Research Article

Identification of membrane proteins in the hyperthermophilic archaeon *Pyrococcus furiosus* using proteomics and prediction programs

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Abstract

Cell-free extracts from the hyperthermophilic archaeon *Pyrococcus furiosus* were separated into membrane and cytoplasmic fractions and each was analyzed by 2D-gel electrophoresis. A total of 66 proteins were identified, 32 in the membrane fraction and 34 in the cytoplasmic fraction. Six prediction programs were used to predict the subcellular locations of these proteins. Three were based on signal-peptides (SignalP, TargetP, and SOSUISignal) and three on transmembrane-spanning α -helices (TSEG, SOSUI, and PRED-TMR2). A consensus of the six programs predicted that 23 of the 32 proteins (72%) from the membrane fraction should be in the membrane and that all of the proteins from the cytoplasmic fraction should be in the cytoplasm. Two membrane-associated proteins predicted to be cytoplasmic by the programs are also predicted to consist primarily of transmembrane-spanning β -sheets using porin protein models, suggesting that they are, in fact, membrane components. An ATPase subunit homolog found in the membrane fraction, although predicted to be cytoplasmic, is most likely complexed with other ATPase subunits in the membrane fraction. An additional three proteins predicted to be cytoplasmic but found in the membrane fraction, may be cytoplasmic contaminants. These include a chaperone homolog that may have attached to denatured membrane proteins during cell fractionation. Omitting these three proteins would boost the membrane-protein predictability of the models to near 80%. A consensus prediction using all six programs for all 2242 ORFs in the *P. furiosus* genome estimates that 24% of the ORF products are found in the membrane. However, this is likely to be a minimum value due to the programs' inability to recognize certain membrane-related proteins, such as subunits associated with membrane complexes and porin-type proteins. Copyright © 2001 John Wiley & Sons, Ltd.

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Introduction

The advent of genome sequencing has revolutionized the study of microbial physiology. Single protein characterizations and biochemical pathway studies have been augmented by our ability to determine the functional relationships between different pathways and the roles of novel proteins.

This approach, known as functional genomics, typically involves the use of DNA microarrays and proteomics (Dove, 1999; Southern *et al.*, 1999). Furthermore, insight into function may be gained from the three-dimensional structure of a protein determined by structural genomics, which involves the cloning and expression of known and unknown ORFs on a genome-wide scale and analyses of the

product by NMR and/or X-ray crystallography (Burley, 2000).

An extremely important parameter in obtaining recombinant proteins, as well as in all aspects of both functional and structural genomics, is the subcellular location of a protein. Specifically, is the native form membrane-associated or cytoplasmic? This can be assessed in one of three ways. First, the sequences of putative proteins can be compared with those of characterized proteins of known location. However, approximately 50% of all of the predicted ORFs within 44 microbial genomes examined encode (conserved) hypothetical proteins (<http://www.tigr.org/tigr-scripts/CMR2>) and, therefore, cannot be assigned a subcellular location by simple sequence comparisons. Second, subcellular location of an ORF product may be predicted on the basis of sequence analyses for signal peptide sequences and transmembrane spanning α -helices (Andersson *et al.*, 1992; Nielsen *et al.*, 1999). In a few cases, this has been done on a genome-wide basis. 21 genome sequences (four archaea, 14 bacteria, and three eukaryotes) were analyzed using these prediction algorithms (Kihara and Kanehisa, 2000; Mitaku *et al.*, 1999; Paulsen *et al.*, 2000; Wallin and von Heijne, 1998). They predicted that 15–30% of the ORFs in these genomes encode membrane proteins. Third, the location of proteins can be determined physically by separating cell-free extracts into cytoplasmic and membrane-associated fractions and by assessing the protein species present in each. This typically involves two-dimensional electrophoresis and the identification of proteins using mass spectrometry (Cordwell *et al.*, 2000). Separation of membrane and cytoplasmic proteins prior to proteomic analyses considerably improved the resolution and ease of identification of membrane proteins from the bacterium *Pseudomonas aeruginosa* PAO1 (Nouwens *et al.*, 2000).

Few cases can be found where more than one of the three methods used to investigate subcellular location has been used on a comparative basis to assess the validity of the results. Nouwens *et al.* (2000) used membrane protein algorithms to accurately predict *P. aeruginosa* PAO1 membrane proteins identified by proteomic methods. Hence, while there has been some success in predicting the subcellular locations of eukaryotic and bacterial proteins, investigations of the cellular location of proteins from archaeal sources are sparse and are based primarily on the characterization of purified proteins from such organisms. Understanding the

architecture and physical properties of archaeal membranes is extremely important, since they differ from those of bacterial and eukaryotic membranes. In fact, Nielsen *et al.* (1999) have stated that it is unclear whether existing algorithms are adequate for predicting the subcellular locations of archaeal proteins.

To date, the genome sequences of eight hyperthermophilic (i.e., microorganisms with an optimum growth temperature above 80°C (Stetter, 1999)) archaea have been determined, three of which belong to the genus *Pyrococcus*. *Pyrococcus furiosus*, a fermentative sulfur reducer (Fiala and Stetter, 1986) whose genome sequence was recently completed (Robb *et al.*, 2001), is among the most thoroughly studied of the hyperthermophilic archaea (Adams, 1999). The relatively large biochemical literature available for *P. furiosus* together with access to the complete genome sequence suggests that this microbe could serve as a model for archaeal membrane proteins. Cytoplasmic and membrane proteins were isolated from *P. furiosus* cells and the most abundant proteins in each subcellular fraction were separated electrophoretically and identified based on peptide mass comparisons with the predicted amino acid sequences from the *P. furiosus* open reading frames (Robb *et al.*, 2001). The predicted amino acid sequences for the identified proteins were also categorized as membrane or cytoplasmic using various programs that predict signal peptides and transmembrane-spanning α -helices. The objective was to assess their efficacy in membrane protein prediction. A consensus approach was applied genome-wide using six programs to predict the (minimum) number of membrane proteins in *P. furiosus*.

Materials and methods

Subcellular fractionation of proteins

Pyrococcus furiosus (DSM 3638) was grown in a 20-liter fermentor containing 15 liters of medium, which was prepared as described previously (Adams *et al.*, 2001; Verhagen *et al.*, 2001). The basic medium contained 0.5% (w/v) of yeast extract (Difco), casein hydrolysate (enzymatic, Difco), and maltose (Sigma); trace minerals, and the oxygen indicator resazurin in an artificial seawater solution. Where indicated, cultures also contained 0.1% (w/v) elemental sulfur (S⁰). Cultures were also grown on

the same medium except that 0.5% maltose and 0.05% yeast extract were used as the carbon sources. The headspace of the fermentor was flushed with N_2 - CO_2 (80:20), and L-cysteine-HCl· H_2O and Na_2S · $9H_2O$ were added as reducing agents to remove residual O_2 . The pH (measured at room temperature) was adjusted to 6.8 and maintained at 5.9 ± 0.1 and $95^\circ C$ during the incubation.

Cells were harvested in the late-logarithmic phase (1×10^8 to 2×10^8 cells· ml^{-1}) and were cooled to room temperature by pumping them through a glass cooling coil bathed in an ice-water slurry. They were harvested by centrifugation at $10,000 \times g$, resuspended in 15 to 20 ml of Buffer A (degassed 50 mM Tris-HCl (pH 8.0) plus 2 mM sodium dithionite and 2 mM dithiothreitol), and frozen under Ar at $-80^\circ C$. All sample transfers and manipulations were carried out in an anaerobic chamber and all buffers were degassed and flushed with Ar. The cell suspension was thawed, and DNase I in Buffer A was added to a final concentration of 0.0002% (w/v). The cells were disrupted anaerobically by sonication for 30 min. Debris and unbroken cells were removed by centrifugation ($10,000 \times g$ for 15 min), and the supernatant was decanted and centrifuged at $100,000 \times g$ for 45 min. The supernatant was used as the cytoplasmic protein fraction. The membrane pellet was suspended in Buffer A, homogenized using a glass tissue grinder, and then centrifuged at $100,000 \times g$ for 45 min. This procedure was repeated three times, and Buffer A in the final step contained 4 M KCl. The washed membrane pellet was suspended and homogenized in Buffer A, and this formed the membrane protein fraction. All protein fractions were then frozen in liquid N_2 and stored at $-80^\circ C$.

Glutamate dehydrogenase (GDH) activity was determined spectrophotometrically by the reduction of 0.4 mM $NADP^+$ measured at 340 nm ($\epsilon = 6,220$ (M·cm) $^{-1}$) and $80^\circ C$ in 100 mM EPPS buffer (pH 8.4) using 6 mM sodium glutamate as the substrate (Robb *et al.*, 1992). The protein concentration of each fraction was estimated using the Bradford method (Bradford, 1976) with bovine serum albumin as a standard.

Two-dimensional electrophoretic protein separation

Cytoplasmic and membrane protein samples were prepared for two-dimensional gel electrophoresis by mixing thawed fractions with an equal volume

of a solution containing 9 M urea, 2% (v/v) 2-mercaptoethanol, 2% ampholytes (pH 8-10, Bio-Rad; v/v), 4% (v/v) Nonidet P40, and protease inhibitors (Complete Mini Protease Inhibitor Cocktail, Boeringer Mannheim). The samples were spun at $435,000 \times g$ for 10 min (TL100 tabletop ultracentrifuge, Beckman) to remove debris. Protein concentrations of the decanted supernatant were determined using the Ramagli modification of the Bradford protein assay (Ramagli *et al.*, 1985).

First-dimension isoelectric focusing (IEF) was carried out using a 1:7 mixture of pH 3-10 and pH 5-7 carrier ampholytes (Bio-Rad) in 18-cm tube gels with a diameter of 1.5 mm for 14 000 V-h, as described previously (Anderson and Anderson, 1978a). Aliquots containing 20 μg of protein for silver staining or 300 μg of protein for Coomassie Blue staining were loaded onto each IEF gel. After IEF, the gels were equilibrated in a buffer containing sodium dodecyl sulfate (SDS) as described by O'Farrell (1975) and then loaded onto 10-17% polyacrylamide gradient slab gels (Anderson and Anderson, 1978b). The second-dimension separation was performed using the Laemmli buffer system (O'Farrell, 1975). The proteins were then fixed and stained in the gels using 0.2% (w/v) Coomassie blue R-250 in 2.5% phosphoric acid and 50% (v/v) ethanol (Giometti *et al.*, 1987), or fixed in 50% (v/v) ethanol with 0.1% formaldehyde and 1% acetic acid for subsequent staining with silver nitrate (Giometti *et al.*, 1991).

Mass-spectrometric protein identification

Matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry and capillary liquid chromatography-electrospray tandem mass spectrometry (μ -LC-ESI-MS/MS) were used to identify cytoplasmic and membrane *P. furiosus* proteins extracted from 2DE gels. Protein spots to be identified were cut from 1 to 4 replicate gels stained with Coomassie Blue R250 depending on the abundance of individual proteins. The excised spots were then reduced with either dithiothreitol (Sigma) at $60^\circ C$ or tris(2-carboxyethyl)phosphine (Pierce, Rockport, IL) at room temperature, alkylated with iodoacetamide (Sigma), and digested *in situ* overnight with sequencing grade modified porcine trypsin (Promega Corp., 12.5 ng/ μl). The digested peptides were extracted three times with equal parts of 25 mM ammonium bicarbonate and acetonitrile and then twice with equal parts of 5%

(v/v) formic acid and acetonitrile. The resulting extracted tryptic peptides were used directly for μ -LC-ESI-MS/MS without further purification, but were desalted and concentrated with commercial ZipTip C₁₈ pipette tips (Millipore, Bedford, MA) prior to MALDI-TOF peptide mass mapping analysis. Proteins were identified and confirmed by analyzing a same tryptic peptide sample with both MALDI-TOF peptide mass mapping and μ -LC-ESI tandem mass spectrometry followed by database searching.

For MALDI-TOF peptide mass mapping, tryptic digest samples, mixed with α -cyano-4-hydroxycinnamic acid (Sigma), were spotted onto a MALDI target plate and then transported into a Voyager DE-STR MALDI-TOF mass spectrometer equipped with delayed extraction and reflectron (PE Biosystem, Framingham, MA). MALDI-TOF mass spectra for each sample spot were generated by averaging 64–126 N₂ laser shots. Proteins were then identified using PROQUEST, a peptide mass mapping database search algorithm developed in Yates Laboratory, by comparing experimentally obtained mass-to-charge (*m/z*) values with theoretically calculated *m/z* values of tryptic peptides of proteins from a *P. furiosus* open reading frame database (<http://comb5-156.umbi.umd.edu>, GeneMate).

For μ -LC-ESI tandem mass spectrometry, tryptic digests were directly loaded onto a 10–15 cm 365 $\times$ 100 μ m fused silica capillary (FSC) column packed with 10 μ m POROS 10 R2 reverse-phase packing material (PE Biosystem, Framingham, MA) by using a helium-pressurized stainless-steel bomb (Gatlin *et al.*, 1998). The tryptic peptides on the column were separated in thirty minutes by performing liquid chromatography employing the linear gradient of 2–60% solvent B (A: 0.5% acetic acid, B: 80% acetonitrile/0.5% acetic acid), and then introduced to LCQ ion trap mass spectrometer (Finnigan MAT, San Jose, CA). The flow rate at the tip of 200–300 nl/min was maintained during the liquid chromatography by using a precolumn splitter. Tandem mass spectra were automatically acquired in data-dependent mode during the 30-min LC-MS runs by picking three most abundant ions above predefined threshold intensity from previous full MS scan. Obtained MS/MS spectra were then directly searched against a *P. furiosus* open reading frame database with SEQUEST database search algorithm (Eng *et al.*, 1994) without need of prior manual MS interpretation. SEQUEST identified proteins by correlating

experimentally obtained MS/MS spectra to protein sequences in the *P. furiosus* database (Link *et al.*, 1997) and the identified proteins were further verified by manually checking every sequence matched with high cross-correlation scores by SEQUEST.

Membrane protein prediction models

The predicted amino acid sequences for the ORFs identified through proteomics and in the complete genome (Robb *et al.*, 2001) were used to assess the accuracy of various membrane protein prediction programs. The programs are SignalP v1.1 (<http://www.cbs.dtu.dk/services/SignalP>; Nielsen *et al.*, 1997), TargetP v1.0 (<http://www.cbs.dtu.dk/services/TargetP>; Emanuelsson *et al.*, 2000), TSEG (<http://www.genome.ad.jp/SIT/tseg.html>; Kihara *et al.*, 1998), SOSUI and SOSUISignal (<http://sosui.proteome.bio.tuat.ac.jp/cgi-bin/sosui.cgi>?; Hirokawa *et al.*, 1998), and PRED-TMR2 (<http://o2.db.uoa.gr/PRED-TMR2>; Pasquier and Hamodrakas, 1999). SignalP, TargetP, and SOSUISignal are based on the presence of a membrane-anchor/secretory signal peptide sequence at the N-terminus. TSEG, SOSUI, and PRED-TMR2 predict the locations of transmembrane α -helices in the peptide sequence using hydropathy models and homologies to known transmembrane α -helices, and exclude hydrophobic segments associated with globular proteins. The default settings were used for each program, except TSEG where the '5 discriminant functions' setting was used. The SOSUISignal program was run using both 'eukaryote' and 'prokaryote' settings. The ORF product was manually designated as either membrane or cytoplasmic based on the consensus results of the various programs. An ORF product was designated as membranous if at least three of the six membrane protein prediction programs yielded a positive result for that ORF.

The amino acid sequences were also analyzed using MacVector v7.0 software (Oxford Molecular Group, Symantec Corporation). With this program, the amino acid composition was determined and predictions of structure were made using plots of Kyte/Doolittle hydrophobicity (window=7), von Heijne transmembrane (window=21), amphiphilic helix (window=21), amphiphilic sheet (window=7), and Robson-Garnier secondary structure (window=7).

Genome-wide membrane protein estimations

For a genome-wide estimate of membrane-encoding ORFs, each *P. furiosus* protein sequence was run through each of the six membrane-protein prediction web sites. In order to accomplish this the analysis was broken down into three main steps: download & pre-processing, sequence submission and result parsing. In the first step, the *P. furiosus* genome sequence was download in FASTA format from GeneMate and this file was used to produce six more FASTA files that were pre-processed according to each web site's instructions. Two methods were used for sequence submission since the format of the web sites varied. The first method (SOSUISignal, TSEG, SOSUI and PRED-TMR2) relied on a web browser automation program written using Microsoft Visual Basic 6 (VB). The program was designed to submit one sequence at a time to the web site through the browser and wait for a response. Once the program received the requested web page (the results) the information on the page was automatically parsed and saved to a text file for later analysis. In the second method (SignalP and TargetP), each site's e-mail server was utilized where large blocks of the genome were submitted at a time. The returned e-mail results were parsed and saved to a text file using another program written in VB. All six result files were then imported into a Microsoft Excel spreadsheet to aid in calculation. A final prediction of protein cellular location was based on a manual consensus analysis of the results for the six membrane protein models, as described above.

Results

Proteins identified by proteomics

Membrane and cytoplasmic fractions were prepared from *P. furiosus* cells grown on 0.5% (w/v) each of maltose, yeast extract, and casein hydrolysate with and without S^o and from cells grown on 0.5% maltose and 0.05% yeast extract without S^o. For the culture grown with casein but without S^o, the glutamate dehydrogenase (GDH) activities in the cytoplasmic and membrane fractions were 85% and <0.1%, respectively, of the activity in unfractionated cell-free extract. The cytoplasmic and membrane fractions contained 59 and 9%, respectively, of the total protein present in the extract. For the culture grown with casein and S^o, cytoplasmic and

membrane fraction GDH activities were 58 and 0.2%, respectively, of the activity in the cell-free extract, and 45 and 7% of the total protein was recovered in the cytoplasmic and membrane protein fractions, respectively. The cytoplasmic and membrane fractions from cells grown with maltose, no casein and without S^o had GDH activities that were 81 and 0.2%, respectively, of the total activity in the cell-free extract, with 62 and 17% of the total protein recovered in the cytoplasmic and membrane fractions, respectively.

A total of 25 proteins from the membrane fraction of cells grown with casein and without S^o were identified using two-dimensional gel electrophoresis to separate the proteins and mass spectrometry of tryptic peptides to identify their corresponding coding sequence in the open reading frame database (Figure 1). A total of 12 proteins from the membrane fraction of cells grown with S^o were identified, producing a total of 32 unique proteins (5 proteins were present in both samples: Table 1). Of these 32 proteins, one had been characterized previously from *P. furiosus*, 13 were conserved hypothetical proteins, and the remaining 18 were homologs of characterized proteins. At least three of these 32 proteins (Pf336067, Pf1459639, and Pf1825269) were also found in the cytoplasmic fraction or show homology to known cytoplasmic proteins, suggesting that they could be cytoplasmic contaminants in the membrane preparations. The most abundant membrane proteins found on the gels were maltose- and dipeptide-binding protein homologs (Figure 1). A total of 34 proteins were identified in the cytoplasmic fraction (Table 2) and 8 of these were enzymes characterized previously from *P. furiosus*. Of the remainder, 11 were conserved hypothetical proteins, one was a unique hypothetical protein, and the other 14 were homologs of characterized proteins. For each fraction, all of the proteins identified using MALDI-TOF were likewise identified using μ -LC-MS/MS. However, more proteins were identified using the latter method due to the increased sensitivity of the procedure.

Predictability of membrane and cytoplasmic proteins

The signal-peptide recognition programs (SignalP, TargetP, and SOSUISignal) predicted that 17 to 20 of the proteins (59–69%, when excluding the three predicted cytoplasmic contaminants) identified in

Figure 1. Two-dimensional gel electrophoresis pattern of membrane proteins isolated from *P. furiosus* grown without sulfur. Proteins were stained with Coomassie Blue R250. Numbers shown refer to the open reading frame (ORF) designation from the *P. furiosus* genome sequence (Robb et al., 2001). Table I shows the annotation for the ORFs indicated

the membrane fraction contained a signal peptide (Table 1). The transmembrane α -helix recognition programs (TSEG, SOSUI, and PRED-TMR2) predicted that 13 to 23 of these proteins (45–79%) contained at least one transmembrane α -helix. A consensus of the six programs predicted that 23 of these proteins (79%) were in the membrane. The signal-peptide recognition programs predicted that 31 to 34 of the proteins (91–100%) identified in the cytoplasmic fraction did not contain a signal peptide (Table 2). The transmembrane-segment recognition programs predicted that 32 to 33 of these proteins (94–97%) did not contain at least one transmembrane α -helix. A consensus of the six programs predicted that all 34 of these proteins were cytoplasmic. A chi-squared statistical test was

performed on the protein location predictions. For both the membrane and cytoplasmic fractions the predictions were significant (i.e., not random; $0.01 < p < 0.025$ and $p < 0.005$, respectively).

For membrane-associated proteins that were predicted to be cytoplasmic by the above programs, additional analyses were made using MacVector v7.0 sequence analysis software. Two of the nine proteins (Pf159491 and Pf1871822) are predicted to consist primarily of hydrophobic residues with a high degree of β -sheet structure (Figure 2). The Pf1871822 protein contains high concentrations of alanine (11.7%), proline (11.1%) and glycine (15.0%), and the Pf159491 protein contains moderately high concentrations of the same residues (5.9, 3.2, and 15.0%, respectively). The remaining 7

Table 1. Proteins identified by proteomics in the membrane cellular fraction, and their predicted locations based on signal peptide and trans-membrane sequence models

ORF #	Pyrococcus furiosus Genome Annotation	Closest Characterized Homolog (except for conserved hypotheticals)	Id./Sim. (%)	Signal Peptide			Pred. # of Transmem. Seg.				Consensus Pred.
				SignalP	TargetP	SOSUI	TSEG	SOSUI	PRED- TMR2		
123510	Putative periplasmic sugar binding protein	Thermococcus litoralis AAC38136	19/32	Yes	Yes	Yes	1	1	0	Mem.	
136581	Conserved hypothetical protein	Pyrococcus abyssi F75209	37/51	Yes	Yes	No	1	1	1	Mem.	
159491	Conserved hypothetical protein	Pyrococcus abyssi H75211	78/88	No	No	No	0	0	0	Cyto.	
164091	Oligosaccharyl transferase	Saccharomyces cerevisiae BAA06079	13/27	Yes	No	No	10	14	14	Mem.	
188579	SST3 subunit related protein										
188579	ATPase subunit K	Desulfurococcus sp. SY AAB64412	45/56	Yes	Yes	Yes	2	4	4	Mem.	
189210	ATPase subunit E	Desulfurococcus sp. SY AAB64413	59/76	No	No	No	0	0	0	Cyto.	
190329	ATPase subunit C	Desulfurococcus sp. SY AAB64414	55/73	No	Yes	Yes	1	1	0	Mem.	
203524	Conserved hypothetical protein	Pyrococcus horikoshii BAA31089	81/90	Yes	Yes	Yes	2	3	2	Mem.	
336067	Related to inosine monophosphate dehydrogenase	Methanococcus jannaschii Q59011	17/32	No	No	No	0	0	0	Cyto.	
372029	Dipeptide-binding protein	Bacillus subtilus AAA62358	19/31	Yes	Yes	Yes	2	5	2	Mem.	
476242	Conserved hypothetical protein	Pyrococcus abyssi B75005	44/61	Yes	Yes	Yes	1	1	0	Mem.	
1074447	Conserved hypothetical protein	Archaeoglobus fulgidus F69482	44/56	No	No	No	0	0	0	Cyto.	
1136394	Conserved hypothetical protein	Aquifex aeolicum AAC07600	22/40	No	No	No	0	0	0	Cyto.	
1137142	Conserved hypothetical protein	Pyrococcus abyssi A75157	48/63	No	No	No	0	0	0	Cyto.	
1151577	Oligopeptide ABC transporter (oligopeptide-binding protein)	Bacillus subtilus P42061	20/33	Yes	Yes	Yes	2	2	1	Mem.	
1186264	Conserved hypothetical protein	Pyrococcus abyssi D75122	30/40	No	No	No	1	1	1	Mem.	
1314580	Putative ATPase, vanadate-sensitive	Methanococcus voltae A38542	22/37	Yes	Yes	Yes	2	1	0	Mem.	
1324824	Probable dipeptide-binding protein	Escherichia coli P23847	19/31	Yes	Yes	Yes	1	1	0	Mem.	
1346821	Conserved hypothetical protein	Pyrococcus horikoshii E71018	75/85	Yes	Yes	Yes	1	1	0	Mem.	
1348779	Conserved hypothetical protein	Methanococcus jannaschii F64490	36/53	No	No	Yes	2	3	2	Mem.	
1406506	Conserved hypothetical protein	Pyrococcus horikoshii BAA30439	52/71	Yes	Yes	Yes	2	2	2	Mem.	
1432096	Stomatin homolog	Rhizobium etli AAA84990	34/51	Yes	Yes	Yes	5	5	7	Mem.	
1459639	DNA directed RNA polymerase subunit B	Thermococcus celer P31814	84/91	No	No	No	0	0	0	Cyto.	
1575846	Hypothetical lipoprotein	Borrelia burgdorferi CAA65879	24/37	Yes	Yes	No	1	2	0	Mem.	
1616027	Trehalose/maltose binding protein (MalE)	Thermococcus litoralis AAC38136	93/93	Yes	Yes	Yes	1	2	0	Mem.	
1650532	Iron (III) ABC transporter, ATP-binding protein (HemV-2)	Vibrio anguillarum P11460	26/44	Yes	Yes	Yes	1	1	1	Mem.	
1732661	O-linked GlcNAc transferase	Rat P56558	71/12	Yes	Yes	No	4	4	4	Mem.	
1783837	Amylopullulanase	Pyrococcus furiosus AF016588	100/100	Yes	Yes	Yes	1	0	0	Mem.	

Table I. Continued

ORF #	Pyrococcus furiosus Genome Annotation	Closest Characterized Homolog (except for conserved hypotheticals)	Id./Sim. (%/%)	Signal Peptide		Pred. # of Transmem. Seg.				Consensus Pred.
				SignalP	TargetP	SOSUI	TSEG	SOSUI	PRED- TMR2	
1788984	Putative sugar binding protein (MalE-like)	Salmonella typhimurium P19576	28/45	Yes	Yes	No	1	1	0	Mem.
1817174	Conserved hypothetical protein	Pyrococcus horikoshii F71220	84/91	Yes	Yes	Yes	1	1	1	Mem.
1825269	Thermosome, single subunit	Desulfurococcus sp. SY AAB35235	87/90	No	No	No	0	0	0	Cyto.
1871822	Conserved hypothetical protein	Pyrococcus abyssi D75078	65/74	No	No	No	0	0	0	Cyto.
Totals ^a :				20 (69.0%)	20 (69.0%)	17 (58.6%)	23 (79.3%)	22 (75.9%)	13 (44.8%)	23 (79.3%)

^aThree of the 32 proteins from the membrane fraction are presumed to be cytoplasmic contaminants and were not included for the percentage calculation.

proteins from the membrane fraction identified as "cytoplasmic" had low concentrations of hydrophobic residues.

Genome-wide estimate of membrane proteins

The SignalP, TargetP and SOSUISignal programs predicted that 389 (17%), 465 (21%) and 270 (12%), respectively, of the ORFs in the *P. furiosus* genome encode for proteins that contain a membrane target signal. The TSEG, SOSUI and PRED-TMR2 programs predict that 563 (25%), 628 (28%) and 452 (20%), respectively, of the ORFs encode for proteins that contain at least one transmembrane α -helix. Consensus predictions using the results from all six programs suggest that 533 (24%) of the ORFs in *P. furiosus* encode proteins located in the membrane.

Discussion

The near absence of glutamate dehydrogenase activity, a major cytoplasmic protein in *P. furiosus* (Robb *et al.*, 1992), in the membrane fraction suggests that the method of subcellular fractionation is effective. Our results indicate that between 7 and 17% of the total cellular protein is located in the membrane fraction of *P. furiosus*, depending on the growth conditions. For the proteins identified in the membrane fraction, the consensus of the signal peptide and transmembrane α -helix programs predicted that $\geq 79\%$ should be membrane proteins. For the proteins identified from the cytoplasmic fraction, the programs predicted that all are cytoplasmic. Those proteins in the membrane fraction that were not predicted to be associated with the membrane may 1. be composed of transmembrane β -sheets rather than α -helices, 2. form complexes with other proteins that are in the membrane, 3. represent cytoplasmic contaminants, or 4. be novel membrane proteins. At present, we have no suitable method to distinguish between these possibilities. In a genome-wide analysis using the six programs, it is estimated that 533 ORFs (24%) in the genome of *P. furiosus* encode for membrane proteins, although, based on the sequence analyses of the proteins identified by proteomics, this is may underestimate the actual number of membrane proteins by up to 20%.

Eight proteins from *P. furiosus* whole cell lysates were previously identified based on co-migration

Table 2. Proteins identified by proteomics in the cytoplasmic cellular fraction, and their predicted locations based on signal peptide and trans-membrane sequence models

ORF #	Pyrococcus furiosus Genome Annotation	Closest Characterized Homolog (except for conserved hypotheticals)	Id./Sim. (%/%)	Signal Peptide		Pred. # of Transmem. Seg.				Consensus Pred.
				SignalIP	TargetP	SOSUI	TSEG	SOSUI	PRED- TMR2	
209264	Conserved hypothetical protein	Pyrococcus horikoshii BAA31083	89/93	No	No	No	0	0	0	Cyto.
232621	Enolase (2-phosphoglycerate dehydratase)	Bacillus subtilis P37869	59/71	No	No	No	0	0	0	Cyto.
275295	2-amino-3-ketobutyrate CoA ligase (glycine acetyl transferase)	Bacillus subtilis CABI 3573	47/67	No	No	No	0	0	0	Cyto.
283928	Alpha-amylase	Pyrococcus furiosus AAA72035	99/99	No	No	No	0	0	0	Cyto.
327695	ADP-dependent glucokinase	Pyrococcus furiosus AAD48401	100/100	No	No	No	0	0	0	Cyto.
336067	Related to inosine monophosphate dehydrogenase	Methanococcus jannaschii Q59011	17/32	No	No	No	0	0	0	Cyto.
392566	Conserved hypothetical protein	Pyrococcus abyssi CAB50420	96/98	No	No	No	0	0	0	Cyto.
518483	Conserved hypothetical protein	Pyrococcus horikoshii BAA29892	91/95	No	No	Yes	0	0	0	Cyto.
630041	Conserved hypothetical protein	Pyrococcus horikoshii BAA29833	62/73	No	No	No	0	0	0	Cyto.
635169	Conserved hypothetical protein	Pyrococcus abyssi CAB50211	98/99	No	No	No	0	0	0	Cyto.
703868	Flavoprotein	Mb. thermoautotrophicum S66533	34/53	No	No	No	0	0	0	Cyto.
857495	Conserved hypothetical protein	Pyrococcus horikoshii BAA29889	58/65	No	No	No	0	0	0	Cyto.
925374	Pyruvate ferredoxin oxidoreductase subunit beta-2	Pyrococcus furiosus X85250	100/100	No	No	Yes	0	0	0	Cyto.
926380	Pyruvate ferredoxin oxidoreductase subunit alpha-2	Pyrococcus furiosus X85250	100/100	No	Unknown	No	1	0	0	Cyto.
928888	2-ketoglutarate ferredoxin oxidoreductase subunit alpha-2	Pyrococcus furiosus X85250	100/100	No	No	No	0	0	0	Cyto.
930439	Pyruvate ferredoxin oxidoreductase/2-ketoglutarate ferredoxin oxidoreductase gamma subunit	Pyrococcus furiosus X85250	100/100	No	No	No	0	0	0	Cyto.
987716	Methionyl-tRNA synthetase	Mb. thermoautotrophicum O26687	40/59	No	No	No	0	0	1	Cyto.
996088	Conserved hypothetical protein	Pyrococcus abyssi CAB50022	62/84	No	No	No	0	0	0	Cyto.
1132938	Conserved hypothetical protein	Pyrococcus horikoshii BAA29300	38/55	No	No	No	0	0	0	Cyto.
1197153	Translation initiation factor (eIF-5A)	Neurospora crassa P38672	31/55	No	No	No	0	0	0	Cyto.
1199768	Cystathionine gamma-lyase (gamma-cystathionase)	Bacillus subtilis CABI 4467	37/55	No	No	No	0	0	0	Cyto.
1210188	Superoxide reductase	Pyrococcus furiosus AAF03227	100/100	No	No	No	0	0	0	Cyto.
1210814	Rubryerythrin	Pyrococcus furiosus AAF03227	100/100	No	No	No	0	0	0	Cyto.
1308192	Phosphoglycerate dehydrogenase	Methanococcus jannaschii Q58424	31/39	No	No	No	0	0	0	Cyto.
1368790	L-asparaginase (L-asparagine aminohydrolase)	Bacillus subtilis P26900	35/52	No	No	No	0	0	0	Cyto.
1425928	Ethylene-inducible protein homolog	Emicella nidulans AF133101	54/65	No	No	No	0	0	0	Cyto.

Table 2. Continued

ORF #	Pyrococcus furiosus Genome Annotation	Closest Characterized Homolog (except for conserved hypotheticals)	Id./Sim. (%/%)	Signal Peptide		Pred. # of Transmem. Seg.				Consensus Pred.
				SignalP	TargetP	SOSUI	TSEG	SOSUI	PRED- TMR2	
1509633	Myo-inositol-1-phosphate synthetase (InoI)	Drosophila melanogaster AAD02819	16/29	No	No	Yes	0	0	0	Cyto.
1548241	Imidazoleglycerol-phosphate synthetase, cyclase subunit (HisF)	Azospirillum brasilense P26721	57/74	Yes	No	No	0	0	0	Cyto.
1562997	Putative 3-isopropylmalate dehydratase large subunit	Aspergillus nidulans Q92412	38/56	No	No	No	1	1	1	Cyto.
1692502	Conserved hypothetical protein	Pyrococcus horikoshii D71189	89/93	No	No	No	0	0	0	Cyto.
1721830	Conserved hypothetical protein	Pyrococcus abyssi CAB49302	91/95	No	No	No	0	0	0	Cyto.
1782302	Conserved hypothetical protein	Pyrococcus abyssi F75216	77/86	No	No	No	0	0	0	Cyto.
1825269	Thermosome, single subunit	Desulfurococcus sp. SY AAB35235	87/90	No	No	No	0	0	0	Cyto.
1889233	Hypothetical protein	-	-	No	No	No	0	0	0	Cyto.
Totals:				1 (2.9%)	0 (0%)	3 (8.8%)	2 (5.9%)	1 (2.9%)	2 (5.9%)	0 (0%)

with purified *P. furiosus* proteins on two-dimensional electrophoresis gels (Giometti *et al.*, 1995). The present work, however, represents the first broad-based membrane-protein identification from a hyperthermophilic archaeon. The most abundant membrane proteins in *P. furiosus* within the pI and size ranges analyzed appear to be associated with binding proteins for either peptides or maltose (Figure 1). This is consistent with the fact that this organism uses such compounds as its primary carbon source. Some of the *P. furiosus* membrane proteins did not resolve well on two-dimensional gels, especially in the first dimension, indicating that specialize solubilization conditions are needed to optimize the resolution of hyperthermophilic membrane proteins as has been observed for numerous mesophilic organisms as well (Chevallet *et al.*, 1998). Membrane-spanning proteins are generally perceived as containing hydrophobic α -helices of >20 residues, but this model excludes integral membrane proteins that are highly polar, lack hydrophobic segments, and consist predominantly of β structure (Cowan and Rosenbusch, 1994). This latter family of proteins consists primarily of porins, which are gated channels across the membrane that facilitate the diffusion of small polar solutes. Further analysis of membrane-associated proteins of *P. furiosus* predicted by the programs to be cytoplasmic suggests that two of these proteins (Pf159491 and Pf1871822) may be membrane proteins consisting primarily of β -structure (Figure 2). The Pf1871822 protein, annotated as a conserved hypothetical protein, is a S^o-responsive protein that dramatically increases in relative abundance when cultures are grown with S^o (J. Holden and M. Adams, unpublished data; Schut *et al.*, submitted). The protein is repeatedly found in the membrane fraction using the fractionation methods described herein. Hydrophilicity, transmembrane, amphiphilic helix and sheet, and secondary structure models predict that the protein is mostly transmembrane and β -sheet structure (Figure 2A). The protein contains high concentrations of alanine, proline and glycine, which are highly abundant in porin β -barrel proteins (Li *et al.*, 1996). The product of Pf159491, another conserved hypothetical protein, may also be a β -structure membrane protein, as it is predicted to be predominantly transmembrane with up to 14 β -sheet structures, and contains moderately high concentrations of the same three amino acid residues (Figure 2B).

Figure 2. Predictions of Kyte/Doolittle hydrophility, von Heijne transmembranes, amphiphilic helices and sheets, and Robson-Garnier secondary structure for the putative membrane proteins encoded by Pf1871822 (A) and Pf159491 (B). The predictions are derived from the MacVector v7.0 sequence analysis software

The product of Pf189210 is homologous to subunit E of a characterized membrane-bound V-type ATPase from the hyperthermophilic archaeon *Desulfurococcus* sp. SY (Shibui *et al.*, 1997). Although predicted to be a cytoplasmic protein by the programs used, this protein was among the proteins identified in the membrane fraction after separation by 2DE. Two other subunits of this ATPase (subunits K from Pf188579 and C from Pf190329), predicted to be membranous, were also identified in the membrane fraction. Therefore, the Pf189210 protein may form a membrane complex with the Pf188579 and Pf190329 proteins. Two *P. furiosus* operons encoding the characterized membrane hydrogenase and the Nuo homologs also contain a mixture of known and presumed membrane and cytoplasmic proteins (Sapra *et al.*, 2000; R. Sapra and M. Adams, unpublished data). The 'cytoplasmic' components of these complexes may also be associated with the membrane and in this case the prediction that they are cytoplasmic proteins by the programs that were used herein would be inaccurate.

The Pf1074447 protein, a conserved hypothetical,

is predicted to be part of a 13 ORF operon consisting entirely of conserved hypothetical ORFs (assuming an operon contains two or more ORFs that are separated by less than 16 nucleotides). Using the membrane prediction programs, at least one of the other ORFs in this operon (Pf1071624) encodes for a membrane protein. Therefore, the Pf1074447 protein could potentially form a complex with the Pf1071624 protein in the membranes. None of the ORFs encoding the other cytoplasmic-predicted proteins from the membrane fraction were in putative operons with a predicted membrane-protein-encoding ORF. However, these proteins may complex with other membrane proteins or have a unique membrane protein structure. Three of the proteins identified in the membrane fraction of *P. furiosus* that were predicted to be cytoplasmic (Pf336067, Pf1459639 and Pf1825269) show homology to known cytoplasmic proteins (i.e., inosine monophosphate dehydrogenase, DNA directed RNA polymerase subunit B, and thermosome, respectively). Pf336067 and Pf1825269 were also identified in the cytoplasmic protein fraction (Table 2). It seems likely that they are cytoplasmic

proteins that are contaminating the membrane fraction. For example, the Pf1825269 protein that is highly homologous to a known hyperthermophilic archaeal chaperonin may have attached to membrane proteins that denatured during the fractionation process. Nouwens *et al.* (2000) did not find any cytoplasmic contaminants in their membrane protein fractions. The presence of cytoplasmic contaminants in our membrane protein preparations may be explained by our method of protein detection and identification (i.e., μ -LC-MS/MS and MALDI-TOF), which is significantly more sensitive than their method (MALDI-TOF only).

In two instances, there was agreement among signal peptide predictions and among transmembrane sequence predictions, but disagreement between the two sets of predictions. For the Pf186264 protein from the membrane fraction, no signal peptide was recognized, but one transmembrane sequence was predicted. The Pf186264 protein shows 30% identity with the *P. abyssi* protein D75122 when analyzed across the entire protein, but aligns with the C-terminal end of the protein with 53% identity and 71% similarity. Pf184685 is 344 bp upstream of Pf186264 and aligns with the N-terminal end of the same *P. abyssi* protein with 29% identity and 51% similarity. The same trends were observed when these two ORF products were compared with *P. horikoshii* protein B71009. The Pf184685 protein contains a putative signal peptide sequence. Therefore, there appears to be a frameshift in the *P. furiosus* nucleotide sequence and Pf184685 and Pf186264 may belong to the same ORF. For the Pf1562997 protein from the cytoplasmic fraction, the predicted transmembrane sequence is between residues 115 and 137 (out of 380) and most likely does not represent a membrane protein. These results underscore the importance of using the consensus of multiple models coupled with other analyses on questionable ORF product predictions to accurately predict protein subcellular locations.

Using a consensus of all six membrane protein prediction models, it was estimated that 24% of the ORF's in the *P. furiosus* genome encode for a membrane protein. This value is similar to those predicted for other hyperthermophilic archaea for which complete genome sequences are available, which include *P. horikoshii*, *Archaeoglobus fulgidus*, and *Methanococcus jannaschii* (Kihara and Kanehisa, 2000; Mitaku *et al.*, 1999; Pasquier and

Hamodrakas, 1999; Paulsen *et al.*, 2000; Wallin and von Heijne, 1998). For each of these predictions only one analysis program was used, and it was predicted that 14–23% of the ORF's in their genome encode membrane proteins. Based on our sequence analyses of proteins identified by proteomics, it appears that the most accurate estimate of subcellular location occurs when a consensus is made between the six programs.

The signal peptide and transmembrane α -helix programs used in this study generally predict accurately the subcellular location of the proteins identified by proteomics in membrane and cytoplasmic fractions when used in a consensus fashion. Therefore, they are useful for "first pass" predictions of membrane proteins, although they do not detect membrane proteins that consist of β -sheets rather than α -helices, nor do they recognize 'cytoplasmic' proteins that form complexes with membrane proteins. Therefore, careful follow-up analyses are necessary for a complete census of membrane proteins. Nielsen *et al.* (1999) stated that much more is known about membrane proteins from organisms other than archaea, and that their analyses with *M. jannaschii* suggest unique membrane proteins structures may exist in this organism, as well as in other archaea. Very little is known about membrane proteins in *P. furiosus*. Accurate membrane protein prediction models are vital for research on *P. furiosus*, especially for on-going structural genomic efforts and other analyses involving the membrane.

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