Identification of a Gene Cluster for the Formation of Extracellular Polysaccharide Precursors in the Chemolithoautotroph *Acidithiobacillus ferrooxidans*

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A cluster of five genes, proposed to be involved in the formation of extracellular polysaccharide (EPS) precursors via the Leloir pathway, have been identified in the acidophilic autotroph *Acidithiobacillus ferrooxidans*. The order of the genes is *luxA-galE-galK-pgm-galM*, encoding a LuxA-like protein, UDP-glucose 4-epimerase, galactokinase, phosphoglucomutase, and galactose mutarotase, respectively. The *gal* cluster forms a single transcriptional unit and is therefore an operon. Two other putative genes of the Leloir pathway, *galU*, potentially encoding UDP-glucose pyrophosphorylase, and a gene designated *galT-like*, which may encode a galactose-1-phosphate uridylyltransferase-like activity, were found unlinked in the genome. Using semiquantitative reverse transcription-PCR, the genes of the *gal* operon were shown to be expressed more during growth in iron medium than in growth in sulfur medium. The functions of *galE*, *pgm*, *galU*, and the *galT-like* gene were validated by complementation of *Escherichia coli* mutants and by in vitro enzyme assays. The data suggest that *A. ferrooxidans* is capable of synthesizing the EPS precursors UDP-glucose and UDP-galactose. In addition, genes *rfaA*, *-B*, *-C*, and *-D* were identified in the genome of *A. ferrooxidans*, suggesting that it can also synthesize the EPS precursor dTDP-rhamnose. Since EPSs constitute the major bulk of biofilms, this study may provide an initial model for the metabolic pathways involved in biofilm formation in *A. ferrooxidans* and aid in understanding the role of biofilms in mineral leaching and the formation of acid mine drainage.

Acidithiobacillus ferrooxidans is an acidophilic, chemolithotrophic, mesophilic, γ -proteobacterium that thrives at pH 2 and functions as part of a consortium of microorganisms for the industrial recovery of metals such as copper and gold (13, 25). In the environment, the microorganism is found in mine drainage, coal wastes, and other acidic sites, especially where pyrite (FeS_2) is available as an energy source. *A. ferrooxidans* can obtain its energy and electron requirements from the oxidation of various forms of reduced sulfur and ferrous iron. It can also fix nitrogen and carbon dioxide.

The attachment and adherence of *A. ferrooxidans* to mineral surfaces and the subsequent formation of biofilms are prerequisites to mineral dissolution, both in industrial operations and in natural environments (27). Biofilm formation is accompanied by the production of extracellular polysaccharides (EPSs) (28). Whereas the role of biofilm formation by *A. ferrooxidans* in metal solubilization has been actively studied (8, 28), little is known regarding the underlying genetics, biochemistry, and regulation of EPS formation by this microorganism.

Many organisms use UDP-glucose, UDP-galactose, and dTDP-rhamnose as precursors or building blocks of EPS biosynthesis (33). The galactosides UDP-glucose and UDP-galactose are synthesized from glucose-1-phosphate by two enzymes of the Leloir pathway, GalU (glucose 1-phosphate-pyrophosphorylase) and GalE (UDP-glucose 4-epimerase) (1, 2, 4, 5, 9,

20, 32). Glucose-1-phosphate is also converted to dTDP-rhamnose by the enzymes RfaA, *-B*, *-C*, and *-D* (14, 17). In addition, glucose-1-phosphate serves as a hub for channeling sugars to the formation of glycogen and connects these pathways with glycolysis, gluconeogenesis, and (ultimately) CO_2 fixation.

Since nothing was known about the early steps in EPS biosynthesis in *A. ferrooxidans* and how these might be connected to general sugar management, we undertook bioinformatic and functional analyses of potential genes involved in both the formation of glucose-1-phosphate and its metabolic connections to the biosynthesis of the EPS precursors UDP-glucose, UDP-galactose, and dTDP-rhamnose.

This study addresses the genetic and biochemical underpinnings of the formation of EPS precursors in *A. ferrooxidans*. Since EPS typically constitutes the major mass of biofilms, it is hoped that this study will lay the foundation for understanding the formation of biofilms by this organism in both natural habitats and industrial metal recovery operations.

MATERIALS AND METHODS

Bacterial strains, plasmids, and media. *A. ferrooxidans* ATCC 23270 was grown at 30°C, pH 3.5, in minimal-salt medium 9K (iron free) supplemented with elemental sulfur (11, 29). *Escherichia coli* strains and plasmids used are listed in Table 1. They were grown in Luria broth (LB) or plated on LB solidified with 12 g of agar per liter. Selection for *E. coli* cells, transformed with the appropriate plasmids, was performed by using LB plates with of 100 μg of ampicillin per ml. MacConkey-Gal plates, containing MacConkey agar base (Difco Laboratories) and 5 g/liter of galactose, and MM-gal, a minimal medium (19) supplemented with 5 g/liter galactose, were used to determine the ability of galactose utilization by these strains. Red colonies in MacConkey-Gal plates and grown in MM-gal were scored as galactose positive (22).

DNA isolation and sequencing, plasmid constructions, and other manipula-

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TABLE 1. Bacterial strains and plasmids used

Strain or plasmid	Relevant genotype or phenotype ^a	Source or reference
Strains		
<i>A. ferrooxidans</i> ATCC 23270 (type strain)		American Type Culture Collection
<i>Escherichia coli</i> DH5 α	F ⁻ Δ (<i>lacZYA-argF</i>) <i>endA1 hsdR17</i> (r _K m _K ⁺) <i>supE44 thi-1 recA1 gyrA96 relA1 ϕ80lacZ ΔM15</i>	
S491	F ⁻ <i>galT22 his lac proA tonA uraP</i>	National Institute of Genetics; SHIGEN
HB101	F ⁻ λ^- <i>ara-14 galK2 hsdS20</i> (r-B,m-B) <i>lacY1 leu ml-1 proA2 recA13 rpsL20 supE44 thi xyl-5</i>	1
CSH41	F ⁺ Δ (<i>lac-pro</i>) <i>galE thi</i>	National Institute of Genetics; SHIGEN
JW1091	λ^- <i>lacZ125 relA1 spoT1 trp-49 zbf-507::Tn10 pgm/zbf-507::Tn10</i>	National Institute of Genetics; SHIGEN
FF4001	MC4100 <i>galU95</i>	21
Plasmids		
PKK233-3	Amp ^r	Pharmacia Biotech
pUC18	Amp ^r	Gibco BRL
pgalT-1	Vector pUC18 with sequence of <i>A. ferrooxidans</i> that expresses GalT activity	This study
pgalU-1	Vector PKK 233-3 with <i>galU</i> of <i>A. ferrooxidans</i>	This study
ppgm-1	Vector PKK 233-3 with <i>pgm</i> of <i>A. ferrooxidans</i>	This study
pgalE	Vector PKK 233-3 with <i>galE</i> of <i>A. ferrooxidans</i>	This study

^a Amp^r, ampicillin resistant.

tions. Chromosomal DNA was isolated from *A. ferrooxidans* ATCC 23270 as described previously (34). Other DNA manipulations, plasmid preparations, electroporations, and DNA sequencing were performed according to standard procedures as described by Sambrook et al. (26). The expression vector pKK233-3 was digested with EcoRI and subsequently repaired with the Klenow fragment of DNA polymerase I to generate blunt ends, which were dephosphorylated with calf intestinal alkaline phosphatase (New England Bio Labs) prior to being cloned. PCR fragments that correspond to *galE*, *galU*, *pgm*, and *luxA* genes were amplified with DNA primers whose sequences appear in Table 2. Each PCR fragment and the pKK233-3 vector were ligated using T4 DNA ligase. The constructed plasmids were named pgalE, pgalU, ppgm, and pgalT (Table 1). Restriction enzyme digestion EcoRI (Invitrogen), Klenow (large) fragment of DNA polymerase I (Invitrogen), T4 DNA ligase (Promega), and calf intestinal alkaline phosphatase were used under conditions recommended by the suppliers.

TABLE 2. Primers used for PCR experiments and gene cloning and RT-PCR experiments

Gene	Position ^a	Primer used for:
PCR		
<i>luxA</i>	A	5' ATGCGGCATTTATTTTGGAGG 3'
	B	5' CGGTGACCAGCACCTGC 3'
<i>galE</i>	A	5' ATGCAGGTGTTGGTCACCGCG 3'
	B	5' CAAAGTTCATAAGCCATTCTGCCACG 3'
<i>pgm</i>	A	5' ATGCGCGTATTGCAGATTG 3'
	B	5' TCAGGTGATGACAGTGGGG 3'
<i>galU</i>	A	5' ATGGCTGAAGTGCAGCAAGGC 3'
	B	5' ACCCACCACCGGACGGGC 3'
RT-PCR		
<i>rhn</i>	1	5' ATGTCCCGAAAGCGATGGT 3'
<i>luxA</i>	2	5' TAAATGCCGCATCCACCCG 3'
	3	5' ATGCGGCATTTATTTTGGAGG 3'
<i>galE</i>	4	5' CGGTGACCAGCACCTGC 3'
	5	5' ATGCAGGTGTTGGTCACCGCG 3'
<i>galK</i>	6	5' TCACAAGGCAAGCCCTGGGTG 3'
	7	5' GACAACCTTGCTCATCATATTCCTGC 3'
<i>pgm</i>	8	5' TGCAATACCGCCATGGCA 3'
	9	5' ATGGCGGTATTGCAGATTG 3'
<i>galM</i>	10	5' GCCAATGGTATCGAGCAGT 3'
	11	5' GACAACCGCATGGGCTAC 3'
<i>mgt</i>	12	5' CACTTCCGTGGGCTC 3'

^a Position relative to genes shown in Fig. 1A and B.

Isolation of a *galT*-like complementing gene. DNA of *A. ferrooxidans* was partially digested with Sau3A (Gibco BRL) according to the manufacturer's directions, yielding fragments with an average size of approximately 1 to 3 Kb. These fragments were cloned into pUC18, and the resulting library was used to transform *E. coli* strain S491 (Table 1). After overnight incubation at 37°C on MacConkey-Gal plates, red colonies were picked and grown in MM-gal with 100 μ g of ampicillin/ml (21). A colony, galT-1, was selected and DNA was prepared for DNA sequencing.

Isolation of RNA and RT-PCR. Total RNA was isolated from cells of *A. ferrooxidans* grown in 9K medium supplemented with S or FeSO₄ to mid-log phase by the method of Hagen and Young (12) as modified by Guacucano et al. (10). Reverse transcription-PCR (RT-PCR), with various control reactions, was carried out as previously described (10). The sequences of the RT and PCR primers used are provided in Table 2.

Preparation of cell extracts. Fifty milliliters of a fresh LB culture of *E. coli* grown for 12 h (late log phase) were harvested by centrifugation (6,000 rpm for 20 min; 4°C), and the cell pellet was suspended and washed twice with 0.05 M potassium phosphate, pH 7. Cells were suspended in 20 ml of 0.05 M potassium phosphate, pH 7, and disrupted by sonication (sonic power, 375 W; output control, 10) at 4°C for a total of 6 min (12 30-s sonication pulses and 30 s of rest in an ice bath with a Bronson 450 sonifier) (7). Cell debris was removed by centrifugation (6,000 rpm for 20 min; 4°C). The protein concentration in the supernatant was determined using the Bio-Rad protein assay based on the method of Bradford (3).

Enzyme assays. All enzyme assays were performed at 30°C in a total volume of 1 ml with freshly prepared cell extracts. The formation of NADPH (α -phosphoglucosyltransferase activity and galactose 1-phospho-uridyltransferase activity) or NADH (UDP-galactose 4-epimerase activity and UDP-glucose pyrophosphorylase activity), products of the coupled reaction, were determined by measuring the increase in absorbance at 340 nm at different times (7). The blank consisted of the reaction mixture without cell extract.

α -Phosphoglucosyltransferase (EC 2.7.5.1) activity was measured in a reaction mixture consisting of 179 mM glycylglycine (pH 7.4), 0.67 mM β -NADP, 0.02 mM glucose 1,6-diphosphate, 30 mM MgCl₂, 43 mM L-cysteine, 1 U glucose 6-phosphate dehydrogenase, and 300 μ l of cell extract. The reaction was initiated by the addition of 5.0 mM α -glucose 1-phosphate (7).

UDP-galactose 4-epimerase (EC 5.1.3.2) activity was measured in a reaction mixture consisting of 400 mM Tris-HCl buffer (pH 8.5), 5 mM MgCl₂, 0.5 mM NAD, 0.015 U UDP-glucose dehydrogenase, and 300 μ l of cell extract. The reaction was started by the addition of 0.2 mM UDP-galactose (7).

UDP-glucose pyrophosphorylase (EC 2.7.7.9) activity was measured in a reaction mixture consisting of 50 mM Tris-HCl buffer (pH 7.5), 8 mM MgCl₂, 1.58 mg cysteine hydrochloride (pH 7.5), 0.5 mM NAD, 1.25 mM UTP, 0.015 U UDP-glucose dehydrogenase, and 300 μ l of cell extract. The reaction was initiated by the addition of 1 mM α -glucose 1-phosphate (7).

Galactose 1-phospho-uridyltransferase (EC 2.7.7.12) activity was measured

TABLE 3. List of candidate genes proposed to be involved in the formation of EPS precursors in *A. ferrooxidans* and their proposed activities

Gene ^a	Proposed enzyme activity ^b	Best BlastP hit ^c	% Sim ^d	Score ^e	E value ^f	Motif(s) ^g
<i>galE</i>	UDP-glucose epimerase (EC 5.1.3.2)	<i>Mycobacterium avium</i>	43	93	7e-18	pfam 1370; COG 1087
<i>galK</i>	Galactokinase (EC 2.7.1.6)	<i>Streptomyces coelicor</i>	48	106	5e-22	PD 339735
<i>pgm</i>	Phosphoglucosyltransferase (EC 5.4.2.2)	<i>Synechococcus elongatus</i>	77	711	0.0	pfam 2878; COG 0033
<i>galM</i>	Aldose-1 epimerase (EC 5.1.3.3)	<i>Sinorhizobium meliloti</i>	52	33.9	1.1	
<i>galT-like</i>	Possible galactose-1-phosphate uridylyltransferase (EC 2.7.7.10)	<i>Kineococcus radiotolerans</i>	42	32	4.3	COG 1085
<i>galU</i>	UDP-glucose pyrophosphorylase (EC 2.7.7.9)	<i>Burkholderia fungorum</i>	73	371	2e-73	pfam 0483; COG 0451; PD 01252
<i>luxA-like</i>	Possible coenzyme F420-dependent reductase	<i>Bradyrhizobium japonicum</i>	44	157	3e-37	pfam 0296; COG 2141
<i>rfaA</i>	d-TDP-glucose pyrophosphorylase (EC 2.7.7.24)	<i>Azotobacter vinelandii</i>	80	406	1e-112	pfam 0483; COG 1209
<i>rfaB</i>	d-TDP-glucose 4,6-dehydratase (EC 4.2.1.46)	<i>Pseudomonas putida</i>	72	443	1e-123	pfam 1370; COG 1088
<i>rfaC</i>	d-TDP-4-dehydrorhamnose 3,5-epimerase (EC 5.1.3.13)	<i>Pseudomonas stutzeri</i>	79	245	3e-64	pfam 0908; COG 1898
<i>rfaD</i>	d-TDP-4-dehydrorhamnose reductase (EC 1.1.1.133)	<i>Synechococcus elongatus</i>	63	261	1e-68	pfam 4321; COG 1091

^a Proposed gene name.^b Proposed enzyme activity.^c Organism with the best BlastP hit to the candidate gene.^d Percentage of similarity (% Sim) of candidate gene to that found in the organism listed in row (c).^e Score of BlastP match.^f E value of BlastP match.^g Motif and domains identified in the candidate proteins. Pfam, protein families; PD, Prodom (protein domains).

in a reaction mixture consisting of 100 mM TEA buffer (pH 7.8) containing 10 mM MgCl₂, 1 mM β-NADP, 0.25 mM glucose 1,6-diphosphate, 5 U of glucose 6-phosphate dehydrogenase, 3 of phosphoglucosyltransferase, and 300 μl of cell extract. The reaction was started by the addition of 1 mM UDP-glucose and 1 mM galactose 1-phosphate (15).

Bioinformatic analysis. Known metabolic pathways involved in galactose catabolism were obtained from BIOCYC (www.biocyc.org), KEGG (www.genome.ad.jp/kegg/), and ERGO (http://ergo.integratedgenomics.com/ERGO/). Amino acid sequences derived from genes identified as being involved in galactose metabolism were used as query sequences to search the partial genome sequence of *A. ferrooxidans* ATCC 23270 in the TIGR (www.tigr.org/) and ERGO databases using TBLASTN and BlastP, respectively. When a prospective candidate gene was identified in TIGR or ERGO, its predicted amino acid sequence was then used to formulate a BlastP (www.ncbi.nlm.nih.gov) search of the nonredundant database at NCBI. Only bidirectional best hits were accepted as evidence for putative orthologs. Candidate genes and their translated proteins were further characterized employing the following bioinformatic tools: Block Maker (http://blocks.fhrc.org/blocks/make_blocks.html), Pfam, Prosite, and domain predictions (http://motif.genome.jp/).

Nucleotide sequence accession numbers. The following nucleotide sequences reported in this paper have been assigned GenBank accession numbers: AY751082 (*galT-like*) and AY789510 to AY789512 for *galE*, *galU*, and *pgm*, respectively.

RESULTS

Identification and characterization of candidate genes involved in the formation of EPS precursors. The following candidate genes and their predicted protein products, potentially involved in the biosynthesis of glucose-1-phosphate and its conversion to the EPS precursors (UDP-glucose, UDP-galactose, and dTDP-rhamnose), were identified in the genome sequence of the type strain of *A. ferrooxidans* ATCC 23270 by bioinformatic analysis: *galM* (galactose mutarotase; EC 5.1.3.3), *galK* (galactokinase; EC 2.7.1.6), *galU* (glucose 1-phosphate-pyrophosphorylase; EC 2.7.7.9), two copies of *galE* (UDP-glucose 4-epimerase; EC 5.1.3.2), *pgm* (phosphoglucosyltransferase; EC 5.4.2.2), *rfaA* (dTDP glucose pyrophosphorylase; EC 2.7.7.24), *rfaB* (dTDP glucose-4,6-dehydratase; EC

4.2.1.4.6), *rfaC* (dTDP 4-hydroxymannose-3,5-epimerase; EC 5.1.3.13), and *rfaD* (dTDP hydroxymannose reductase; EC 1.1.1.133) (Table 3). The similarity of these candidate genes with their database matches, the predicted presence of diagnostic functional motifs and domains, and their suggested cluster of orthologous genes (COG) designations are listed in Table 3. No candidate gene with statistically significant similarity to *galT* was detected in the genome of *A. ferrooxidans* by bioinformatic analysis. However, a potential gene encoding a GalT-complementing activity (galactose 1-phosphate-uridylyltransferase; EC 2.7.7.12) was experimentally isolated (see below).

Organization and expression of candidate genes potentially involved in the metabolism of glucose-1-phosphate. Candidate genes *luxA-like*, *galE*, *galK*, *pgm*, and *galM* proposed to be involved in glucose-1-phosphate metabolism were found to be organized in a gene cluster (Fig. 1A). The cluster was flanked by the potential genes *rbn* and *mgt*, potentially encoding an oligoribonuclease and magnesium transporting ATPase, respectively. A possible pseudo-tRNA gene was detected between *galM* and *mgt*. RT-PCR experiments showed that *luxA-like*, *galE*, *galK*, *pgm*, and *galM* are cotranscribed, demonstrating that they form an operon, whereas the flanking *rbn* and *mgt* genes did not constitute part of this operon (Fig. 1B).

A comparison of the organization of the *A. ferrooxidans* *gal* operon with other known or putative *gal* operons indicates a considerable degree of conservation of genes, although not necessarily preservation of gene order (Fig. 1C). This evidence supports the proposed function of the *A. ferrooxidans* *gal* operon and permits a focus to be placed on identifying genetic regulatory features in future studies.

Expression of the *gal* operon during growth in sulfur and iron. Several genes have been demonstrated to be differentially

FIG. 1. Organization and cotranscription of the *gal* operon of *A. ferrooxidans*. (A) The *gal* operon consists of *luxA-like*, *galE*, *galK*, *pgm*, and *galM* flanked by the genes *rbn* and *mgt*. A potential pseudo-tRNA is located between *galM* and *mgt*. The direction of transcription is indicated by the block arrows. Locations of PCR primers are shown below the operon. (B) Determination of cotranscription of the genes comprising the *gal* operon by RT-PCR. Gel electrophoresis of DNA fragments amplified by RT-PCR using purified RNA as a substrate (a) or PCR using genomic DNA as a substrate (b). Arrows indicate the predicted sizes of the amplified DNA fragments in base pairs. (C) A comparison of the organization of the *gal* operon of *A. ferrooxidans* with the *gal* operons of *E. coli* (18), *Lactobacillus casei*, and *Klebsiella pneumoniae*. The organization of these operons was derived from the Integrated Genomics web site (www.integratedgenomics.com).

expressed in *A. ferrooxidans* grown in either iron or sulfur (16). With the goal of determining whether the oxidizable energy source also influenced the expression of genes potentially involved in galactose metabolism, a comparison was made by semiquantitative PCR of the expression of genes in the *gal* operon when *A. ferrooxidans* was grown in the presence of iron or sulfur. As shown in Fig. 2, four of the five genes of the *gal* operon (*galE*, *galK*, *pgm*, and *galM*) showed higher levels of

transcription when cells were grown in iron rather than in sulfur-containing medium. The fifth gene of the *gal* operon, the *luxA-like* gene, was not tested.

Analysis of the *luxA-like* gene. The predicted protein potentially derived from *luxA-like* exhibits similarity with the LuxA family of bacterial luciferase-like monooxygenases and contains the pfam00296 domain characteristic of this family (6, 31). It also contains COG2141, characteristic of the coenzyme

FIG. 2. Expression of several genes of the Gal cluster of *A. ferrooxidans* grown in the presence of sulfur (S⁰) or iron (Fe²⁺) as determined by semiquantitative PCR using 20, 25, or 30 cycles, respectively. PCR primers used are shown in Table 2.

F_{420} -dependent $N5,N10$ -methylene tetrahydromethanopterin reductase and related flavin-dependent oxidoreductases. For example, the domain is found in glucose-6-phosphate dehydrogenase from *Rhizobium* sp. (30). The *A. ferrooxidans* LuxA-like protein exhibited weak similarity (47%) with this protein (BlastP score = 79; E value = $1e-13$), which initiates the pentose phosphate pathway (23, 24). Therefore, a possible function of the LuxA-like protein could be to connect the Leloir pathway with the pentose phosphate pathway. Supporting this conjecture is the presence of the putative *pgm* gene in the same operon as *luxA-like*, encoding a PGM protein involved in the synthesis of glucose-6-phosphate. The latter is a major point of entry of carbon into the pentose phosphate pathway.

An alternative hypothesis is that *luxA-like* encodes the missing GalT function typically found in the *gal* operon but not detected in the *A. ferrooxidans* *gal* operon. However, in an in vitro enzyme assay, LuxA-like did not exhibit galactose 1-phosphate-uridylyltransferase activity, diminishing the likelihood of this conjecture.

Identification of a *galT-like* gene. Since no candidate *galT* gene was detected in the genome of *A. ferrooxidans* by bioinformatic analysis, it was decided to explore the possibility of the existence of GalT activity by direct experimental techniques. DNA of *A. ferrooxidans* was partially digested with Sau3A (1- to 3-Kb fragments) and was inserted into pUC18. The resulting recombinant library was used to transform a strain of *E. coli* (S491) lacking *galT*. A plasmid that was able to complement the missing GalT function was isolated and designated GalT-1. The insert in GalT-1 was sequenced, and the resulting sequence was used to identify a corresponding open reading frame in the partial genome sequence of *A. ferrooxidans* (TIGR) using BlastN. This open reading frame was designated *galT-like*.

Further analysis of the partial genome sequence revealed that *galT-like* was embedded in a gene cluster that potentially encodes other genes that may be involved in galactose metabolism, although their bioinformatic identification by BlastP was below a statistically significant threshold and could only be detected by Psi-BLAST (analysis not shown). The predicted GalT-like product was 120 amino acids long, which is shorter than most GalT proteins (340 amino acids). It also lacked the pfam01230 domain that includes the characteristic histidine triad of the GalT family (18).

Complementation of *E. coli* mutants with genes from *A. ferrooxidans*. *E. coli* strains CSH41(lacking *galE*), JW1091(lacking *pgm*), FF4001(lacking *galU*), and S491(lacking *galT*) were independently transformed with pKK233-3 containing cloned candidate genes from *A. ferrooxidans* potentially encoding GalE, PGM, GalU, and GalT, respectively, and giving rise to the recombinant plasmids pgalE-1, ppgm-1, pgalU-1, and pgalT-1, respectively. In the first three cases, the cloned genes were prepared by PCR of *A. ferrooxidans* genomic DNA using primers designed from the predicted start sites to the predicted stop sites as identified by bioinformatic analysis of the genome sequence. Expression of these cloned inserts was presumed to occur from the vector promoter. Each of these recombinant plasmids was capable of complementing the respective *E. coli* mutant, as shown by comparing growth with that of wild-type *E. coli* DH5 α (Fig. 3).

FIG. 3. Growth at 37°C in minimal medium supplemented with 0.5% galactose of *E. coli* mutants with or without vectors containing *A. ferrooxidans* genes. $\square$, wild-type *E. coli* DH5 α ; $\blacksquare$, mutant *E. coli* strain CSH41 (lacking *galE*), JW1091 (lacking *pgm*), FF4001 (lacking *galU*), or S491 (*galT*); $\blacklozenge$, CSH41 (lacking *galE*) with pgalE-1; $\times$, JW1091 (lacking *pgm*) with ppgm-1; $\blacktriangle$, FF4001 (lacking *galU*) with pgalU-1; and $\circ$, S491 (lacking *galT*) with pgalT-1.

In contrast, pgalT-1 was derived by complementation of *E. coli* S491 (lacking *galT*) using a randomly prepared library of *A. ferrooxidans* genomic DNA in pUC18. It was capable of complementing *E. coli* S491 (lacking *galT*) (Fig. 3), but growth of the recombinant strain was slower than that of *E. coli* DH5 α . It is possible that the proposed *galT* was transcribed from an *A. ferrooxidans* promoter, resulting in lower levels of transcription or, alternatively, that the *A. ferrooxidans* GalT was less efficient than *E. coli* GalT for reasons of translation inefficiency, instability, or reduced enzyme activity.

Assays of enzyme activities from *E. coli* harboring cloned *A. ferrooxidans* genes. *E. coli* strains CSH41(lacking *galE*), JW1091(lacking *pgm*), FF4001(*galU*), and S491(lacking *galT*) harboring the respective complementing plasmids pgalE-1, ppgm-1, pgalU-1, and pgalT-1 were grown overnight, and the enzymatic activities of GalE, PGM, GalU and GalT were analyzed as described in Materials and Methods (Table 4). These activities were compared to those derived from the appropriate *E. coli* mutant strain and from wild-type strain DH5 α (Table 4). *E. coli* mutant strains harboring the proposed *galE*, *pgm*,

TABLE 4. Enzyme activity measurements

Strain (relevant mutation)	Enzyme activity ^a		
	Negative control ^b	Positive control ^c	+ Complementing plasmid ^d
CSH41 (<i>galE</i>)	0	68 $\pm$ 5	109 $\pm$ 2 (pgalE-1)
JW1091 (<i>pgm</i>)	27 $\pm$ 1	60 $\pm$ 12	121 $\pm$ 7 (ppgm-1)
FF4001 (<i>galU</i>)	0.5 $\pm$ 0.1	19 $\pm$ 6	45 $\pm$ 7 (pgalU-1)
S491 (<i>galT</i>)	5 $\pm$ 1.0	10 $\pm$ 2	18 $\pm$ 1 (pgalT-1)

^a Enzyme activity expressed as the number of nanomoles per minute per milligram of total cell protein. Each value is the average of three measurements ($\pm$ standard deviation). All strains were grown in 50 ml of LB medium at 37°C for 12 h with 0.5% galactose at 30°C.

^b Enzyme activities in mutant strains transformed with the vector pKK233-3 but lacking the *A. ferrooxidans* insert.

^c Enzyme activities of *E. coli* DH5 α (wild type for each of the respective enzyme activities).

^d Enzyme activities of mutant strains complemented with the plasmid indicated in brackets carrying the respective *A. ferrooxidans* insert.

FIG. 4. Schematic representation of proposed pathways (solid arrows) and potential genes involved in the biosynthesis of glucose-1-phosphate (circled) and its role in the production of the precursors of EPS formation in *A. ferrooxidans*. Dotted arrows indicate suggested connections between glucose-1-phosphate and glycogen anabolism and catabolism, pentose phosphate pathway, glycolysis, gluconeogenesis, and the formation of EPS via glycosyltransferases (GTs).

and *galU* of *A. ferrooxidans* exhibited more activity for each of the respective enzymes than *E. coli* DH5 α , possibly because of the presence of more enzyme mass due to expression of a multicopy plasmid. *E. coli* S491 harboring the potential *galT-like* gene of *A. ferrooxidans* also exhibited slightly more activity for GalT than did *E. coli* DH5 α . Enzyme activities are measured in late-log cells where the growth of *E. coli* S491 containing *galT-like* was similar to that of *E. coli* DH5 α (Fig. 3), which suggests that the initial reduction in growth rate observed for *E. coli* S491 containing *galT-like* could reflect the less-efficient induction of the *A. ferrooxidans* *galT-like* gene in a heterologous host than that of the native gene and not to some intrinsic property of the GalT-like enzyme.

DISCUSSION

Inspection of the partial genome sequence of *A. ferrooxidans* provided by the Institute for Genome Research and by Integrated Genomics reveals the presence of a number of potential genes that exhibit significant similarity to genes known to be involved in the metabolism of glucose-1-phosphate and its conversion to the EPS precursors UDP-glucose, UDP-galactose, and dTDP-rhamnose via the Leloir pathway (Table 3). Figure 4 shows a working model for how these genes and their products might be connected in the formation of EPS precursors in *A. ferrooxidans*.

The genes that have been identified include *galE*, *galK*, *pgm*, and *galM*, organized in a gene cluster that is cotranscribed (Fig. 1A and B) and thus constituting an operon (*gal* operon) in *A. ferrooxidans*. This *gal* operon exhibits conservation of gene organization with other well-described *gal* operons (Fig. 1C). Two of the genes of the *gal* operon, *galE* and *pgm*, were shown

to complement *E. coli* mutants lacking these functions (Fig. 3) and also to encode GalE and Pgm activities, respectively, as determined by enzyme assays (Table 4). Another gene that has been identified in this study that probably plays a role in glucose-1-phosphate metabolism but does not form part of the *A. ferrooxidans* *gal* operon is a potential *galU*. The putative GalU protein has significant sequence similarity to GalU proteins in other organisms and exhibits GalU enzymatic activity in vitro (Table 4). It also complements an *E. coli* mutant lacking *galU* (Fig. 3).

Semiquantitative PCR experiments demonstrate that the genes of the *A. ferrooxidans* *gal* operon are expressed more when the microorganism is grown in the presence of Fe²⁺ as an energy source than with S⁰ (Fig. 2). It has been established that the energy substrate used by *A. ferrooxidans* influences the quantity and chemical composition of the EPS formed (8, 26). It was proposed that variations in EPS formation demonstrate the need for different mechanisms of adhesion of the microorganism to diverse substrates encountered in the environment. We speculate that the differential expression of genes of the *gal* operon reported here reflect this need and suggest how it might be accomplished.

A *galT* gene could not be detected initially in the genome sequence of *A. ferrooxidans* by bioinformatic analysis. However, an *A. ferrooxidans* sequence (*galT-like*) complementing growth of *E. coli* S491 (Fig. 3) and providing GalT enzymatic activity (Table 4) was experimentally identified from a shotgun plasmid library of *A. ferrooxidans* DNA cloned into *E. coli* S491. The sequence of *galT-like* revealed very weak similarity with known *galT* genes (Table 3), although it was considerably shorter than typical *galT* and did not exhibit the GalT prosite motif.

Putative genes *rfbA*, *-B*, *-C*, and *-D* potentially encoding enzymes that convert glucose-1-phosphate to dTDP-rhamnose were identified by bioinformatic analysis in the genome of *A. ferrooxidans* and exhibit significant sequence similarity to enzymes in other organisms (Table 3). However, their function awaits experimental validation.

More troublesome is the identification of the function of the *luxA*-like gene that forms part of the *A. ferrooxidans* *gal* operon. It exhibits weak similarity to an F_{420} -dependent glucose-6-phosphate dehydrogenase that catalyzes the conversion of glucose-6-phosphate to 6-phosphogluconolactone. *pgm* also forms part of the *gal* operon; its role is the conversion of glucose-1-phosphate to glucose-6-phosphate, which could then be converted to 6-phosphogluconolactone by the hypothetical product of the *luxA*-like gene and subsequently enter the pentose phosphate pathway. Thus, a speculative function for the LuxA-like product could be to connect glucose-1-phosphate metabolism with the pentose phosphate pathway (Fig. 4). If LuxA-like exhibited reverse activity, it could catalyze the conversion of 6-phosphogluconolactone to glucose-6-phosphate, and then it could help channel products of CO_2 fixation towards the formation of EPS precursors.

Taking these results as a whole, including sequence similarities, genetic organization, and (in several instances) experimental validation, it appears that *A. ferrooxidans* has the genetic potential to encode the EPS precursors UDP-glucose, UDP-galactose, and dTDP-rhamnose using glucose-1-phosphate as a central starting point (Fig. 4). Glucose-1-phosphate serves as a hub of carbohydrate metabolism, channeling sugars to the formation of extracellular polysaccharides (EPS) and glycogen and connecting these pathways with glycolysis, gluconeogenesis, perhaps the pentose phosphate pathway, and ultimately CO_2 fixation (Fig. 4). Future work will be directed towards an examination of the mechanisms involved in the regulation of the genes involved in glucose-1-phosphate formation in *A. ferrooxidans* and the pathways involved in the formation of EPS from the EPS precursors suggested in this study. It is anticipated that this information will provide a working model for understanding the formation of biofilms by *A. ferrooxidans* and help in understanding the role of this microorganism in mineral leaching and the formation of acid mine drainage.

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