Production of Recombinant and Tagged Proteins in the Hyperthermophilic Archaeon *Sulfolobus solfataricus*


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Many systems are available for the production of recombinant proteins in bacterial and eukaryotic model organisms, which allow us to study proteins in their native hosts and to identify protein-protein interaction partners. In contrast, only a few transformation systems have been developed for archaea, and no system for high-level gene expression existed for hyperthermophilic organisms. Recently, a virus-based shuttle vector with a reporter gene was developed for the crenarchaeote *Sulfolobus solfataricus*, a model organism of hyperthermophilic archaea that grows optimally at 80°C (M. Jonuscheit, E. Martusewitsch, K. M. Stedman, and C. Schleper, Mol. Microbiol. 48:1241–1252, 2003). Here we have refined this system for high-level gene expression in *S. solfataricus* with the help of two different promoters, the heat-inducible promoter of the major chaperonin, thermophilic factor 55, and the arabinose-inducible promoter of the arabinose-binding protein AraS. Functional expression of heterologous and homologous genes was demonstrated, including production of the cytoplasmic sulfur oxygenase reductase from *Acidianus ambivalens*, an Fe-S protein of the ABC class from *S. solfataricus*, and two membrane-associated ATPases potentially involved in the secretion of proteins. Single-step purification of the proteins was obtained via fused His or Strep tags. To our knowledge, these are the first examples of the application of an expression vector system to produce large amounts of recombinant and also tagged proteins in a hyperthermophilic archaean.

The homologous and heterologous expression of genes is a prerequisite for most biochemical studies of protein function. A vast variety of systems have been developed for protein production in members of the *Bacteria* and *Eukarya*, using numerous combinations of vector and promoter systems. Members of the *Archaea*, the third domain of life, are much less amenable to genetic manipulation. Transformation tools for the production of recombinant proteins exist for only a few species (for a review, see reference 2). A shuttle vector has been described for the expression of bacterial and methanococcal genes in *Methanococcus maripaludis* (12). Heterologous expression of bacterial and eukaryotic genes as well as of homologous genes has been achieved in the genetically most accessible archaea, the mesophilic, salt-dependent *Halobacterium* spp. (14, 17, 33). However, no such system has existed so far for thermophilic or hyperthermophilic archaea. Mesophilic hosts, in particular *Escherichia coli*, have been used to produce thermostable proteins for biochemical characterization and crystallographic studies (e.g., see references 22, 23, 29, and 34). However, a considerable number of proteins of hyperthermophiles fold into their native state only under natural conditions of high temperature or in the presence of their native cofactors. Furthermore, the production of recombinant and tagged proteins in native thermophilic hosts allows the identification of associated factors or even larger protein complexes.

The crenarchaeote *Sulfolobus solfataricus* has developed into an important model organism for molecular and biochemical studies of hyperthermophilic archaea. It grows optimally at 80°C and pH 3 under aerobic and heterotrophic conditions. Since many studies on the transcription, translation, and replication of this extremophile have been performed in vitro (4, 5, 10, 40), it is highly desirable to develop genetic tools for in vivo studies and for high-level production of proteins in this organism. Initial transformation systems and selectable markers have been established for *Sulfolobus solfataricus* in some laboratories (3, 6, 7, 9, 16, 41). We have recently developed a reporter gene system based on the virus SSV1 as well as the selectable marker genes *pyrEF* for the complementation of uracil auxotrophic mutants. The latter genes allow stabilization of the propagation of the vector by growing transformants under selective conditions (16). Moreover, strong, heat-inducible expression of the reporter gene *lacS*, coding for β-galactosidase, was demonstrated when it was placed under the control of the promoter of the major heat shock chaperonin gene *tf55α* (16).

In this study, we have used and improved this vector system for the heterologous and homologous expression of genes in *S. solfataricus*. We have constructed a set of entry vectors and introduced a transcriptional terminator and a second inducible promoter. Our system allows for the high-level production of functional and tagged cytoplasmic and membrane-associated proteins.
TABLE 1. Plasmids used for this study

<table>
<thead>
<tr>
<th>Plasmid</th>
<th>Description</th>
<th>Purpose of construction</th>
<th>Reference or source</th>
</tr>
</thead>
<tbody>
<tr>
<td>pMJ02</td>
<td>SSV1 in pUC18 with lacS under control of tf55alpha promoter</td>
<td>Promoter studies in Sulfolobus (heat shock induction)</td>
<td>16</td>
</tr>
<tr>
<td>pMJ03</td>
<td>pyrEF cassette inserted into pMJ02</td>
<td>Selectable marker added for stability</td>
<td>16</td>
</tr>
<tr>
<td>pMJ05</td>
<td>pyrEF cassette, tf55alpha promoter, lacS from pBR05 inserted into pMJ02, replacing tf55alpha and lacS from pMJ02</td>
<td>Destination vector for E. coli</td>
<td>This study</td>
</tr>
<tr>
<td>pMJ11</td>
<td>Promoter studies in Sulfolobus (heat shock induction)</td>
<td>S. Albers and M. Jonuschkeit, unpublished</td>
<td></td>
</tr>
<tr>
<td>pBR-05</td>
<td>pBR322 containing pyrEF cassette and lacS under control of tf55alpha promoter, with BssHIII site introduced (Fig. 1)</td>
<td>E. coli entry vector with heat shock promoter</td>
<td>This study</td>
</tr>
<tr>
<td>pASK5</td>
<td>pASK75 containing sor gene fused to C-terminal Strep tag</td>
<td>PCR template for amplification of Streptagged sor</td>
<td>37</td>
</tr>
<tr>
<td>pMJ05-sor</td>
<td>pyrEF cassette, tf55alpha promoter, sor from pBR05-sor inserted into pMJ02, replacing tf55alpha and lacS from pMJ02</td>
<td>sor expression in Sulfolobus</td>
<td>This study</td>
</tr>
<tr>
<td>pSVA1</td>
<td>SSO2316-6his inserted into pSVA1 (Fig. 1)</td>
<td>E. coli entry vector with terminator sequence separated</td>
<td>This study</td>
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<tr>
<td>pSVA2</td>
<td>SSO2316 insertion into pSVA1 (Fig. 1)</td>
<td>E. coli donor vector for SSO2316</td>
<td>This study</td>
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<tr>
<td>pSVA5</td>
<td>pSVA1 containing the lacS gene under control of araS promoter (Fig. 1)</td>
<td>E. coli entry vector with ara promoter</td>
<td>This study</td>
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<td>pSVA6</td>
<td>tf55alpha promoter, SSO2316 from pSVA2 inserted into pMJ05, replacing tf55alpha and lacS from pMJ05</td>
<td>SSO2316 expression in Sulfolobus</td>
<td>This study</td>
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<tr>
<td>pSVA9</td>
<td>araS promoter, lacS from pSVA5 inserted into pMJ05, replacing tf55alpha promoter and lacS from pMJ05</td>
<td>Promoter studies in Sulfolobus (arabinose induction)</td>
<td>S. Albers, unpublished</td>
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<td>pSVA14</td>
<td>SSO2680-10his inserted into pSVA5</td>
<td>E. coli donor vector for SSO2680</td>
<td>This study</td>
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<td>pSVA15</td>
<td>SSO2680-10his from pSVA14 inserted into pMJ05, replacing tf55alpha and lacS from pMJ05</td>
<td>SSO2680 expression in Sulfolobus</td>
<td>This study</td>
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<td>pSVA30</td>
<td>SSO287-Strep-10his inserted into pSVA5</td>
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<td>This study</td>
</tr>
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<td>pSVA31</td>
<td>araS promoter, SSO287-Strep-10his from pSVA30 inserted into pMJ05, replacing tf55alpha and lacS from pMJ05</td>
<td>SSO287 expression in Sulfolobus</td>
<td>This study</td>
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proteins in *S. solfataricus*, enabling biochemical characterizations and in vivo studies of protein function.

**MATERIALS AND METHODS**

**Strains and culture conditions.** An *S. solfataricus* pyrEF mutant (PH1-16) (25) was grown at 80°C and pH 3 in Brock’s medium with or without 10 μg ml⁻¹ uracil and with 0.1% tryptone and 0.2% arabinose or as indicated. The optical densities of liquid cultures were monitored at 600 nm (OD₆₀₀).

**Plasmid construction.** The plasmids used for this study are listed in Table 1. For the construction of pBR05, the genes pyrEF and lacS as well as the promoter region of the TSSαa gene were amplified by using the primers pyrEF-f-AvrII (CGGAAATACCCCTAGGGAATAAGTC), pyrEF-r-Nhel (GTGGTGCTAGCCCTCCTGTTAG), pRS5-f-AvrII (GCCATATCCCCTAGGGTGATTTCG), pRS5-r-BssHII (GACTGGCGCGCCCATACCTCA), S-r-EagI, thus generating a fragment of 3.6 kbp. The PCR product was used as template DNAAs in a second PCR using the flanking primers pyrEF-f-AvrII and lacS-r-EagI. From complementary overlaps of the plasmids, all PCR products were used as template DNAs in a second PCR using the flanking primers pyrEF-f-AvrII and lacS-r-EagI, thus generating a fragment of 3.6 kbp. The PCR product was purified by using a QIAgene Gene-Clean kit, restricted with Nhel and EagI, and ligated into pBR322 restricted with Nhel and EagI. The Sulfolobus-E. coli shuttle vector pBR05 was constructed by ligating the Nhel/EagI fragment of pBR05 to the XbaI/EagI-restricted and dephosphorylated SSV1/pUC18 shuttle vector pMJ02 (16).

The sor gene of *Acidulos ambivalens*, including a 10-amino-acid-encoding Strep tag at the C terminus, was PCR amplified by using the primers Sor-Strep-BssHIII (CTAGATAACGCGCCGACAAAAATCGCCG) and Sor-C-Strep-EagI (CATGCCCGCCGAGCGGGCTTATACATGCATCTCAGCGCA). The construct pASK5 (37), containing the sor gene fused to a C-terminal Strep tag, was used as the DNA template. The PCR product was restricted with BssHIII and EagI and ligated into pBR05, which yielded vector pBR-sor. This plasmid was cleaved with Nhel and EagI, and the sor gene with the promoter was ligated to XbaI/EagI-restricted and dephosphorylated pBR05, yielding the Sulfolobus expression vector pMJ05-sor.

An additional restriction site (Apal) was introduced into pBR05 such that other genes could be introduced by leaving the terminator from lacS in the vector. This was done with a quick mutagenesis kit I (Stratagene). A PCR was performed with the primer ApaIF (GCCATTAAAGGCACTAAAGGCCCACT TTCCTAAGTCTC) and its complement ApaIR (containing an ApaI site), with pBR05 as the template, yielding pSVA1. This vector was used as a template to replace the lacS gene with SSO2316 (fut). The latter was amplified by PCR from the *S. solfataricus* genomic DNA using the primer pair 2316f (CCCCCGCG CGACGTAGTGTTATGAGGATATTAC) and 2316r (CCCCCGCG CCCTTAAAGTCTAGTATGAGGATATTAC), thereby fusing a C-terminal six-His tag to the encoded product. The PCR product was cloned into the BssHIII and Apal sites of pSVA1 to yield pSVA2. For the construction of pSVA5, the promoter region 241 bp upstream of the araS gene (SSO2316) was amplified from *S. solfataricus* genomic DNA with the primer pair araSF (GGGGGCCATGGACTCATTTCCAAATAAC) and araSR (CCCCCCGGG CGGGAATGTTTTAGCTTTTTTTATGAGGATATTAC), containing a Blaln and an NcoI restriction site, respectively. In the same manner, the lacS gene was amplified with the primer pair lasf (GGGGGCCATGGACTTTTCATTTTTATGAGGATATTAC) and lacSR (CCCCCGCGCCGCTTTTCTTTTAGTTACATGAGGATATTAC), thereby fusing a C-terminal six-His tag to the encoded product. The PCR product was digested with NcoI and Eagl, and the resulting ligation product was digested...
with BlnI and Apal and inserted into pSVA1 to yield pSVAS, which contained the lacS gene under the control of the araS promoter.

SSO2680 and SSO0287 were amplified from genomic DNA by PCR using the primer pairs SSO2680f/SSO2680r and SSO0287f/SSO0287r, respectively. This resulted in the introduction of Neo and Apal restriction sites in the flanking regions of these genes and the presence of a C-terminal 10-His tag in SSO2680 and a StreptII tag and 8-His tag at the C terminus of SSO0287. Both genes were cloned into pSVAS using the Neo/Apal sites, yielding pSVAS1 and pSVAS30, which carried SSO2680 and SSO0287, respectively. To transfer the araS promoter together with the gene to be expressed into the virus-based vector, the BlnI/EagI inserts from pSVA14 and pSVA30 were ligated into pMJ05, resulting in plasmids pSVAS1 and pSVAS31, respectively.

**Transformation of *S. solfataricus***

Electroporation of *S. solfataricus* PH1-16 and the isolation of single transformants were done as described previously (16, 31). Integration of the viral vector into the genome was confirmed by Southern analysis using standard procedures.

**Expression and purification of Sor.** Cells containing the pMJ05-aro construct were grown in 300 ml Brock’s medium supplemented with 0.1% tryptone and 0.2% d-arabinose. After 3 days of cultivation, cells of strain (1 g) were harvested by centrifugation and resuspended in 25 ml 10 mM Tris-HCl, pH 8.0. The cells were lysed by sonication, and after centrifugation to remove cell debris and the membrane fraction, the supernatant was used for Streptactin affinity chromatography. Twenty milliliters of supernatant was applied to a Streptactin Superflow column (10 ml; IBA, Göttingen, Germany) and subsequently with a Speed-Vac machine.

**Expression of FlaI.** Cells containing pSVA6 were inoculated into 50 ml Brock’s medium supplemented with 0.1% tryptone and 0.2% d-arabinose. At an OD600 of 0.5, the cells were diluted into 400 ml of the same medium and grown to an OD600 of 0.8. Cultures were then shifted to 88°C for 24 to 48 h, and the cells were harvested by centrifugation and resuspended in 10 ml Tris-HCl, pH 8, and 100 mM NaCl. Lysis of the cells and the isolation of membranes were done as described below.

**Expression and purification of Ssr.** Cells containing pSVA15 were inoculated into 50 ml Brock’s medium supplemented with 0.1% tryptone and 0.2% d-arabinose. At an OD600 of 0.5, 10 ml of cells was transferred to 400 ml medium containing 0.1% tryptone and 0.2% d-arabinose to induce the expression of SSO2680. After 2 days of growth (OD600 ~0.5), 10 ml of cells was transferred to 400 ml medium containing 0.1% tryptone and 0.2% d-arabinose to induce the expression of SSO2680. After 2 days of growth (OD600 ~0.8), the cells were harvested and resuspended in buffer A (50 mM NaPi, pH 8, 100 mM NaCl, and 5 mM imidazole). Lysis of the cells and the isolation of membranes were done as described below. Membranes were resuspended in buffer A and solubilized with 2% n-dodecyl-β-maltopyranoside (DDM) at a protein concentration of 4 mg/ml for 45 min at room temperature. Nonsolubilized protein was removed by high-speed centrifugation, and the supernatant containing the solubilized membrane protein was applied to a His-Select 1-ml column (Sigma) to a tetramer of 240 kDa in vivo (28), was demonstrated by the presence of the start codon of *S. solfataricus* lacS. This promoter has a strong basal activity and is additionally inducible by shifting cultures from 78 to 88°C. In order to improve the pMJ03-based vector system for the homologous and heterologous expression of genes in *S. solfataricus*, entry vectors were constructed with restriction sites that allowed us to replace the lacS gene with other genes of interest by cloning them directly behind the promoter sequence. The putative transcriptional terminator sequences of lacS were left intact (Fig. 1). Moreover, the heat-inducible tf55α promoter was replaced with an arabinose-inducible promoter in order to allow the induction of high-level expression without imposing stress on the host cells. For this purpose, a 241-bp region upstream of the open reading frame of araS, encoding the arabinose-binding protein, a subunit of an ABC sugar transporter in *S. solfataricus* (11), was selected for cloning. In contrast to the original tf55α-lacS promoter-gene fusion in pMJ03, this construct did not contain the five initial codons of the tf55α gene. Instead, the start codon of araS coincided with the start codon of the gene to be expressed (Fig. 1). High-level expression of β-galactosidase, which forms a tetramer of 240 kDa in vivo (28), was demonstrated by the separation of cell extracts of transformants in a non-denaturing blue native protein gel and subsequent activity staining with ONPG (24).

**RESULTS**

**Entry vectors and introduction of the sugar-inducible araS promoter.** In the recently described shuttle vector pMJ03 (16), the complete DNA of the virus SSV1 (15 kb) was combined with pUC18 for replication/selection in *Escherichia coli*, the genes pyrEF of *S. solfataricus* for complementation of uracil auxotrophic mutants of *Sulfolobus*, and the reporter gene lacS, coding for β-galactosidase. The system was used to show the expression of the β-galactosidase gene (lacS) under the control of the heat-inducible promoter of the α subunit of the thermodomosome of *S. solfataricus* (TF55α). This promoter has a strong basal activity and is additionally inducible by shifting cultures from 78 to 88°C. In order to improve the pMJ03-based vector system for the homologous and heterologous expression of genes in *S. solfataricus*, entry vectors were constructed with restriction sites that allowed us to replace the lacS gene with other genes of interest by cloning them directly behind the promoter sequence. The putative transcriptional terminator sequences of lacS were left intact (Fig. 1). Moreover, the heat-inducible tf55α promoter was replaced with an arabinose-inducible promoter in order to allow the induction of high-level expression without imposing stress on the host cells. For this purpose, a 241-bp region upstream of the open reading frame of araS, encoding the arabinose-binding protein, a subunit of an ABC sugar transporter in *S. solfataricus* (11), was selected for cloning. In contrast to the original tf55α-lacS promoter-gene fusion in pMJ03, this construct did not contain the five initial codons of the tf55α gene. Instead, the start codon of araS coincided with the start codon of the gene to be expressed (Fig. 1). High-level expression of β-galactosidase, which forms a tetramer of 240 kDa in vivo (28), was demonstrated by the separation of cell extracts of transformants in a non-denaturing blue native protein gel and subsequent activity staining with ONPG (24). A quantitative estimation of promoter strengths using the LacS reporter activity showed that the arabinose promoter exhibited a considerably lower basal activ-
ity than the tf55α promoter in the absence of an inducer (Table 2). Upon the addition of arabinose to the medium, the activity levels of β-galactosidase increased about 13-fold. The absolute value of 3.3 U/mg protein was comparable to that obtained with the tf55α promoter after heat shock (Table 2, second column). However, transformants with a tf55α promoter that contained an Ncol site at the translation start site (comparable to the situation in the araS construct) exhibited an even higher activity, at ca. 12 U/per mg of protein, in crude extracts (Table 2, construct pMJ11). A more detailed analysis of promoter variants and the effects of mutations on the initiation of transcription and translation is currently in progress.

**Heterologous expression and purification of the sulfur oxygenase reductase from Acidianus ambivalens.** The sulfur oxygenase reductase (Sor) from the chemolithoautotrophic crenarchaeote *A. ambivalens* is a cytoplasmic enzyme that catalyzes the initial step in the dissimilatory sulfur oxidation pathway in this organism (19, 20). Sulfite, thiosulfate, and hydrogen sulfide are simultaneously produced from elemental sulfur in an oxygen-dependent reaction. Sor is an icosatetrameric protein with a molecular mass of 840 kDa that is composed of identical subunits of 35.2 kDa (38) and contains a mononuclear non-heme iron center as a cofactor (37). *S. solfataricus* lacks a sor homolog (32).

The sor gene, including codons for a C-terminally fused Strep tag, was cloned under the control of the tf55α promoter. Single transformants of *S. solfataricus* PH1-16 containing the pMJ05-sor construct were grown at 78°C and subsequently shifted to 88°C to induce the expression of the sor gene. The yield was 0.5 mg of enriched Sor protein per liter of culture after Streptactin affinity chromatography. A band corresponding to a molecular mass of 36 kDa was visible on SDS gels.
containing the purified protein fraction (Fig. 3A). The protein was confirmed to be identical to the *A. ambivalens* Sor protein plus the additional amino acids from the Strept tag by MALDI-TOF analysis. Other bands were visible besides the Sor band. A prominent band with an apparent molecular mass of 55 kDa was identified by MALDI-TOF analysis as the large subunit of the biotinylated *S. solfataricus* acetyl-coenzyme A (acetyl-CoA)/propionyl-CoA carboxylase (also called AccC or SSO2466 [15]). The specific activities in the Sor preparation, measured as the production of thiosulfate and hydrogen sulfide from elemental sulfur, were 4.5 and 1.2 U mg protein \(^{-1}\) (oxygenase and reductase activities, respectively). These values are comparable to those obtained for the enzyme isolated from its native host (Fig. 3B), indicating the successful assembly of the homomultimeric enzyme and the incorporation of the iron cofactor.

**Homologous expression of the ATPase FlaI.** *flaI* (SSO2316) is part of the potential flagellin operon of *S. solfataricus* and encodes a 59-kDa protein with a nucleotide-binding domain. FlaI is homologous to other ATPases present in archaeal flagellin operons, which may fulfill a role in the assembly of the archaean flagellum. Gene inactivation experiments with *Halobacterium salinarium* and *Methanococcus voltae* have previously shown that the FlaI protein is essential for flagellum formation (27, 36). In *S. solfataricus*, *flaI* is expressed only in the stationary growth phase (1). The protein was expressed in *E. coli* earlier, purified to homogeneity, and shown to exhibit divalent cation-dependent ATP-hydrolyzing activity (1). In order to produce FlaI in *Sulfolobus*, the gene, including codons for a six-His tag, was cloned under the control of the tf55 promoter to yield pSVA6. Single transformants containing the vector were isolated and grown at 78°C with a subsequent shift to 88°C. Strain PH1-16 (*pyrEF lacS* double mutant) was grown as a control. Samples were taken from both cultures at the start of heat incubation and after 1 and 2 days. Cells were harvested, separated into membrane and cytoplasmic fractions, and analyzed by SDS-polyacrylamide gel electrophoresis (SDS-PAGE) and immunoblotting using antibodies directed against purified FlaI (1). While the heterologously expressed FlaI protein in *E. coli* had been isolated from the soluble cytoplasmic fraction

<table>
<thead>
<tr>
<th>Construct</th>
<th>Promoter</th>
<th>Beta-galactosidase sp act (U/mg protein) (^a)</th>
<th>Fold induction</th>
<th>Inducing agent</th>
</tr>
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<tbody>
<tr>
<td>pMJ03</td>
<td>tf55 alpha</td>
<td>1.46 (0.13)</td>
<td>5.09 (0.16)</td>
<td>3.50 (0.33)</td>
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<tr>
<td>pMJ11(^b)</td>
<td>tf55 alpha (NcoI)(^c)</td>
<td>1.18 (0.13)</td>
<td>11.83 (0.70)</td>
<td>10.02 (0.67)</td>
</tr>
<tr>
<td>pSVA9</td>
<td>araS (NcoI)(^c)</td>
<td>0.28 (0.09)</td>
<td>3.31 (0.51)</td>
<td>12.97 (3.84)</td>
</tr>
</tbody>
</table>

\(^a\) Data from 5 (pMJ03), 3 (pMJ11), and 10 (pSVA9) independent experiments (numbers in parentheses indicate standard deviations).

\(^b\) Unpublished construct.

\(^c\) NcoI site introduced at the start codon of *lacS*.

**FIG. 3.** Purification and activity of *A. ambivalens* Sor isolated from *S. solfataricus* transformed with pMJ05-sor. (A) SDS-PAGE of different fractions (E1 and E2) eluted from the Streptactin column. M, molecular mass marker proteins. Solid arrow, Sor band; dashed arrow, AccC subunit of the acetyl/propionyl-CoA carboxylase (both identified by MALDI-TOF analysis). (B) Sulfur oxygenase and reductase activities of pooled eluate fractions showing time-dependent increases in the amounts of thiosulfate and hydrogen sulfide in the assay mixture (19). Background, nonenzymatic production of thiosulfate and hydrogen sulfide from sulfur disproportionation under the same assay conditions without the enzyme (19).
(1), the protein produced in *Sulfolobus* could only be detected in the membrane fraction of induced cells (Fig. 4). This observation was confirmed using antibodies directed against the C-terminal His tag of FlaI (data not shown).

**Homologous expression and purification of SSO2680.**

SSO2680 is a hydrophilic nucleotide-binding protein of 59 kDa. It has been shown to be encoded in an operon with five other genes (1) that encode proteins with motifs typically found in subunits of bacterial protein secretion systems. These so-called type II and type IV secretion machineries, as well as type IV pilin assembly systems, are involved in the assembly or secretion of multimeric substrate proteins such as pili. The protein substrate of this putative secretion operon in *Sulfolobus* is unknown. Interestingly, the operon contains a gene for one small protein of 15 kDa, SSO2681, which contains a type IV pilin signal peptide and might therefore be a subunit of a pilus structure (1). SSO2680 shows sequence similarity to bacterial secretion ATPases, such as Virb11 and PilT, which power the secretion process (30, 42). SSO2680 was cloned behind the *araS* promoter with codons for a C-terminal 10-His tag and transferred to the viral vector, yielding pSVA15. After transformation of *Sulfolobus*, single transformants were grown on 0.1% tryptone and subsequently transferred to the same medium supplemented with 0.2% arabinose. After 2 days of growth, cells were harvested, lysed, and separated into membrane and cytoplasmic fractions. These fractions were analyzed by SDS-PAGE and immunoblotting using antibodies directed against the His tag (Fig. 5). Although SSO2680 is predicted to be a soluble protein, it was recovered with the membrane fraction. In *E. coli*, the protein had been recovered with inclusion bodies, and only a small fraction could be isolated from the soluble fraction (1). SSO2680 was purified from detergent-solubilized membranes of *S. solfataricus* by His-Select Ni affinity chromatography, yielding ~1 mg purified enzyme per liter of cells (Fig. 5A). Immunodetection with His tag-specific antibodies confirmed that the purified protein corresponded to the recombinant tagged protein (lower part of Fig. 5A). SSO2680 showed divalent cation-dependent ATPase activity at a high temperature and a preference for Mn^2+ over Mg^2+ (Fig. 5B), in agreement with previous studies using the heterologously expressed protein purified from *E. coli* (1). Interestingly, the specific activity of the protein produced and purified from *S. solfataricus* was six times higher (50 nmol of P_i released mg protein^-1 min^-1) than that of the enzyme isolated from *E. coli* (8 nmol of P_i released mg protein^-1 min^-1) (1), which indicated a more native conformation of the homologously expressed protein. In order to analyze the effect of medium composition on the expression level, cells were grown with different amounts of tryptone and arabinose, and the membrane and cytoplasmic fractions of lysed cells were analyzed by SDS-PAGE and immunoblotting. The relative yield of recombinant protein did not change when the arabinose concentra-

![FIG. 4. Western analysis of FlaI expression. *S. solfataricus* cells transformed with pSVA6 and control PH1-16 cells were grown for 2 days at 88°C. Samples were taken at the beginning of the temperature shift from 78 to 88°C and after 1 and 2 days of growth at 88°C. Membranes were isolated from the cells and were analyzed by SDS-PAGE and immunoblotting using FlaI-specific antibodies. The arrow indicates the detected expression product.](image-url)

![FIG. 5. Overexpression, purification, and activity of SSO2680. (A) Coomassie blue-stained SDS-PAGE gel of His tag-specific affinity chromatography fractions from solubilized membranes derived from *S. solfataricus* cells transformed with pSVA15. In the lower panel, the corresponding Western blot of the same samples, using His tag-specific antibodies, is shown. M, molecular mass marker; St, starting material; FT, flowthrough; W, wash; E, elution fraction. (B) ATPase activity of purified SSO2680 in the presence of EDTA, Mg^2+, or Mn^2+. (C) Overexpression of SSO2680. Membranes of a single pSVA15 transformant were separated in a Coomassie blue-stained SDS-PAGE gel before (−) and after (+) induction with 0.4% arabinose. The arrow indicates SSO2680. In the lower panel, the corresponding Western blot of the same samples, using His tag-specific antibodies, is shown.](image-url)
tion was varied between 0.2 and 0.4%. Although the amount of tryptone affected the growth rate, it did not affect the expression level (data not shown). We occasionally observed very high expression levels, comparable to those obtained with *E. coli* expression systems, in different transformants of *Sulfolobus*. One such example is shown for SSO2680 in Fig. 5C. The conditions which led to these increased expression levels are currently under investigation.

**Expression of SSO0287.** SSO0287 (or ABCE1) is a cytoplasmic protein of 68 kDa that contains two nucleotide-binding domains and is predicted to harbor two iron-sulfur clusters. It is homologous to eukaryotic ABC proteins that are essential for cell function and involved in ribosome biogenesis and protein translation (18, 43) but whose specific function is unknown.

The gene was cloned behind the *araS* promoter with codons for a C-terminal tandem tag, including a Strep tag followed by an eight-His tag. Single transformants were grown on 0.1% tryptone and subsequently transferred to medium containing both 0.1% tryptone and 0.2% arabinose. After 2 days, cells were harvested and lysed. The cytosolic fraction was applied to either a Streptactin or His-Select Ni affinity column. Both methods resulted in single-step purification of the tagged protein (Fig. 6). The identity of the protein was verified by means of immunoblotting using antibodies directed against the Strep tag and the His tag and polyclonal antibodies directed against ABCE1 itself. The purified protein exhibited divalent cation-dependent ATPase activity, with a high temperature optimum of 85°C, close to the optimal growth temperature of *S. solfataricus* (S. Dinkelaker and R. Tampe, manuscript submitted).

As described above, SSO0287 contains two putative sites for the binding of iron-sulfur clusters. Previous attempts to obtain correct assembly of these clusters by heterologous expression in *E. coli* failed (unpublished results), whereas the homologically produced protein in *S. solfataricus* appeared to contain the iron-sulfur clusters. The protein was subjected to gel filtration chromatography, and elution was monitored at two different wavelengths, 280 and 410 nm (Fig. 6B), to detect the protein and the iron-sulfur clusters, respectively. The isolated protein eluted as a single monodisperse symmetric peak that was devoid of aggregates. The elution profiles recorded at 280 and 410 nm coincided, suggesting that the recombinant SSO0287 protein contained correctly assembled iron-sulfur clusters.

The expression of SSO0287 in *S. solfataricus* was scaled up from 400-ml cultures to fermentation in 45 liters. A fermentor containing medium with 0.1% tryptone and 0.2% arabinose was inoculated with 1.6 liters of SSO0287-expressing *S. solfataricus* culture. After 2 days of growth, the cells were collected at an OD_{600} of ~0.8, and the protein was purified. The yields per liter (0.75 to 1 mg) were comparable to those from small cultures, but when the cells were grown to higher densities (ODs of about 1.5), degradation products of the purified protein were observed.

**DISCUSSION**

We report here the construction of a virus-based shuttle vector for the heterologous and homologous expression of genes in the hyperthermophilic archaean *S. solfataricus*. A summary of the expression studies is shown in Table 3. Two promoters were used to drive expression, namely, the heat-inducible promoter of the alpha subunit of the chaperonin TF55 (for Sor, FlaI, and β-galactosidase) and an arabinose-inducible promoter (for SSO2680, SSO0287, and β-galactosidase). Since the TF55α promoter requires a shift of cultures to 88°C, which is close to the maximal growth temperature of *S. solfataricus*, this procedure might impose highly stressful conditions on the cells. For routine expression purposes, we therefore favor the use of the arabinose promoter, which is more easily induced and results in less stressful conditions. Similarly, the pBAD system is often used in *E. coli*, which is based on the arabBAD operon controlling the arabinose metabolic pathway (13). This is a tightly controlled system that only allows for expression in the presence of arabinose, thereby preventing adverse effects on growth during the stages that are needed to obtain cell mass. In *S. solfataricus*, several sugar binding proteins that are part of ABC transporters are expressed only in the presence of their respective substrates (11). One of these genes is *araS*, which encodes an arabinose-binding protein. The expression of *araS* is induced only when *S. solfataricus* is grown on arabinose (11). We have therefore chosen the promoter region of *araS* to drive the expression of genes in *S. solfataricus*. A thorough characterization of the regulatory elements in this promoter and the construction of minimal in-
**TABLE 3. Summary of expression studies**

<table>
<thead>
<tr>
<th>Protein</th>
<th>Activity</th>
<th>Clearance beta-galactosidase</th>
<th>Beta-galactosidase</th>
<th>Construct</th>
<th>Tag*</th>
<th>Tag</th>
<th>Promoter</th>
</tr>
</thead>
<tbody>
<tr>
<td>S. solfataricus</td>
<td>ATPase</td>
<td>Naive stain</td>
<td>Strep</td>
<td>pMV6</td>
<td>10-His</td>
<td>pmM5/LYSV5</td>
<td>Strep plus eight-His</td>
</tr>
<tr>
<td>S. solfataricus</td>
<td>ATPase</td>
<td>Native stain</td>
<td>Strept + eight-His</td>
<td>pMV6</td>
<td>10-His</td>
<td>pmM5/LYSV5</td>
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<td>pmM5/LYSV5</td>
<td>Strept plus eight-His</td>
</tr>
</tbody>
</table>

*All tags were positioned at the C terminus.

All proteins produced in this study (except β-galactosidase) were fused either with a His tag, a Strep tag, or both. The data show that *S. solfataricus* tolerates these tags and that they can be used for protein detection and purification. In particular, the use of His-Select material for His tag-specific affinity chromatography resulted in single-step purification to 99% homogeneity, indicating that there is hardly any background of endogenous proteins in *S. solfataricus* that may bind to the affinity matrix. In contrast, the purification of proteins from *Sulfolobus* via a Streptactin column result in coelution of the biotinylated acetyl-CoA/propionyl-CoA carboxylase. This enzyme is involved in the modified 3-hydroxypropionate cycle used for autotrophic CO₂ fixation in crenarchaeota (15). The function of this enzyme in *S. solfataricus* is not clear since all of our experiments were conducted under heterotrophic conditions. Copurification of the carboxylase was observed only when low, and not high, yields of the recombinant protein were obtained (Fig. 3, Sor, versus Fig. 6, SSO0287).

The proteins produced in this study will be used for biochemical and physiological studies. For example, transformants that express the sor gene from *Acidimius ambivalens* will be analyzed for the ability to metabolize sulfur. *S. solfataricus* lacks the sor gene but contains other genes that might encode proteins involved in the metabolism of inorganic sulfur compounds, e.g., a doxA operon encoding a thiosulfate:quinone oxidoreductase, an enzyme which acts downstream in the sulfur oxidation pathway in *A. ambivalens* (26). Therefore, it will be interesting to analyze the sulfur-oxidizing capabilities of various *S. solfataricus* transformants carrying the sor gene. In the case of SSO2680, we will attempt to identify interacting partners by coimmunoprecipitation. Interestingly, this protein exhibited a much higher activity than the heterologously produced protein from *E. coli*, where it was mainly recovered in inclusion bodies, pointing to a folding defect in the bacterial host (1). The homologously expressed protein was solely localized in the membranes of *S. solfataricus* cells. Since the protein is predicted to be a cytoplasmic ATPase subunit of a membrane-bound protein secretion system, this result might indicate that the protein is associated with native lipids or membrane components that are part of the secretion machinery. Similarly, tagged FlaI can be used for the identification of binding partners. The proteins which play an important role in flagellation in archaea are known from transcription and deletion studies (35), but their subunit interactions during assembly of the flagellum are not known. The flaI-expressing recombinant promoters for this expression system are under way (M. Jonuscheit, S. V. Albers, et al., unpublished data).
binant Sulfolobus strain can serve as an important tool to address such questions.

Also, in the case of SSO0287, the tagged version can be used for searches for unidentified interacting proteins. The exact function of this protein is unknown, but it contains a unique ABC domain with two iron-sulfur clusters. Strikingly, it appeared that only 50% of these iron-sulfur clusters assembled when the protein was expressed in E. coli, whereas in S. solfataricus the protein mostly likely followed the endogenous assembly pathway, which ensures the correct incorporation of these iron-sulfur clusters into the protein. The homologous expression of SSO0287 now allows for mutational analysis in S. solfataricus in order to assess its in vivo function using interference assays.

In conclusion, we have established a vector system for protein expression in S. solfataricus, making use of two different promoters. The versatility of the system was demonstrated by various examples of heterologous or homologous proteins with a cytoplasmic or membrane-associated association, and all recombinant proteins were shown to be functional (except for FlaI). This is the first report of the application of an expression system in a hyperthermophilic archaeon. The expression of recombinant proteins were shown to be functional (except for FlaI). This is the first report of the application of an expression system in a hyperthermophilic archaeon. The expression of recombinant proteins were shown to be functional (except for FlaI). This is the first report of the application of an expression system in a hyperthermophilic archaeon.

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