Characterization of *Sinorhizobium meliloti* Triose Phosphate Isomerase Genes

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A Tn5 mutant strain of *Sinorhizobium meliloti* with an insertion in *tpiA* (systematic identifier SMc01023), a putative triose phosphate isomerase (TPI)-encoding gene, was isolated. The *tpiA* mutant grew more slowly than the wild type on rhamnose and did not grow with glycerol as a sole carbon source. The genome of *S. meliloti* wild-type Rm1021 contains a second predicted TPI-encoding gene, *tpiB* (SMc01614). We have constructed mutations and confirmed that both genes encode functional TPI enzymes. *tpiA* appears to be constitutively expressed and provides the primary TPI activity for central metabolism. *tpiB* has been shown to be required for growth with erythritol; however, basal levels of TPI activity present in *tpiA* mutants allow for growth with gluconeogenic carbon sources. Although *tpiA* mutants can be complemented by *tpiB*, *tpiA* cannot substitute for mutations in *tpiB* with respect to erythritol catabolism. Mutations in *tpiA* or *tpiB* alone do not cause symbiotic defects; however, mutations in both *tpiA* and *tpiB* caused reduced symbiotic nitrogen fixation.

Triose phosphate isomerase (TPI) catalyzes the reversible interconversion of glyceraldehyde-3-phosphate (G3P) and dihydroxyacetone phosphate (DHAP). This activity makes TPI a key enzyme of central carbon metabolism, allowing it to play a role in the glycolysis (Embden-Meyerhof-Parnas [EMP]), gluconeogenesis, pentose phosphate (PP), and Entner-Doudoroff (ED) pathways. Past research on the symbiotic soil bacterium *Sinorhizobium meliloti* and other rhizobia has shown that hexose catabolism proceeds through the ED and PP pathways, while the EMP pathway functions at very low levels, if at all (49). Gluconeogenesis, however, is functional, and mutations in gluconeogenic enzymes have been shown to cause complex symbiotic phenotypes, including reduced or abolished nitrogen-fixing activity (6, 15, 16, 19, 24).

Recent work has confirmed the roles of the ED and PP pathways as catabolic pathways in *S. meliloti*. Analyses of carbon flux carried out using labeled carbon compounds and gas chromatography-mass spectrometry have confirmed the absence of the EMP pathway during growth with glucose (17). Another set of experiments done using labeled carbon and nuclear magnetic resonance is in agreement, confirming that glucose is degraded primarily through the ED pathway (22, 40). The gas chromatography-mass spectrometry and nuclear magnetic resonance experiments also confirm that some of the G3P produced by the catabolism of hexoses through the ED pathway is converted back to higher-molecular-weight compounds through TPI and fructose bisphosphate aldolase (FBA) in a cyclic pathway. Because FBA requires G3P and DHAP as substrates, the interconversion of G3P and DHAP is necessary for the cyclic metabolism of hexoses as well as gluconeogene-
sis. An *S. meliloti* strain missing TPI activity would therefore be compromised metabolically.

Based on gene homology, *S. meliloti* has two putative chromosomal genes predicted to encode TPI enzymes (18). The gene *tpiA* (SMc01023) appears to be transcribed independently, while *tpiB* (SMc01614) appears to be transcribed along with two other genes: SMc01615 and *rpiB*. A putative operon upstream of *tpiB* includes homologs of genes involved in the catabolism of erythritol in *Rhizobium leguminosarum* and *Brucella abortus* (45, 50). The *tpiB* gene and its proximity to erythritol catabolic genes are conserved in a number of organisms, which led us to hypothesize that *tpiB* is involved in erythritol catabolism. In this work we describe the isolation and characterization of TPI mutants in *S. meliloti*. We have found that both *tpiA* and *tpiB* encode functional TPI enzymes and that *tpiB* is induced by and specifically required for erythritol catabolism. Mutations in both TPI genes cause a loss of the ability to use gluconeogenic carbon sources, but viability and growth with other carbon sources or combinations of nonpermissive carbon sources remain possible. Mutations in either *tpiA* or *tpiB* alone did not affect symbiosis, while mutations in both TPI-encoding genes were found to cause reduced levels of symbiotic nitrogen fixation.

**MATERIALS AND METHODS**

**Bacterial strains, plasmids, and media.** The bacterial strains and plasmids used and generated in this work are listed in Table 1. Bacteria were grown at 30°C, using either Luria-Bertani broth (LB) or Vincent’s minimal medium (VMM), as previously described (43, 44). All carbon sources used in VMM were filter sterilized and used at a concentration of 15 mM, unless otherwise stated. All sugars were in the D configuration, except D-arabinose, D-fucose, D-rhamnose, and D-lyxose. When required, antibiotics were used at the following concentrations for *Escherichia coli*: tetracycline (Tc), 5 μg ml⁻¹; neomycin (Nm), 200 μg ml⁻¹; chloramphenicol (Cm), 20 μg ml⁻¹; and gentamicin (Gm), 20 μg ml⁻¹. Gm was used at a concentration of 50 μg ml⁻¹ for *S. meliloti*.

**DNA manipulations and plasmid constructions.** Standard techniques were used for DNA isolation, restriction enzyme digests, ligations, transformations, and agarose gel electrophoresis (44).

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To construct pNP150, a 350-bp internal fragment of *tpiB* was PCR amplified using primers 1 and 2 (Table 2) and genomic Rm1021 DNA as a template. The PCR product was gel isolated and cloned into pKNOCK-Gm by use of the 5' and 3' ends of the fragment, respectively. This region was cloned into pBluescriptII with NotI and BamHI, yielding pNP152. Similarly, a 660-bp region downstream of *tpiA* was PCR amplified from Rm1021, adding the restriction sites BamHI and SalI to the 5' and 3' ends of the fragment, respectively. The 660-bp fragment was then subcloned from pNP154 into pJQ200SK (42), and this construct was named pNP154. A 1,360-bp NotI/SalI fragment of pNP152 was then amplified from pNP154, and this PCR product was gel isolated and cloned into pKNOCK-Gm by use of the 5' and 3' ends of the fragment, respectively. This region was cloned into pBluescriptII with NotI and BamHI, yielding pNP154. A 1,360-bp NotI/SalI fragment was subcloned into pBluescriptII with NotI and BamHI, yielding pNP155. To construct pNP155, a 350-bp internal fragment of *tpiB* was PCR amplified using primers 1 and 2 (Table 2) and genomic Rm1021 DNA as a template. The PCR product was gel isolated and cloned into pKNOCK-Gm by use of the 5' and 3' ends of the fragment, respectively. This region was cloned into pBluescriptII with NotI and BamHI, yielding pNP152. Similarly, a 660-bp region downstream of *tpiA* was PCR amplified with primers 5 and 6 (Table 2), which add the restriction sites NotI and BamHI to the 5' and 3' ends of the fragment, respectively. The 660-bp fragment was cloned into pBluescriptII with NotI and BamHI, yielding pNP154. A 350-bp internal fragment of pNP154 was then amplified from pNP154, and this fragment was then cloned into pJQ200SK (42), and this construct was named pNP155.

In order to constitutively express *tpiA* and *tpiB*, both open reading frames were cloned into a broad-host-range vector, pRK7813 (25). Primers 7 and 8 (Table 2) were used to amplify *tpiA* from Rm1021, which adds a ribosome binding site (GGAG), an N-terminal RGS(His)6 tag, and BamHI and EcoRI restriction sites to the gene. Primers 9 and 10 (Table 2) were used to amplify *tpiB*, adding the same features as *tpiA*. Both *tpiA* and *tpiB* were cloned into pRK7813, yielding pNP156 and pNP167, respectively. Both constructs were sequenced entirely, showing that neither had any deviation in sequence from the published Rm1021 genome (data not shown).
Genetic techniques and mutant construction. Mutagenesis of Rm1021 with Tn5 was carried out using PRK602 as previously described (13). Conjugations between E. coli and S. meliloti were carried out using the mobilizing strain MT616 as previously described (16). Transduction with phage φM12 (12) was routinely used to move transposon markers from mutant strains into wild-type parent strains, and 50 to 100 colonies were screened to ensure that phenotypes were 100% linked in transduction and therefore associated solely with the transposon insertion.

Construction of the tpIB single crossover mutant SRmA355 was accomplished by conjugating pNP150 into Rm5000. Single crossovers were selected for by plating on VMM agar containing Rif and Gm. The pKNOCK tpIB insertion was confirmed by transductional linkage analysis with a nearby marker (data not shown). The tpia tpIB mutant SRmA366 was constructed by transducing an SRmA185 φM12 lysate into an SRmA355 background. The chromosomal deletion of tpia in SRmA449 was constructed by mating pNP155 into Rm1021 and selecting for single crossover events by use of Gm. Some of these were plated onto LB containing 5% sucrose to select for double crossover events. A sucrose-sensitive, Gm-resistance colony was purified, and the tpia deletion was confirmed by PCR amplifying the deleted region with primers 3 and 6 (Table 2), sequencing the gel-isolated PCR product, and comparing the result with the expected nucleotide sequence (data not shown). Strain SRmA515 was created by moving the Gm marker from SRmA355 into SRmA449 by use of transduction. The tpib-complementing cosmid pNP163 was isolated by conjugating an S. meliloti cosmid bank with SRmA515 and plating on a defined medium containing erythritol as a sole carbon source. The cosmid pNP163 was then mutagenized by passage through the Tn5-B20-carrying strain EcA101, selecting for cotransfer of the plasmid and Tn5-B20 into EcA100. Individual cosmids carrying Tn5-B20 were then mated into SRmA515 and screened for loss of the ability to complement growth on erythritol. One such mutagenized cosmid was isolated and shown to contain a mutation in tpib by use of a protocol described below (data not shown). Marker exchange of the cosmids-borne tpib mutation into the Rm1021 chromosome was performed as described previously (20), yielding strain SRmA584. The SRmA584 transposon marker was subsequently transferred into SRmA449 by transduction, yielding SRmA585.

Mutant identification. To identify the point of insertion for individual Tn5 mutations, a modification of an arbitrary PCR protocol was utilized, as previously described (32, 41). Briefly, strains containing Tn5 or Tn5-B20 inserts were purified and the genomic DNA was used as a template. Two sequential PCRs were performed, the first with primers 11 and 12 (Table 2) and the second with primers 13 and 14, with the products of the first reaction as the template. The final PCR product was gel isolated and sequenced with primer 13 (Table 2) to locate the point of insertion in the genome.

TPI assays. Enzyme assays were performed on cells grown in rich medium or in defined medium containing succinate, glucose, or erythritol. Cells were grown overnight and resuspended in an extraction buffer containing 100 mM Tris, 5 mM β-mercaptoethanol, and 1 mM MgCl2 at pH 7.6. A French pressure cell (16,000 psi) was used to lyse cells, and the lysate was centrifuged at 20,000 × g for 20 min. The supernatant was then filtered through a 0.22-μm filter and used for in vitro assays. Protein concentration was measured as described previously (32).

RESULTS

Analysis of tpia and tpib. As part of our ongoing interest in rhamnose catabolism and associated symbiotic phenotypes (34, 43), we have screened many S. meliloti mutants for the ability to catabolize rhamnose as a sole carbon source. A number of Tn5-induced Rm1021 mutants unable to use rhamnose were isolated, including one strain, SRmA185, which was able to grow slowly only on plates containing rhamnose and unable to grow on glycerol plates. All mutations except for that of SRmA185 were situated in a contiguous chromosomal locus (systematic identifiers Smc0182 to Smc0235 and Smc0180 to Smc0235) (data not shown). This locus was found to be homologous to a rhamnose utilization locus previously reported for R. leguminosarum (43). An SRmA185 genomic DNA fragment spanning the Tn5 insertion and flanking DNA was amplified with an arbitrary PCR protocol. Sequencing the fragment revealed that the Tn5 insertion was interrupting the putative TPI-encoding gene tpia (data not shown). The genome annotation of S. meliloti predicts that there are two chromosomal TPI-encoding genes, which have been annotated tpia1 and tpia2 (systematic identifiers Smc01023 and Smc01614, respectively) (18). In this study, we have shown that the two genes encode TPIs, but that there is a significant difference in function between them. We are therefore suggesting the names tpia1 and tpia2 to replace tpia1 and tpia2, respectively, as the new names conform more to the standard genetic nomenclature (11). Our suggested gene names are used throughout the paper.

The gene tpia is on the S. meliloti chromosome and appears to be isolated rather than transcribed within an operon (Fig. 1). The gene tpib is also located on the chromosome and appears to be in a small operon downstream of a large operon which contains homologs of the erythritol utilization genes eryA, eryB, and eryC from B. abortus and R. leguminosarum (45, 50). Over the course of our characterization of TPI in S. meliloti, strains carrying several independent tpia and/or tpib alleles were constructed (Table 1).

tpia and tpib are needed for gluconeogenesis. Wild-type strain Rm1021 and the constructed tpia and tpib mutant strains were tested for the ability to grow on a number of types of defined media containing single carbon sources (Table 3). Mutations in tpia caused a loss of the ability to utilize glycerol for growth as well as slow growth on rhamnose compared with that of the wild-type parent strain. Mutations in tpib caused only a loss of the ability to use erythritol as a carbon source. This correlates with the fact that tpib is situated downstream of a predicted erythritol catabolism operon (Fig. 1) and that tpib-dependent TPI activity is induced by erythritol (Table 4). Not only does the tpia tpib strain have a combination of the phenotypes of tpia and tpib strains, but having no TPI genes causes a loss of the ability to utilize fucose, succinate, γ-amino- butyric acid (GABA), acetate, arabinose, glutamate, and xylose. Strains with both tpia and tpib mutations were able to grow with combinations of nonpermissive carbon sources, such as glycerol and succinate or glyceral and arabinose. This is consistent with the hypothesis that gluconeogenesis occurs through FBA, as the permisive combinations of carbon sources supply both DHAP from glyceral and G3P from arabinose or succinate (3, 16, 49).

Both tpia and tpib encode TPIs. To confirm the predicted TPI functions encoded by tpia and tpib, cell extracts were prepared and assayed for enzyme activity. Cells were grown in flasks containing a rich medium or a defined medium with glucose, succinate, or erythritol as the sole carbon source. Due to their inability to grow, extracts were not prepared for

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SRmA355 (tpiB) with erythritol or SRmA366 (tpiA tpiB) with either erythritol or succinate as the sole carbon source. Assays for TPI activity indicate that tpiA mutants had less than 10% of wild-type activity when grown in rich medium or defined medium with either glucose or succinate, as shown in Table 4. However, activity levels were approximately 25% of wild-type activity levels when grown with erythritol. SRmA355 cell extract activity levels were similar to those of the wild type for all growth media tested, while SRmA366 extracts had background levels of TPI activity. Taken together, these data show that both tpiA and tpiB encode TPI enzymes and suggest that tpiA may be constitutively expressed, providing most of the cell’s TPI activity, whereas tpiB is induced by erythritol and expressed at low levels under the other conditions tested.

In vivo complementation of tpiA. Because tpiB was identified as a locus necessary for erythritol catabolism and tpiB-dependent TPI activity was induced by erythritol, it was hypothesized that induction of tpiB in a tpiA strain could rescue the ability to grow on glycerol. To test this, SRmA327 was streaked onto a number of plates of defined medium containing 15 mM glycerol and various amounts of erythritol (from 0.1 mM to 5 mM) to determine the lowest concentration of erythritol that could rescue the ability to grow on glycerol. SRmA327 was found to be able to grow as well as Rm1021 with glycerol when supplemented with at least 0.4 mM erythritol. This amount of erythritol alone does not support the growth of S. meliloti on agar plates, and the tpiA tpiB mutant SRmA366 was not able to grow with glycerol supplemented with erythritol. This indicates that tpiB is inducible by erythritol and has in vivo TPI activity with the ability to complement tpiA. Based on these results, it was hypothesized that suppressors to tpiA would easily be isolated with an up-regulated tpiB. Although more than 10^10 SRmA327 cells were plated on media with glycerol as a sole carbon source, we were unable to isolate any suppressors over the course of several independent experiments.

Constitutively expressed tpiB cannot complement tpiA. As both tpiA and tpiB were found to have in vitro TPI activity, we were interested in why tpiB mutants were unable to grow with erythritol as a sole carbon source. To determine if tpiB was directly required for erythritol catabolism, we complemented tpiA and tpiB mutations with constitutively expressed TPI genes. Both tpiA and tpiB were cloned into a broad-host-range vector, pRK7813, such that the genes were constitutively expressed (25). The resulting plasmids, pNP166 and pNP167, respectively, were mated into various strains and tested for the ability to use glycerol, succinate, arabinose, glutamate, or erythritol as the sole carbon source (Table 5). Constitutively expressed tpiB was able to complement tpiA, tpiB, and tpiA tpiB mutants for growth on all of the carbon sources tested. Constitutively expressed tpiA was able to complement the inability of tpiA mutant strains to grow on glycerol and the inability of tpiA tpiB mutants to grow on glycerol, succinate, arabinose, and glutamate. In contrast to tpiB, tpiA did not complement the inability to use erythritol caused by tpiB mutations. This

![FIG. 1. Physical maps of two S. meliloti Rm1021 chromosomal regions carrying either (top) tpiA or (bottom) tpiB. Boxes represent predicted open reading frames, with the pointed end indicating the direction of transcription.](http://jb.asm.org/)

### TABLE 3. Relevant growth phenotypes of strains used in this work on defined media

<table>
<thead>
<tr>
<th>Strain</th>
<th>Relevant genotype</th>
<th>Growth with carbon source</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td>Glc</td>
</tr>
<tr>
<td>Rm1021</td>
<td>Wild type</td>
<td>+</td>
</tr>
<tr>
<td>SRmA449</td>
<td>tpiA</td>
<td>+</td>
</tr>
<tr>
<td>SRmA584</td>
<td>tpiB</td>
<td>+</td>
</tr>
<tr>
<td>SRmA585</td>
<td>tpiA</td>
<td>tpiB</td>
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</table>

*Grown on VMM media with 15 mM of a given carbon source. Abbreviations: Glc, glucose; Suc, succinate; Gly, glycerol; Rha, rhamnose; Ery, erythritol; Fuc, fucose; Ace, acetate; Ara, arabinose; Glu, glutamate; Rib, ribose; Xyl, xylose; Lyx, lyxose. The symbol indicates the growth phenotype as follows: +, growth same as that of the wild type; ±, slow growth compared to that of the wild type; -, no visible growth.*
indicates that although tpiA and tpiB have the same enzyme activity, tpiB is specifically required for erythritol catabolism in _S. meliloti_. We note that when the same experiment was performed with SRmA584 (tpiB) and SRmA585 (tpiA tpiB), pNP167 (tpiB⁻) complementation resulted in growth on erythritol that was slower than that of the wild type. We are attributing this to the Tn5-B20 insertion in SRmA584 and SRmA585 being polar on _tpiB_ (Fig. 1), which has been shown to be necessary for erythritol catabolism in _R. leguminosarum_ (50). The complementing cosmid containing the genomic DNA surrounding _tpiB_, pNP163, was able to complement each strain, restoring wild-type growth.

**Symbiotic competence phenotypes.** _R. leguminosarum_ rhamnose catabolism mutants have previously been shown to have a deficiency in competition for nodule occupancy (34). It was hypothesized that _tpiA_ mutants would have a similar phenotype, based on their inability to grow with rhamnose at the same rate as wild-type _S. meliloti_. In addition, there have been many reports of defective symbiosis associated with the inability to use succinate as a sole carbon source (6, 15, 19, 24), suggesting that _tpiA_ tpiB strains may be unable to symbiotically fix nitrogen or may have only partial activity.

To test the hypothesis that _tpiA_ mutants would be compromised in the ability to compete with the wild type for nodule occupancy, alfalfa were inoculated with SRmA327 and Rm1021. The relative amount of each strain recovered from nodules was not significantly different from the amount of each strain found in the inoculum (data not shown). Therefore, the ability to compete for nodule occupancy is not compromised in a _tpiA_ strain, under the conditions tested.

Plants inoculated with _tpiB_ strains did not show any significant difference in dry weight compared to that of the wild type. However, plants inoculated with _tpiA_ tpiB strains reached dry weights of approximately 50% that of wild-type-inoculated plants. This reduced nitrogen fixation phenotype is very similar to the phenotype described for the succinate mutant _pckA_ (15, 35). Dry weights of plants inoculated with _Rm5439 (pckA)_ were not significantly different from those of the _tpiA_ tpiB mutant _SRmA585_ (data not shown). In order to test whether the reduced nitrogen-fixing phenotype was due to bacteria that had undergone a suppression or reversion event, _SRmA585_ was isolated from the nodules of plants from one experiment and used as an inoculum in another experiment. _SRmA585_ isolated from nodules had all of the expected genetic markers and phenotypes intact and resulted in similar plant dry weights.

**DISCUSSION**

Despite the central role of TPI in carbon metabolism, very few _tpi_ mutants have been characterized for prokaryotic organisms. In fact, to the best of our knowledge, mutations in _tpi_ have only been described previously for three organisms (2, 31, 37, 52). Of these, two are relatively closely related (2, 37, 52), and the other is associated with autotrophic growth (31). Moreover, well-characterized mutations have been studied in only a single case (52). Given that the number of genome sequences predicting multiple metabolic enzyme activities is increasing, it is imperative that a functional characterization of central metabolic pathways is carried out and not assumed. Our analysis of the two putative _S. meliloti_ genes is only the second study to show two functional _tpi_ genes within any organism and is the first within the _Rhizobiaceae_. As many organisms do not possess two TPI-encoding genes and their functions were initially unclear, we decided to investigate the physiological roles of _tpiA_ and _tpiB_.

The enzyme activities we have reported are consistent with _tpiA_ being constitutively expressed, and we hypothesize that _tpiA_ plays primarily a gluconeogenic role (Table 4). _S. meliloti_ microarray data are consistent with this and have shown that _tpiA_ is expressed at similar levels during growth with glucose, succinate, and rich media and in the bacteroid (4). Our work indicates that _tpiB_ is induced by growth with erythritol and is required for erythritol catabolism. _S. meliloti_ contains homologs of the _ery_ operon, which suggests that erythritol is catalyzed to DHAP as it is in _B. abortus_ (47). We have shown that constitutively expressed copies of _tpiA_ cannot complement the inability of _tpiB_ strains to grow with erythritol (Table 5). Interestingly, while characterizing erythritol catabolism in _B. abortus_, Sperry and Robertson (47) noted that TPI activity could not be completely uncoupled from 3-keto-1-erythronate-4-phosphate decarboxylase activity. This is consistent with the possibility of TpiB and the enzyme(s) of erythritol catabolism.
forming a metabolic complex. The existence of metabolic complexes and sequential reactions has been proposed for a number of pathways, including glycolysis (48).

Each of the genomes of *B. abortus* (8, 23), *Brucella melitensis* (10), *Brucella suis* (38), *Mesorhizobium loti* (26), *Rhizobium etli* (21), *R. leguminosarum* (51), and *S. meliloti* (18) has homologs of *tpiA* and *tpiB*. In each case, *tpiB* is followed by *tpiB* directly downstream, and in all but the genome of *R. etli*, *tpiB* and *tpiB* are directly downstream of operons containing homologs of the erythritol catabolism genes *eryA*, *eryB*, *eryC*, and *eryD*. This leads us to hypothesize that *tpiB* may be necessary for erythritol catabolism in each of these organisms, except for *R. etli*, which does not grow with erythritol and does not carry the erythritol genes (21, 50). In addition to our report that *tpiB* is necessary for growth on erythritol by *S. meliloti*, this has been confirmed for *R. leguminosarum* (50).

The fact that the *S. meliloti* genome annotation predicts that there are no TPI-encoding genes other than *tpiA* and *tpiB*, combined with the observation that TPI activity was essentially depleted in SRmA366 (Table 4), suggests that *tpiA* and *tpiB* strains are unable to perform gluconeogenesis. This is consistent with the fact that *tpiA* and *tpiB* strains are unable to grow with carbon sources that require gluconeogenesis (e.g., succinate, GABA, glutamate, arabinose, acetate, erythritol, and glycerol). That the low levels of TPI activity measured in *tpiA* mutants (Table 4) are able to allow gluconeogenic growth was somewhat surprising but is consistent with the report of *E. coli* revertants growing gluconeogenically with only 5% of wild-type TPI activity (2). Although gluconeogenic growth is possible in a *tpiA* strain, there does not seem to be enough TPI activity to allow catabolism of glycerol, which is thought to proceed through glycerol kinase and glycerolphosphate dehydrogenase forming DHAP, requiring TPI for further catabolism (3, 49).

We hypothesized that *tpiB* would be regulated negatively by the adjacent deOR-type regulator SMc01615. Our inability to isolate spontaneous mutants leading to up-regulation of *tpiB* by platting on minimal media with glycerol was surprising. The fact that over 10^10 cells were plated and no suppressors were isolated suggests that the regulation of *tpiB* may not be solely due to negative regulation by SMc01615. We note that the erythritol catabolism operon upstream of *tpiB* has been shown to be induced by xylitol, adonitol, sorbitol, and erythritol (28). As *tpiB* function is required for erythritol catabolism, these two operons may be coordinately regulated, suggesting a regulatory mechanism involving more than one event.

The first mutation in *tpiA* was isolated by screening for mutants with an impaired ability to utilize rhamnose as a sole carbon source. This led us to test the *tpiA* mutant for a symbiotic competition defect, as has been reported for *rha* mutants. Data from three independent experiments showed that there is no difference in nodulation competition between the two strains (data not shown). However, *tpiA* does not cause a complete loss of the ability to utilize rhamnose as a sole carbon source, so whether rhamnose catabolism mutants of *S. meliloti* are less competitive than the wild type remains an open question.

Consistent with the reported literature for *S. meliloti* mutants unable to grow on succinate, the *tpiA* and *tpiB* mutant was tested and shown to have a compromised symbiotic phenotype. When inoculated on alfalfa, plant dry weights reached less than 50% that of wild-type-inoculated plants, similar to the phenotype of *S. meliloti* gluconeogenic *pckA* mutants (15, 35). As *tpiA* and *tpiB* strains are unable to perform gluconeogenesis, the fact that symbiotic nitrogen fixation is not abolished suggests that gluconeogenesis is beneficial but not required in symbiosis with alfalfa, possibly due to small amounts of plant-supplied sugar reaching the bacteroid. Consistent with this hypothesis, the gluconeogenic enzyme phosphoenolpyruvate carboxykinase (PEPCK), encoded by *pckA*, is not found at detectable levels in *S. meliloti* bacteroids, although mutations in *pckA* cause reduced nitrogen fixation (15). On the other hand, *R. leguminosarum* bacteroids were found to have PEPCK activity, and a PEPCK-deficient mutant was found to have no difference in symbiosis (29).

Mutations in *pckA* of the broad-host-range *Rhizobium* sp. strain NGR234 were found to cause several phenotypes, ranging from Fix^-^ to low rates of nitrogen fixation, depending on the host species (36). It has been suggested previously that the strain- and host-dependent differences in severity of symbiotic phenotypes of bacterial gluconeogenesis mutants may be due to differences in peribacteroid membrane sugar permeability (39).

In this study, we have shown that there are two TPI-encoding genes in the *S. meliloti* genome with different functions, as *tpiA* cannot substitute for *tpiB*. The gene *tpiA* is hypothesized to be constitutively active in a gluconeogenic role, while *tpiB* is required for erythritol catabolism by an unknown mechanism. In addition, *tpiA* and *tpiB* mutants lose the ability to perform gluconeogenesis but retain the ability to symbiotically fix nitrogen at a lower level than the wild type. We are directing our efforts towards understanding the functional difference between *tpiA* and *tpiB* which leads to the specificity of *tpiB* for erythritol catabolism.

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