Type IV prepilin peptidases (TFPPs) are prokaryotic signal peptidases that are required for the correct secretion of affiliated substrates, which are termed type IV prepilins and type IV prepilin-like proteins (42). Several TFPPs from gram-negative and gram-positive bacteria have been previously characterized, and recently, archaeal homologues have also been described (6, 8). All TFPPs have in common that they recognize an unusual membrane-targeting sequence termed the type IV pilin signal peptide. It is located at the amino terminus of a precursor protein and consists of a short positively charged leader followed by a hydrophobic stretch of about 20 amino acids, with a conserved cleavage site located between these two domains. The extreme N terminus of membrane-inserted prepilins has been shown to face the cytoplasm (11, 41). Accordingly, the active site of the processing enzyme is expected to be on the cytoplasmic side of the membrane. After cleavage by the TFPP, the hydrophobic domain remains part of the mature protein. Additionally, most bacterial TFPPs transfer a methyl group to the new amino terminus, whereas the archaeal enzymes seem to possess only cleavage activity (4; unpublished results). Indeed, the peptidase and methyl transferase activities of PilD from *Pseudomonas aeruginosa* are separable, which suggests two independent active sites (33, 40, 42). Processed (i.e., cleaved and methylated) type IV pilins are assembled into the type IV pilus, a long, retractable surface organelle involved in diverse processes, such as surface attachment, twitching motility, and DNA uptake (16, 25, 30). Additionally, type IV pilin-like proteins play an important role in the general (type II) secretion pathway of gram-negative bacteria by forming subunits of the type II pseudopilus (12, 13, 49). In archaea, type IV pilin-like signal peptides were first identified in flagellins, the structural subunits of flagellar filament (15, 46). The

The hyperthermophilic acidophile *Sulfolobus solfataricus*, however, was shown to secrete a more diverse set of proteins with type IV pilin-like signal sequences (3, 6), including flagellin. In this organism, a cleaved type IV pilin-like signal peptide was identified in three membrane-associated sugar-binding proteins which mediate solute uptake by ABC transporters (4, 14). Further analysis of the genome revealed that 12 additional proteins could be equipped with this signal peptide, including 4 putative solute-binding proteins and small proteins of unknown function (3, 6).

Recently, we identified PilD, the *S. solfataricus* enzyme responsible for cleavage of flagellin (FlaB) and glucose-binding protein (GlcS) precursors (pre-FlaB and pre-GlcS, respectively) (6). PilD was discovered on the basis of a phylogenetic relationship with bacterial TFPPs described in the COG database (44). This relationship could not be detected by BLAST searches. Indeed, a multiple linear alignment of whole sequences shows no significant homology between bacterial TFPPs and their archaeal counterparts. Therefore, it is important to determine whether the enzymes from these two prokaryotic domains have similar catalytic mechanisms, which would support the suggested evolutionary relationship. The proteolytic mechanism of TFPPs from *Pseudomonas aeruginosa* and *Vibrio cholerae* has been studied by use of site-directed mutagenesis of conserved residues in the corresponding enzymes. For PilD from *P. aeruginosa*, cysteine residues were shown to be important for the processing of prepilin (40). However, the related enzyme from *Xanthomonas campestris*, which does not contain cysteine residues, could complement a *P. aeruginosa* pilD mutant (19), suggesting that amino acids other than cysteine could be required for the cleavage reaction. Indeed, when either of two universally conserved aspartic acid residues was altered in the TFPP homologue TcpJ from *V. cholerae*, the enzyme showed no activity in vivo or in vitro (28). The same report shows that the domain containing the conserved cysteines is not required for cleavage activity. It is thus evident that TFPPs are aspartic acid proteases and, because the catalytic residues are conserved among bacteria, they should also be present in
archaeal enzymes, provided that the evolutionary connection is true. Recently, the preflagellin peptide FlaK from *Methanococcus voltae* was shown to contain two aspartic acid residues required for in vitro activity, and these residues could correspond to those identified in TcpJ (7). As these residues are found in the amino-terminal half of the protein, no function could be assigned to the carboxyl-terminal portion of the protein, which includes a large cytoplasmic loop.

Here we present further evidence for the relatedness of archaeal TFPs, exemplified by PibD from *S. solfatarius*, and bacterial enzymes. By mutational analysis, two conserved aspartic acid residues are identified as essential for activity. Additionally, a pair of aspartic acid residues in the large cytoplasmic loop of the enzyme is shown to be important for enzyme function. Finally, we suggest a membrane topology model for PibD based on cysteine accessibility studies.

**MATERIALS AND METHODS**

**Strains, growth conditions, and plasmids.** Escherichia coli strain DH5α was used for all cloning steps (17), and strains BL21(DE3) (43) and C43(DE3) (32) were used for the overproduction of protein. The two latter strains additionally carry the pACYC-RIL plasmid, which encodes additional tRNAs to compensate for rare *E. coli* codons (Stratagene, La Jolla, Calif.). Plasmid pZA1 encodes the *flaB* precursor protein, and pZA5 carries *pibD* both are under the control of the T7 promoter and contain a carboxyl-terminal six-histidine tag. pBAD/HisA is corresponding to two times the amount used in each cleavage reaction were serum albumin as a standard. Activity of wild-type and mutant variants of PibD was determined using a DC protein assay kit (Bio-Rad, Hercules, CA) with bovine serum albumin (wt/vol) glycerol. The protein concentrations of the membrane preparations were concentrated of about 20 mg/ml in 50 mM Tris-HCl, pH 7.5, containing 20% (wt/vol) sucrose. After the concentration had been increased to 1.5 mg/ml in 50 mM sodium phosphate, pH 8.0, 150 mM NaCl, 10% (wt/vol) glycerol, and the nonionic detergent dodecylmaltoside (DDM) (Anatrace, Maumee, OH) at 1% (wt/vol). Membranes were solubilized for 15 min at ambient temperature, whereupon the solubilized material was cleared by ultracentrifugation (MLA 80 rotor; 15 min, 20,500 g, 4°C). The supernatant was mixed with 10 mM imidazole, pH 8.0, and 40 μl HIS-select nickel affinity agarose resin (Sigma) equilibrated with wash buffer (50 mM sodium phosphate, pH 8.0, 150 mM NaCl, 10% glycerol, 20 mM imidazole, 0.1% DDM). The resin was incubated 16 h at 4°C with gentle shaking. Next, the resin was collected by centrifugation and washed twice with 1 ml wash buffer. Washed resin was incubated 5 min with 35 μl elution buffer (50 mM sodium phosphate, pH 8.0, 150 mM NaCl, 10% glycerol, 100 mM Na2-EDTA, 0.1% DDM) under gentle shaking and collected by centrifugation. Eluted protein was labeled with fluoresein maleimide (Molecular Probes, Leiden, The Netherlands) as follows. A fresh stock solution was prepared by diluting 50 mM fluorescein maleimide dissolved in dimethyl formamide 10-fold with denitized water. Fluorescein maleimide was added to elution fractions at a final concentration of 0.1 mM. Samples were incubated 5 min at 30°C in the dark, and the reaction was quenched by the addition of 1 mM DTT. For analysis, purified and labeled samples were mixed with SDS sample buffer and separated on a 12% SDS-PAGE gel according to the method of Laemmli (27). Fluorescent bands were detected using an FI Lumi-Imager (Roche Diagnostics, Almere, The Netherlands) with a 520-nm fluorescence filter. Subsequently, the same gel was fixed and stained with Coomassie brilliant blue R-250 (CBB), destained, and recorded with an FI Lumi-Imager in the transillumination mode. Densitometric analysis was carried out using the public-domain software ImageJ (developed by W.S. Rasband, U.S. National Institutes of Health, Bethesda, Maryland). Pixel densities of each fluorescent band were measured and corrected for background signal by use of an area of identical size in the same lane that did not contain protein. Bands in the CBB-stained gel and gel were quantitated in the same way. The intensity of each band (expressed as a ratio of fluorescent band signal and CBB band signal) of A405 versus mock-treated samples was calculated.

**Predictions of TMS.** The PibD primary sequence was submitted to the World Wide Web version of the PSI-Pred/MEMSAT2 (21, 22, 31), HMMTOP (47, 48), TMHMM2.0 (36) (26), and Phobius (23) programs; default settings were used. In the case of Phobius, predictions were also run in the constrained mode with the amino-terminal noncytoplasmic loop setting. However, this did not change the number of transmembrane segments (TMS) predicted by the program.

**RESULTS**

**Chemical inactivation of PibD.** To establish whether aspartic acids are involved in the reaction catalyzed by the type IV pilin-like signal peptidase PibD, the inhibitor 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDAC) in the presence of glycaminide was tested. EDAC is known to modify acidic amino acid residues (i.e., aspartic acid and glutamic acid) and has been used previously to inhibit aspartic acid proteases (18, 51), including TcpJ. *E. coli* inner membranes containing expressed PibD were treated with EDAC-glycaminide and subsequently tested for in vitro cleavage of precursors of flagellin (FlaB) and glucose-binding protein (GlcS)
Identification of catalytic residues in PibD by site-directed mutagenesis. To determine which residues are essential for catalysis, each of the nine aspartate residues was changed to alanine by site-directed mutagenesis. The adjacent residues D187 and D188 were replaced simultaneously. Mutant proteins were expressed in *E. coli*, and membranes containing the enzymes were tested for cleavage of pre-FlaB in vitro (Fig. 2) (6). Expression of the D207A mutant could not be detected by Western immunoblot analysis with antibodies directed against the carboxyl-terminal epitope tag of PibD (data not shown). However, a mutant with an asparagine (D207N) at this position was stable upon expression in *E. coli*. The amounts of PibD protein present in the membrane preparations were estimated by Western immunoblot analysis. Based on this estimation, comparable amounts of enzyme source were used in all experiments. The activity levels of five single mutants (D30A, D157A, D173A, D180A, D207N) were comparable to that of the wild-type enzyme (Fig. 2A). However, no cleavage of preflagellin was observed in D23A, D80A, and D187A/D188A mutants. Therefore, these amino acids represent potential candidates for catalytic residues of the enzyme. The aspartate residues at positions 23 and 80 are part of putative signature motifs that are also found around catalytic residues of other TFPPs and intramembrane aspartic acid proteases (28, 39, 52). Asp23 is embedded between hydrophobic residues, and Asp80 is preceded by an alanine and two glycines.
This resembles the characteristic GxGD motif found in TFPPs and in presenilin aspartic proteases (39). An aspartate-to-glutamate mutation in catalytic sites of aspartic peptidases might restore enzyme activity. However, a D23E mutant of PibD was also inactive, whereas a D80E mutant retained partial activity (Fig. 2B). In order to dissect the defect of the D187A/D188A mutant, the respective single mutants were generated. PibD containing the D187A or D188A mutation was active, although there was an apparent reduction in activity in the D187A mutant (Fig. 2C). In conclusion, the data suggest that Asp23 and Asp80 are the catalytic residues in PibD.

Membrane topology analysis of PibD. Previously, a topology model of PibD was suggested based on the prediction by TMHMM 2.0 (6, 38). In this model, the amino terminus is cytoplasmic and followed by five transmembrane segments and an extracellular carboxyl terminus (Fig. 3A). In order to dissect the defect of the D187A/D188A mutant, the respective single mutants were generated. PibD containing the D187A or D188A mutation was active, although there was an apparent reduction in activity in the D187A mutant (Fig. 2C). In conclusion, the data suggest that Asp23 and Asp80 are the catalytic residues in PibD.

% residual fluorescence

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<th>Cysteine mutations</th>
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<tr>
<td>Cysless</td>
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FIG. 3. Membrane topology models of PibD and accessibility of single-cysteine mutants to AMS and fluorescein maleimide in everted membrane vesicles. (A) Representative membrane topology models predicted with TMHMM2.0 (left) and Phobius (right). Critical aspartic acid residues are highlighted as black circles, and positively charged residues (lysine and arginine) are marked with plus signs. Amino acid positions that were altered to cysteine are Glu28 (circled 1), Asn50 (circled 2), Thr63 (circled 3), Ser74 (circled 4), and Ile236 (circled 5). (B) Fluorescence labeling of PibD single-cysteine mutants. Everted E. coli vesicles without or with preincubation with AMS were solubilized, and protein was purified by immobilized metal affinity chromatography. Eluted protein was labeled with fluorescein maleimide and separated by SDS-PAGE. Fluorescent bands were visualized using an F1 Lumi-Imager (upper panel). Subsequently, the same gel was stained with Coomassie brilliant blue (lower panel). Numbers above lanes correspond to those used in panel A. Residual fluorescence of bands is expressed for each mutant as a ratio of AMS prelabeled samples to mock-treated samples after correction for protein loading and the background signal of the cysteineless mutant.
ized water instead of AMS. Subsequently, the reactions were quenched by addition of DTT, membranes were solubilized with detergent, and the His-tagged PibD proteins were enriched by immobilized metal affinity chromatography. Eluted proteins were labeled with fluorescein maleimide and separated by SDS-PAGE, and the fluorescence intensities of pre-labeled samples and mock-treated samples were compared. A strong decrease in fluorescence signal was observed for the E25C mutant by AMS treatment (Fig. 3B), suggesting that this position is cytoplasmic, as predicted by the six-transmembrane model. According to a model with five TMS, this residue should be located on the extracellular side of the membrane and therefore be protected. Similarly, the S74C mutant showed some reduction of fluorescence in AMS-treated membranes. For the N50C and I236C mutants, little decrease of fluorescence intensity was seen, consistent with these residues being inaccessible from the cytoplasmic side of the membrane. In the case of the T63C mutant, the labeling of the cysteine by fluorescein maleimide was very poor, even without pretreatment of the sample with AMS. This indicates that the residue is buried in the protein structure, possibly within a helical transmembrane segment. Moreover, AMS treatment of membranes containing this mutant had a minor effect on subsequent fluorescein maleimide labeling. Quantification of fluorescent band intensities further supports the observation that residues Glu28 and, most likely, Ser74 are exposed to the cytoplasmic side of the membrane (Fig. 3B). In conclusion, our data support a six-transmembrane model for PibD, with extracellular amino and carboxyl termini and the two catalytic residues located in or close to intracellular loops between TMS 1 and 2 and TMS 3 and 4, similar to the second model presented in Fig. 3A. This model also fits better with the distribution of positively charged residues in the protein (Fig. 3A). Taken together, these results indicate that PibD is an aspartyl protease with an unusual topology consisting of extracellular amino and carboxyl termini and six transmembrane segments.

**DISCUSSION**

The family of type IV prepilin peptidases is divided into two subfamilies of bacterial and archaeal enzymes (MEROPS database identifiers A24A and A24B, respectively) (34). The archaeal subfamily has been named preflagellin peptidases because the flagellin subunits were presumed to be the only substrate of the archaeal enzymes. However, PibD from *S. solfataricus* is involved not only in preflagellin processing but also in the processing of a set of binding proteins with prepilin-like signal sequences and of small pilin-like proteins (2, 3). By screening archaeal genomes for putative type IV pilin-like signal sequences, we found that this type of secretory sequence is in fact more widely used by archaea than previously assumed (Z. Szabo, O. de Andrade Oliveira, J. C. Kissinger, S.-V. Albers, A. J. M. Driessen, and M. Pohlschroder, unpublished results). In the present work, we demonstrate that PibD is an aspartic acid protease, as suggested by the inactivation of the enzyme by EDAC and glycinamide, a procedure previously reported to inhibit bacterial TFPPs (28). Site-directed mutagenesis of the nine aspartic acid residues showed that Asp23 and Asp80 are essential for cleavage. The aspartate-to-alanine mutants were inactive, whereas a glutamate substitution at position 80 resulted in an enzyme with partial activity. The D23E mutant, however, did not cleave preflagellin. In aspartate-to-glutamate mutations, amino acid side chain length increases but the negative charge is conserved. It can be concluded that structural constraints are less tolerated at position Asp23 than at position Asp80. Interestingly, similar mutations in TcpJ led to opposite results: a glutamate mutation at the position corresponding to Asp23 in PibD was tolerated better than a mutation corresponding to D80E (28). For TcpJ, it was concluded that the first catalytic amino acid might coordinate a water molecule necessary for hydrolysis and therefore be structurally more flexible. The second would then directly participate in chemical reduction of the peptide bond. However, taking our results into consideration, no common theme concerning the local structural environments of the two catalytic residues can be deduced, and therefore it is difficult to access their specific function during catalysis.

Site-directed mutagenesis of aspartate residues in PibD also identified a pair of amino acids, Asp187/Asp188, that appeared essential for enzyme function. A double-alanine mutant was inactive, whereas single mutations did not critically affect activity. These residues localize within the large cytoplasmic domain that is typically found in the archaeal PibD homologs. Together with the distal transmembrane segment, this domain defines a unique protein family, “Archaeal Peptidase A24 C terminus Type II” (Pfam accession number, PF06847) (9), which was based on an alignment of 10 archaeal sequences, including those of PibD and *M. voltae* FlaK. Although no absolutely conserved residues can be found in the cytoplasmic domain, it is conspicuous that they all contain small clusters of from two to four successive and negatively charged residues (aspartate and/or glutamate). *M. voltae* FlaK also contains a double aspartate at position 185 and 186 which could correspond to Asp187 and 188 in PibD. A D186A mutant was reported to be active in vitro (7), but the role of Asp185 was not investigated. It is therefore not clear whether these aspartate residues are functionally equivalent to those in PibD. Nevertheless, one may speculate that clusters of acidic residues in the cytoplasmic loop play an important role in archaeal TFPP homologs. They might fulfill a structural function by participating in intramolecular salt bridges. Alternatively, these residues could be essential for recognition of the positively charged leader sequence of the substrate protein. An involvement of negatively charged residues of a protease in recognition of positively charged amino acids in the substrate was reported for the yeast (*Saccharomyces cerevisiae*) mitochondrial processing peptidase (MPP). MPP is a heterodimeric metalloprotease that shows a functional similarity to TFPPs, as it cleaves a positively charged leader peptide of precursor proteins imported into mitochondria (for a review, see reference 20). Biochemical and structural studies of MPP revealed that negatively charged residues in the substrate-binding pocket of the enzyme might form stabilizing electrostatic interactions with the substrate (24, 37, 45). If such electrostatic interactions are also required for the interaction of PibD with its substrates, then Asp187 and 188 are likely to play a role.

Cleavage of substrate proteins by prepilin/preflagellin peptidases is expected to occur at the cytoplasmic side of the membrane (10). To define the topology of PibD, we employed cysteine accessibility studies with everted membrane vesicles.
containing single-cysteine mutants. Two positions (Glu28 and Ser74) were found to be exposed to the cytoplasmic side of the membrane. Two mutants (N50C and I236C) were not accessible to AMS, and therefore it can be concluded that these residues are extracellular or within a transmembrane segment. One mutant (T63C) was inefficiently labeled by fluorescein maleimide in the solubilized state of the protein, even without pretreatment with AMS. This residue is likely to be buried within the protein structure, which prevents access of the bulky sulfhydryl-modifying reagents. In none of the mutants could fluorescein maleimide labeling be completely blocked by pretreatment with AMS. This can be explained by the fact that PibD is expressed very poorly in E. coli, a fact which is also reflected by the low protein yield (Fig. 3). Therefore, PibD represents only a minor fraction of the total membrane protein, which in turn leads to very high background labeling of endogenous cysteine containing proteins. This might be compensated for by increasing the concentration of AMS or extending labeling times beyond those used in previously published work. Attempts to modify the labeling protocol in this way, however, led to the modification of all cysteine mutations, suggesting that the probe leaked through the membrane (not shown). In conjunction with the positive-inside rule for membrane proteins (50), our results are consistent with a model of membrane topology and signal peptide prediction method. J. Mol. Biol. 232:275–278.

REFERENCES


