidase has been described previously (31, 32). In brief, cells were grown to an optical density at 600 nm (OD_{600}) of 0.5, pelleted, and resuspended in an equivalent volume of buffer B (25 mM potassium phosphate, 50 mM KCl, and 1 mM $MgSO_4$ [pH 6.4]). Sample aliquots ranging from 0.1 to 0.5 ml were diluted to 0.8 ml in buffer B and lysed for 30 min at 28°C with the addition of 16 μ l of lysozyme (10 mg/ml). Lysis was followed by the addition of 40 μ l of 10% Triton X-100. Subsequently, a preincubation at 60°C for 15 min was used to inactivate endogenous β -galactosidase. The assay was initiated with the addition of 0.2 ml of *o*-nitrophenyl- β -D-galactopyranoside (ONPG) at a concentration of 4 mg/ml and quenched by the addition of 0.5 ml of 1 M Na_2CO_3 . Absorbance was recorded at 420 nm in a Spectramax Plus spectrophotometer (Molecular Devices).

RESULTS

We chose to construct a xylose-based expression system by incorporating components of the xylose operon from strain W23 of *B. subtilis*. The xylose utilization machinery of this *B. subtilis* strain has been the subject of extensive characterization (5, 6, 12, 13, 16–19). The *xyl* expression system developed in this work includes the xylose repressor encoded by *xylR*, intergenic *xyl* operator sequences, the *xylA* promoter, and the 5' portion of *xylA* containing an optimized CRE (Fig. 1). The system is contained on plasmid pSWEET, a derivative of pDG364 (3), for ready integration of the expression system into the *B. subtilis* chromosome at *amyE*.

Transcriptional regulatory sequences in the *xyl* operon include tandem overlapping operator sequences downstream of the *xylA* transcriptional start site (5) and a CRE located 36 nucleotides into the coding sequence of the xylose isomerase gene (*xylA*) (18). To ensure that the *xyl* system was maximally subjected to catabolite repression, we included the first 58 nucleotides of the gene *xylA* and introduced two mismatches in the CRE site to perfectly match the consensus sequence established previously (18, 34). An in-frame stop codon was placed in the *xylA* gene following the 58th nucleotide, effectively truncating the XylA protein. Downstream of that translational stop, we constructed pSWEET-*bgaB* so that *PacI* (eight-base recognition sequence TTAATTAA) and one of several polylinker enzymes could be used to replace *bgaB* and

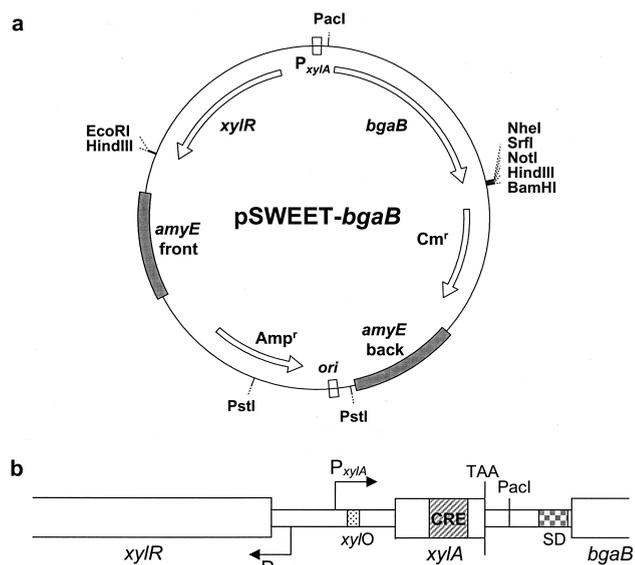


FIG. 1. Map of plasmid pSWEET-*bgaB*. (a) Significant features of the xylose-based expression system. Plasmid pSWEET-*bgaB* is a derivative of pDG364 (3), which allows integration into the *B. subtilis* chromosome at *amyE* via double recombination and selection with CHL (10 μ g/ml). The plasmid also has an *E. coli* origin of replication, denoted *ori*, and an ampicillin resistance cassette (50 μ g/ml) for routine cloning steps. On the outside of the plasmid map, restriction sites of interest are highlighted, including two *PstI* sites for convenient plasmid linearization, a *PacI* site (eight-base recognition sequence TTAATTAA) upstream of *bgaB*, and a polylinker downstream of *bgaB* (*HindIII* is not unique). (b) Close-up of key elements of the xylose expression system (not to scale). Shown are *xylR*, encoding the xylose repressor; the *xyl* intergenic region, including promoters for *xylR* (P_{xylR}), *xylA* (P_{xylA}), and *xyl* operator sequences (*xyLO*); translationally truncated *xylA* (first 58 nucleotides followed by an in-frame TAA), including an optimized CRE in *xylA* (see Materials and Methods and reference 18); *PacI* 5' cloning site; ribosome binding site (SD) native to *B. subtilis tagD*; and gene *bgaB*, encoding a thermostable β -galactosidase from *B. stearothermophilus*.

its associated ribosome binding site with any sequences of interest.

Schroegel and Allmansberger (31) have optimized the use of heat-stable β -galactosidase, *bgaB* from *B. stearothermophilus*, as a reporter gene in *B. subtilis*, which permits inactivation of endogenous background β -galactosidase activity. We employed this reporter system in order to describe the lower limits of transcriptional control afforded by the *xyl* system relative to the *spac* system. For an unbiased comparison of the *xyl* and *spac* expression systems, we constructed pSPAC-*bgaB* and pSWEET-*bgaB* so that their ribosome binding sites and their contexts were identical (see Materials and Methods).

Figure 2a and b demonstrate expression of *bgaB* from the *xyl* and *spac* expression systems (induced with xylose and IPTG, respectively) as indicated by X-Gal hydrolysis in LB agar (EB107 and EB103, respectively). Strain EB104 (*spac* system without *lacI*) also showed significant X-Gal hydrolysis. Strains EB106 (*xyl*) and EB105 (*spac*) were negative-control strains, containing the corresponding expression system but lacking the *bgaB* gene, and showed no evidence of X-Gal hydrolysis. Strains containing both expression systems were plated on X-Gal in the absence of any inducer to assess transcriptional

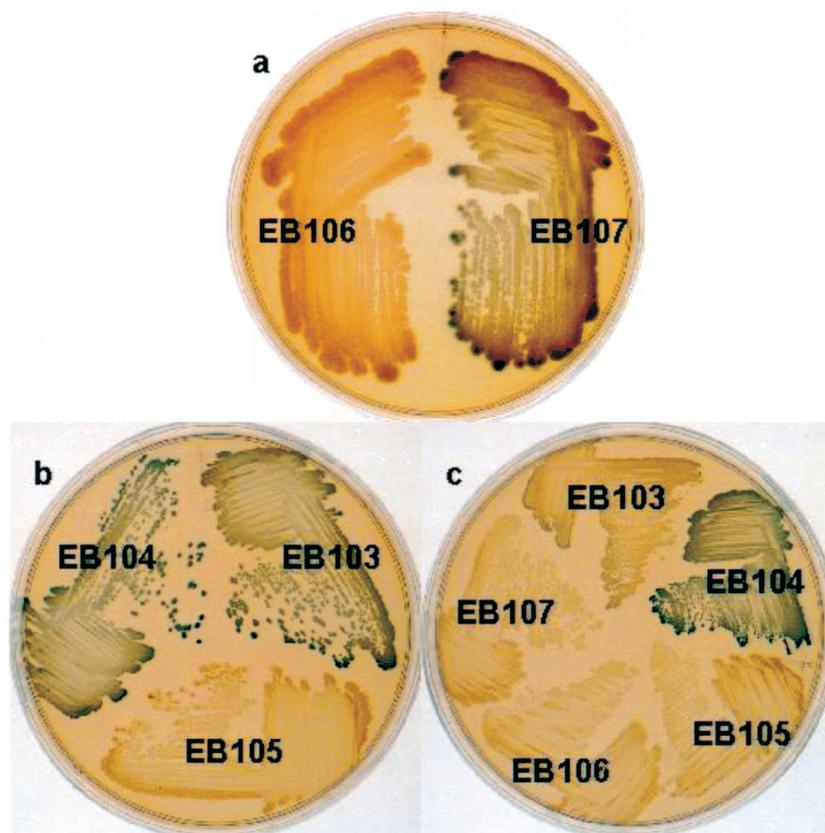


FIG. 2. Detection of reporter gene expression by X-Gal hydrolysis on solid media. (a) Strains EB106 (pSWEET) and EB107 (pSWEET-*bgaB*) were plated on LB-CHL-X-Gal in the presence of 2% xylose. (b) Strains EB103 (pSPAC-*bgaB*), EB104 (pSPAC-*bgaB* Δ *lacI*), and EB105 (pSPAC) were plated on LB-CHL-X-Gal in the presence of 1 mM IPTG. (c) All five strains were plated on LB-CHL-X-Gal in the absence of an inducer. Strains were grown overnight at 37°C and then incubated at 55°C for color development (up to 36 h).

control (i.e., the leakiness of the expression system). Figure 2c shows that the *spac* system (EB103) produced clearly discernible levels of BgaB as revealed by the extent of X-Gal hydrolysis. This cleavage is not due to endogenous β -galactosidase genes, as demonstrated by the lack of color development in the negative-control strain (EB105). The strain lacking *lacI* (EB104) served to illustrate that while the *lac* repressor allowed significant transcriptional control, it did not limit expression beyond the detection of this assay. In contrast, the *xyl* expression system (EB107) showed no discernible expression in the absence of an inducer and showed no significant deviation from the negative-control strain (EB106).

To better characterize the *xyl* and *spac* expression systems, *B. subtilis* strains carrying transcriptional fusions of *bgaB* to P_{xyl} and P_{spac} , respectively, were assayed for β -galactosidase activity after growth in liquid media. Figure 3 shows an example (β -galactosidase activities of EB104 under inducing conditions) of our efforts to ensure that the assay was linear both with time and with the volume of cells. Under the conditions used in this work, *o*-nitrophenol production remained linear for 240 min and was directly dependent on the amount of sample added to the assay. Having established parameters for a linear response in the assay of thermostable β -galactosidase, EB103 and EB107 were grown in both rich (LB) and minimal media (plus 0.2% arabinose) in the presence or absence of an

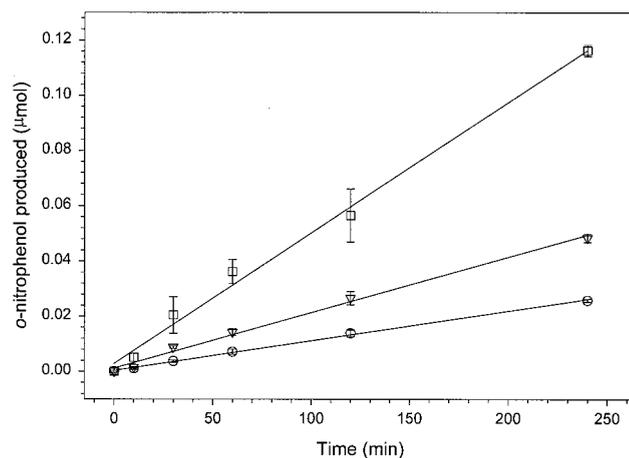


FIG. 3. Linearity of heat-stable β -galactosidase assay. Strain EB104 was grown (LB-CHL with 1 mM IPTG) to mid-log phase ($OD_{600} = 0.5$) and assayed for BgaB activity (see Materials and Methods) at different sample volumes (\circ , ∇ , and \square denote 0.1, 0.2 and 0.5 ml of culture, respectively) for up to 4 h. β -Galactosidase activity was defined as micromoles of *o*-nitrophenol released after 0, 10, 30, 60, 120, and 240 min. Slopes for 0.1, 0.2, and 0.5 ml of culture were 0.11, 0.20, and 0.47 nmol/min. Errors are standard deviations from three separate experiments.

TABLE 3. Induction and repression of *xyl* and *spac* expression systems^a

Expression system	Medium	β-Gal activity (mean ± SD)			Induction/repression ratio
		Induced ^b	Uninduced	Repressed	
<i>xyl</i> (EB107)	Minimal	12,795 ± 858	89 ± 19	52 ± 17 ^c	246
<i>xyl</i> (EB107)	Rich	12,824 ± 658	46 ± 21	ND ^d	279
<i>spac</i> (EB103)	Minimal	953 ± 43	111 ± 22	ND	9
<i>spac</i> (EB103)	Rich	814 ± 48	34 ± 20	ND	24

^a Saturated cultures EB107 (pSWEET-*bgaB*) and EB103 (pSPAC-*bgaB*) were diluted (100-fold) into fresh rich LB or minimal medium, grown to mid-log phase, and assayed as described in Materials and Methods. Errors are standard deviations from at least three separate experiments. β-Gal, β-galactosidase.

^b Medium supplemented with 2% xylose (EB107) or 1 mM IPTG (EB103).

^c Medium supplemented with 0.2% glucose.

^d ND, not determined.

inducer. The *xyl* expression system showed a significantly higher ratio of induction to repression than the *spac* system (ratios of 279 versus 24 in rich media, respectively), which can be attributed to higher levels of expression under inducing conditions (Table 3). Our analysis of the *spac* system indicates a somewhat lower induction/repression ratio than that previously recorded (140-fold) for the comparable *spac* system of pMUTIN1 (33), perhaps due to differences in the genetic contexts of each system. Overall, the *xyl* system demonstrated a 16-fold increase in expression relative to the *spac* system (β-galactosidase units of 12,824 and 814 in rich media for the *xyl* and *spac* systems, respectively). Under noninducing conditions, however, there was no significant difference between the *xyl* and *spac* expression systems using this assay (β-galactosidase units of 46 ± 21 [mean ± standard deviation] and 34 ± 20 in rich medium, respectively). Indeed, significant noise in this assay, which was particularly troublesome at the lower limits of detection, may have precluded our detection of otherwise significant differences between these two expression systems.

The role of the CRE in catabolite repression of the *xyl* operon and its capacity for optimization through mutation has been well documented (13, 16, 18, 34). Furthermore, glucose-specific, XylR-mediated repression has also been reported (18). In Table 3, we report about a 250-fold difference in reporter gene expression, comparing growth on minimal medium with xylose (2%) to growth on minimal medium plus glucose (0.2%). This result is consistent with previous studies of the *xyl* regulon in which induction/repression ratios as high as 260-fold have been reported (13). We also examined reporter gene expression after growth in minimal medium with xylose (0.2%) and one of three sugars (glucose, fructose, or glycerol) at a level of 0.2%. Regardless of the type of sugar added, we observed about a 100-fold decrease in reporter gene expression (data not shown). These findings are also consistent with a previous study in which the CRE sequence used in our studies (G→T and A→T at positions 3 and 10, respectively) resulted in similar levels of catabolite-mediated repression and masked to a large extent the glucose-specific effect seen with wild-type CREs (18). Kim et al. (17) have described a xylose-inducible integration vector using *xyl* regulatory sequences from *Bacillus megaterium*. Induction/repression ratios ranging from 150- to 200-fold were reported for that system, which lacked the CRE sequence and was not subject to catabolite

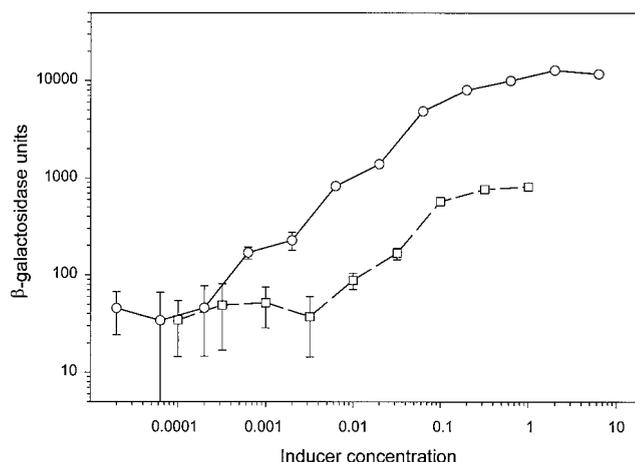


FIG. 4. Modulation of the *xyl* and *spac* expression systems. Saturated cultures of EB107 (pSWEET-*bgaB*) and EB103 (pSPAC-*bgaB*) were inoculated (1/100) into fresh LB-CHL media supplemented with increasing concentrations of inducer, i.e., 0.00002 to 6.3% xylose (○) and 100 nM to 1 mM IPTG (□). Cultures were grown to mid-log phase and assayed for BgaB activity (see Materials and Methods). β-Galactosidase units are picomoles of *o*-nitrophenol released per minute per milliliter of culture at an OD₆₀₀ of 0.5. Errors are standard deviations from at least three separate experiments.

repression. Eichenbaum et al. have also published a comparison study of a plasmid-based *B. subtilis* xylose expression system with *spac*, *lac*, and *nisA* promoters (9). In that work, the *xyl* and *spac* systems demonstrated similar induction/repression ratios in *B. subtilis* (11- and 16-fold, respectively), though few details were published regarding the construction of that xylose-based expression system.

Having determined the extremes of expression, we wanted to assess the ability of each system to modulate expression in response to an inducer. Strains EB107 and EB103 were grown in various concentrations of inducers (xylose and IPTG, respectively) in rich liquid media, and expression levels were assessed using the β-galactosidase assay (Fig. 4). The xylose-induced expression system showed a particular capacity for modulation, as β-galactosidase activity varied from 30 units to about 11,000 units over an inducer concentration range of 3.5 log units (0.0002 to 0.63% xylose). In contrast, the *spac* system was modulated over an inducer concentration range of only 1.5 log units (0.003 to 0.1 mM IPTG).

As a test case for the capacity of the *xyl* expression system to provide conditional complementation of essential functions, we attempted to complement a temperature-sensitive mutant (*tag-12*) previously attributed to *B. subtilis tagD* (23), which encodes glycerol-3-phosphate cytidyltransferase. Again, for comparative purposes, *tagD* was cloned into both pSWEET and pSPAC in order to create EB123 and EB127, which expressed wild-type *tagD* in the temperature-sensitive background under *xyl* and *spac* control, respectively. We assessed the ability of each system to complement the temperature-sensitive mutant at the nonpermissive temperature. As a control, we included the parental *tag-12* temperature-sensitive strain (EB4) and the isogenic *tag*⁺ strain (EB6) in these experiments. As expected, EB6 grew well at both the permissive (30°C) and nonpermissive (47°C) temperatures, while EB4

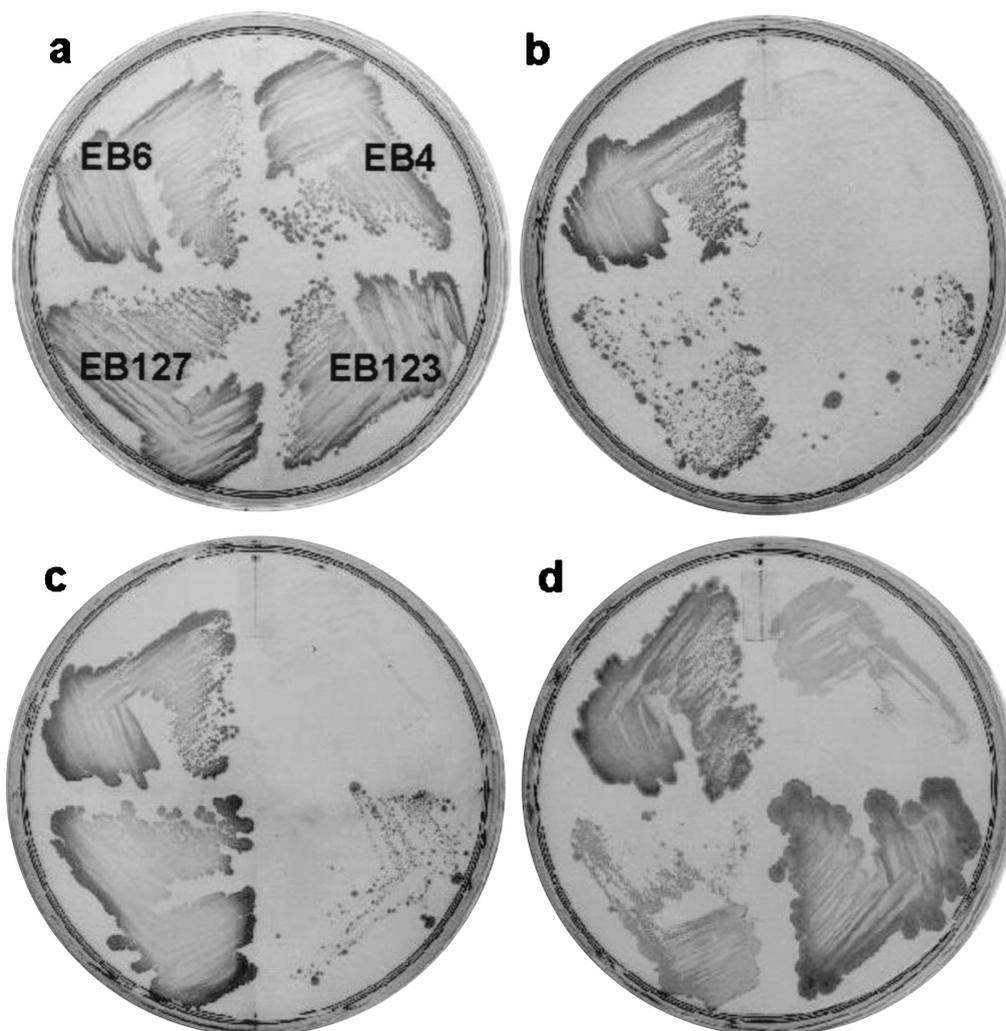


FIG. 5. Xylose-dependent complementation of a temperature-sensitive mutant in teichoic acid biosynthesis. Strains were plated on LB agar in the presence or absence of an inducer and incubated at 30°C (permissive temperature) or 47°C (nonpermissive temperature). Strains EB4 (*tag-12*), EB6 (*tag*⁺), EB123 (*tag-12* pSWEET-*tagD*), and EB127 (*tag-12* pSPAC-*tagD*) were plated at 30°C (a), plated at 47°C (b), supplemented with 1 mM IPTG at 47°C (c), and supplemented with 2% xylose at 47°C (d). Strains shown in panels b through d were plated as indicated in panel a.

showed almost no growth at the nonpermissive temperature (Fig. 5a and b). Both EB123 and EB127 were able to complement the mutant at the nonpermissive temperature in the presence of their respective inducers (Fig. 5c and d). In the absence of added inducer, the *spac* expression system (EB127) showed substantial growth at the nonpermissive temperature, while the *xyl* system (EB123) showed only very slight growth relative to the control EB4 (Fig. 5b). Interestingly, we have consistently noted that in the absence of inducer and with a heavy inoculum, a mixed population of large and small colonies is evident with strains EB123 and EB127 (e.g., Fig. 5b). These large colonies subsequently demonstrate a temperature-insensitive and inducer-independent growth phenotype and may have arisen from recombination of the temperature-sensitive copy of *tagD* at the *tag* locus with the wild-type copy of *tagD* resident at *amyE*.

To further assess the ability of the *xyl* system to conditionally complement the *tag-12* mutant, a growth curve was examined

using strain EB123 (expression of *tagD* under *xyl* control in the *tag-12* background) grown in the presence and absence of the inducer xylose (0.2%) (Fig. 6). Again, the parental strains EB4 and EB6 were included as controls. The cultures were initially grown at the permissive temperature (30°C) for 300 min and were then shifted to the nonpermissive temperature (47°C) and monitored for up to 25 h. The *tag-12* mutant (EB4) showed a lytic phenotype upon temperature shift, as indicated by the steady decrease in OD after 660 min. In the absence of xylose, strain EB123 duplicated the growth and lysis exhibited by EB4. In the presence of the inducer, EB123 showed slightly better growth than EB6, likely due to the presence of a rich carbon source (xylose). Nevertheless, we are certain that this complementation was due to the expression of *tagD* and not the presence of xylose, since, when grown at the nonpermissive temperature and in the presence of xylose, EB4 showed only very slight growth (Fig. 5d). We also tested EB127 (expression of *tagD* under *spac* control in the *tag-12* background) in an

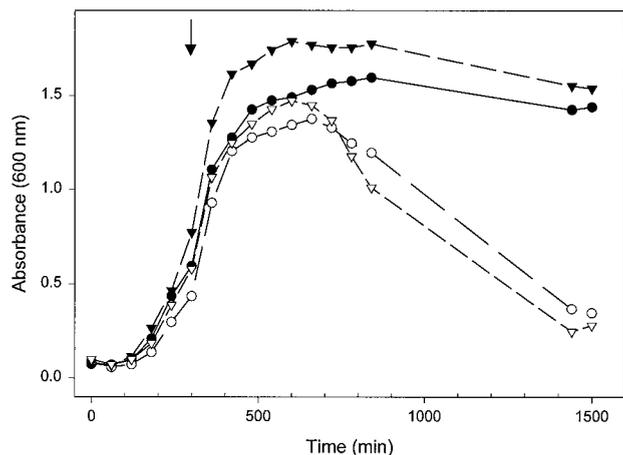


FIG. 6. Growth profile of the *B. subtilis tag-12* mutant and conditional complementation with *tagD* under xylose control. Overnight cultures (LB) of EB4 (*tag-12*), EB6 (*tag+*), and EB123 (*tag-12 pSWEET-tagD*) were inoculated (1/100) into fresh medium (LB) and grown for 25 h, with periodic monitoring of growth by OD₆₀₀. Growth curves for EB4 and EB6 are represented by the symbols ○ and ●, respectively. EB123 was grown in the presence (▼) and absence (▽) of 0.2% xylose. Growth was at 30°C (permissive temperature for *tag-12*) until mid-log phase, when cultures were shifted to 47°C (restrictive temperature for *tag-12*) at 300 min (denoted by the arrow).

analogous experiment and found that the strain exhibited a growth curve which was indistinguishable from that seen for EB123 (data not shown).

DISCUSSION

We have undertaken the development and characterization of a system for efficient expression in *B. subtilis* in order to characterize putatively indispensable functions in teichoic acid biosynthesis. To that end, we have taken advantage of an extensive body of work on the xylose utilization operon of *B. subtilis* W23, which has described in detail the machinery for xylose induction and catabolite repression in that system (5, 6, 12, 13, 16, 18, 19). Accordingly, we chose to construct a xylose-based expression system by incorporating components of the xylose operon from strain W23. The expression system contains the xylose repressor encoded by *xylR*, intergenic *xyl* operator sequences, the *xylA* promoter, and the 5' portion of *xylA* containing an optimized CRE. These components are present on plasmid pSWEET, a derivative of pDG364 (3), for integration into the *B. subtilis* chromosome at *amyE*.

We have made significant contributions to the improvement of xylose-based systems (9, 17) with the inclusion of *xyl* sequences important in catabolite repression (CREs) and with an extensive characterization of the expression system constructed. The *xyl* expression system developed was effective in achieving very low levels of induction in the absence of an inducer, was capable of a wide range of induction levels, and had the capacity for modulated expression over a broad scale of inducer concentrations. We have carefully compared the performance of the *xyl* expression system to that of the widely used *spac* system as a point of reference and found it to be superior to *spac* in each of these characteristics.

The tight control of expression afforded by the *xyl* expression

system is an attractive feature in the study of null mutations, where strongly regulated expression of a complementing copy of the gene of interest can lead to an unambiguous interpretation of phenotype. Indeed, this expression system has particular utility in the study of null mutations in essential genes. In the studies detailed here, the temperature-sensitive mutant *tag-12* showed little or no growth at the restrictive temperature. Significant growth was seen, however, at the nonpermissive temperature for the *tag-12* mutant with *spac-tagD* at *amyE* in the absence of IPTG, whereas only very slight growth was evident with the *xyl-tagD* complementation system in the absence of xylose. Were this a *tagD* null mutant, complementation experiments with the *spac* expression system might incorrectly indicate that a phenotype of impaired growth, not lethality, is associated with a *tagD* knockout.

The particular capacity for modulation of gene expression with the *xyl* system is arguably a very attractive feature for the deliberate expression of cloned genes in *B. subtilis*. While the *spac* system modulated expression over a narrow range of inducer concentrations (1.5 log units), the *xyl* system controlled expression over a >3.5-log span of xylose concentrations. When combined with the large induction/repression ratio characteristic of the *xyl* system, this broad response to inducer concentration should facilitate exploration of the effects of expression level of a cloned gene on phenotype. Indeed, protein expression levels can be important to phenotypic analyses in bacteria for a wide variety of reasons, including an exquisite sensitivity, in some cases, to protein stoichiometry, such as that observed for the *E. coli* proteins FtsZ and FtsA (7, 8).

Despite the apparent superiority of the *xyl* promoter-operator system over that of *spac*, both systems showed considerable capacity for control of expression in *B. subtilis*. Together, these two systems may be particularly useful in instances where more than one induction system may be warranted. Plasmid pMUTIN, for example, provides the means for facile inactivation of a target gene while placing downstream genes under the control of the *spac* promoter to test for polar effects (33). Complementation of the inactivated gene by placing a copy under *spac* transcriptional control would not provide the means to distinguish the effects of transcription of downstream genes from those of complementation. Use of pSWEET for this purpose would facilitate an unequivocal analysis of any phenotype(s) associated with the null mutant by placing the target gene *in trans* under the control of the xylose promoter.

The *tag* genes are responsible for the synthesis of poly(glycerol phosphate), the predominant cell wall-linked polyanionic polymer of *B. subtilis* strain 168. A considerable body of work indicates an essential role in this strain for poly(glycerol phosphate) synthesis (2, 23, 24, 26, 27). Paradoxically, two other wall polymers, poly(glucose *N*-acetylgalactosamine phosphate) and teichuronic acid, are also produced and are capable of at least partially substituting for the predominant polymer (10, 11). In fact, certain strains of *B. subtilis* have been reported to completely replace wall teichoic acid with phosphate-free teichuronic acid under phosphate-limiting conditions (20). Therefore, while teichoic acid biosynthesis may have great potential as a therapeutic drug target in gram-positive bacterial physiology, a clear understanding of the putatively essential role for this polymer, even in the model organism *B. subtilis* 168, has remained elusive.

An essential role for *tagD*, encoding glycerol-3-phosphate cytidylyltransferase (25), was indicated by the localization of two thermosensitive mutations, *tag-11* and *tag-12*, to *tagD* (23). In addition, glycerol-3-phosphate cytidylyltransferase activity in extracts of the *tag-11* mutant was shown to be thermolabile and significantly reduced relative to that of wild-type extracts (27). In the work reported here, we noted a pronounced growth defect in the *tag-12* mutant, such that this strain showed little or no growth at the restrictive temperature and demonstrated a significant drop in OD soon after a temperature shift from 30 to 47°C. The decrease in cell density upon shifting to the nonpermissive temperature is remarkable in its similarity to the lytic response normally reserved for defects in peptidoglycan biosynthesis and is consistent with previous studies detailing gross morphological changes associated with teichoic acid mutants (1, 28). Using the *xyl* expression system developed here, we have demonstrated, for the first time, *trans* complementation of a teichoic acid biosynthesis mutant. Rescue of the *tag-12* mutant at the restrictive temperature with *tagD* under *xyl* control was xylose dependent, unequivocally indicating a role for *tagD* in this growth defect. This work sets the stage for further analysis of teichoic acid biosynthesis genes in *B. subtilis* through the construction of null mutants therein and conditional complementation using pSWEET.

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ERRATA

Development and Characterization of a Xylose-Dependent System for Expression of Cloned Genes in *Bacillus subtilis*: Conditional Complementation of a Teichoic Acid Mutant

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Volume 67, no. 1, p. 403–410, 2001. Page 405, Fig. 1: the plasmid map depicted in Fig. 1a is not correct; the orientation of the plasmid backbone is the reverse of that shown in the published article. Figure 1 should read as shown below.

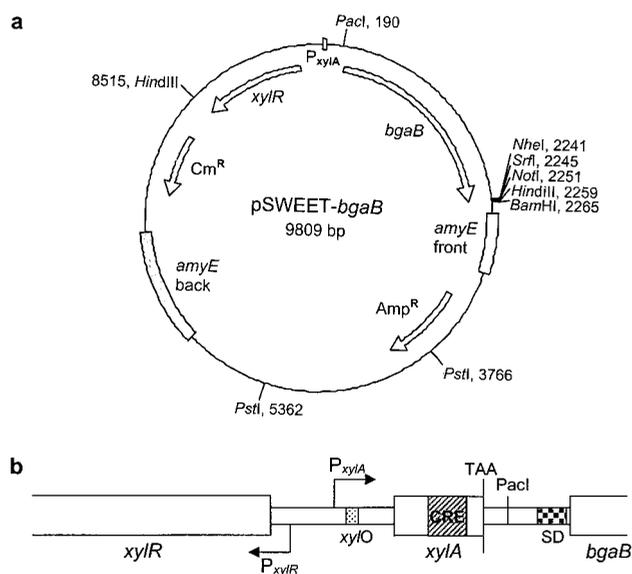


FIG. 1. Map of plasmid pSWEET-*bgaB*. (a) Significant features of the xylose-based expression system. Plasmid pSWEET-*bgaB* is a derivative of pDG364 (3), which allows integration into the *B. subtilis* chromosome at *amyE* via double recombination and selection with CHL (10 $\mu\text{g/ml}$). The plasmid also has an *E. coli* origin of replication and ampicillin resistance cassette (50 $\mu\text{g/ml}$) for routine cloning steps. On the outside of the plasmid map restriction sites of interest are highlighted, including two *PstI* sites for convenient plasmid linearization, a *PacI* site (eight-base recognition sequence TTAATTAA) upstream of *bgaB*, and a polylinker downstream of *bgaB* (*HindIII* is not unique). (b) Close-up of key elements of the xylose expression system (not to scale). Shown are *xyIR*, encoding the xylose repressor; the *xyI* intergenic region, including promoters for *xyIR* (*P_{xyIR}*), *xyIA* (*P_{xyIA}*), and *xyI* operator sequences (*xyIO*); translationally truncated *xyIA* (first 58 nucleotides followed by an in-frame TAA), including an optimized CRE in *xyIA* (see Materials and Methods and reference 18); *PacI* 5' cloning site; ribosome binding site (SD) native to *B. subtilis tagD*; and gene *bgaB*, encoding a thermostable β -galactosidase from *B. stearothermophilis*.

Biosorption of Copper by a Bacterial Biofilm on a Flexible Polyvinyl Chloride Conduit

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Volume 67, no. 9, p. 4349–4352, 2001. Page 4351, legend to Fig. 3: “×, effluent” should read “×, cell-free control.”