Genomics of the thermo-acidophilic red alga \textit{Galdieria sulphuraria}

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ABSTRACT

Extremophilic organisms dwell in environments that are characterized by high or low temperatures (thermophiles or psychrophiles), very low or high pH-values (acidophiles or alkalophiles), high salt concentrations (halophiles), high pressure (barophiles), or extreme drought (xerophiles). Many extremophiles are microbes, and many also belong to the prokaryota. \textit{Galdieria sulphuraria}, however, is a member of a group of extremophilic eukaryotes that are named Cyanidiales. Cyanidiales are unicellular red micro-algae that occur worldwide in hot acidic waters, volcanic calderas, and in human-made acidic environments such as acidic mine drainage. \textit{G. sulphuraria} has a unique position within the Cyanidiales because, in contrast to the other obligate photoautotrophic members of this group, it is able to grow photoautotrophically, mixotrophically, and heterotrophically. It is not only resistant to acid (pH 0) and heat (56ºC), but also to high salt (1.5 M NaCl), toxic metals, and many other abiotic stressors. This unusual combination of features such as thermophily, acidophily, resistance to a wide array of abiotic stressors, and an extraordinary metabolic plasticity make \textit{G. sulphuraria} highly interesting model organism to study adaptation to extreme environments. We have started a genomics approach to gain insight into the biology of \textit{G. sulphuraria} and to identify genes and gene products critical for survival under extreme conditions. To this end, we pursue a whole-genome, shotgun sequencing approach towards unraveling the genome sequence of \textit{G. sulphuraria}. We report here on the status quo of the genome-sequencing project and we summarize what we have learned to date from the genome sequence about the biology of this truly unique extremophile.

Keywords: Extremophilic, thermophilic, acidophilic, photosynthesis, red alga, genome sequencing, Cyanidiales, metabolic plasticity

1. INTRODUCTION

1.1 Systematics and phylogenetic position of \textit{G. sulphuraria}

\textit{G. sulphuraria} belongs to a small group of evolutionary anciently diverged, unicellular micro-algae called Cyanidiales. Based on molecular clock estimates, Cyanidiales are probably more than 1.5 billion years old and thus likely some of oldest extant eukaryotic organisms. Cyanidiales form an ancient monophyletic group within the red algae that is located at the root of secondary endosymbiosis \cite{1,2}. These findings are supported by detailed studies of the evolution of starch metabolism in apicomplexa, rhodophytes, and chlorophytes \cite{3}. Cyanidiales exist in many parts of the world in hot acidic habitats, both natural and human-made. This group consists of the three genera \textit{Cyanidioschyzon}, \textit{Cyanidium}, and \textit{Galdieria}. Based on detailed morphological studies and the mode of cell division, six independent species have been established: \textit{Cyanidioschyzon merolae}, \textit{Cyanidium caldarium}, \textit{Galdieria maxima}, \textit{Galdieria partita}, \textit{Galdieria daedala}, and \textit{Galdieria sulphuraria} \cite{4}. The taxonomic positions of \textit{Cyanidium}, \textit{Cyanidioschyzon} and \textit{Galdieria} have been recently updated \cite{5} and the proposed taxonomy is in concordance with recent molecular evidence \cite{6,7}. In this manuscript, we will address \textit{Galdieria sulphuraria} (Galdieri) Merola \cite{6}, the metabolically most versatile member of the Cyanidiales. It is important to note that much of the classic literature on \textit{Cyanidium caldarium} actually applies to \textit{Galdieria sulphuraria}, due to previous uncertainties in the phylogenetic positions of both organisms.

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1.2 Cyanidiales are the only photosynthetic organisms able to thrive in hot acid
Most extremophiles are microorganisms. For example, the presently known upper temperature limits are 113°C for archaeal, 95°C for bacterial, and 62°C for eukaryotic microorganisms. With few exceptions, metazoans are unable to grow above 50°C.8,9. Surprisingly, photosynthetic prokaryotes such as the cyanobacteria are completely absent from hot acid waters. Instead, thermo-acidophilic photosynthetic unicellular red algae of the Cyanidiales are the principal photosynthetic organisms found in these ecological niches.8,11,12. These obligate acidophilic eukaryotic microorganisms can grow at pH 0 and temperatures up to 56°C.13,14.

Figure 1: Confocal laser scanning image of three G. sulphuraria cells. Excitation wavelength was 543 nm and a 650 nm long pass emission filter was used. The image shows chlorophyll autofluorescence of the tubular shaped chloroplasts in each cell. The scale bar to the lower right is 2 µm.

1.3 Galdieria sulphuraria is an extraordinarily stress tolerant and metabolically versatile microorganism
G. sulphuraria naturally inhabits volcanic environments, such as hot sulfur springs and solfatara soils (pH 0 to 4 and temperatures up to 56°C), but also equally hostile environments originating from human activities such as strip or opencut mining. Galdieria frequently represents up to 90% of the total biomass and almost 100% of the eukaryotic biomass under these conditions. Cyanidiales are the principle photosynthetic organisms in these ecological niches.

Galdieria sulphuraria is able to grow photoautotrophically, mixotrophically, and heterotrophically. It thrives on more than 50 different carbon sources such as sugars, sugar alcohols and amino acids, a metabolic flexibility matched by few other organisms.

In addition, G. sulphuraria tolerates up to 9% (w/v) salinity and its cells possess a very rigid protein-rich cell wall. Galdieria thrives in pure CO₂ and elevated atmospheric pressure, leading to the proposal that it could survive even on Venus.22.

Acidic environments are frequently characterized by high concentrations of iron, cadmium, aluminum, nickel, and other metals that are highly toxic to almost all organisms. G. sulphuraria is resistant to high levels of toxic metal ions including 200 mM aluminum. The resistance is likely due to active secretion of metal ions but such a mechanism has yet to be confirmed.

Surprisingly, G. sulphuraria, in contrast to acidophilic green algae such as Dunaliella acidophila, is able to maintain a proton gradient of 1:1 million across its plasma membrane during several months of continuous darkness, indicating an energy independent proton-exclusion mechanism. The mechanism by which this temporary impermeability to protons is achieved is also unclear. It has been hypothesized, based on studies with archaeabacteria, that the incorporation of sterols, saturated fatty acids, bipolar tetraether lipids, and proteins could account for this impermeability.27-30. Very long chain dicarboxylic acids are thought to have similar roles in some eubacteria.31-33. Possibly also the rigid, protein-rich cell wall is involved in excluding protons.

1.4 Life in hot acid and its relation to protein structure
Photosynthesis in hot, acidic environments is challenging because of an increased oxygenation reaction of ribulose 1,5-bisphosphate carboxylase/oxygenase (RubisCO) at high temperature, and a strong decrease of dissolved inorganic carbon availability at low pH. Therefore, specific adaptations of the carbon assimilatory apparatus to these extreme conditions are to be expected. Galdieria RubisCO is relatively heat stable.34. The crystal structure of RubisCO from G. partita was recently solved and it was demonstrated that this enzyme had the highest specificity in the direction of the carboxylation reaction of RubisCOs hitherto reported.35. Galdieria Rubisco has a unique hydrogen bond between the main chain oxygen of Val332 on loop 6 and the epsilon-amino group of Gln386 of the same large subunit. It was proposed that this interaction is crucial for stabilizing loop 6 in the closed state and for generating a higher affinity for anionic ligands.36. The enhanced thermo-stability of Galdieria proteins should facilitate efforts to gain structural information. We expect Galdieria proteins to become an important resource for structural studies of soluble and membrane proteins; in particular those, that do not have orthologs in prokaryotic extremophiles. Functional genomics of G. sulphuraria and other eukaryotic extremophiles will open up the way to...
explore the molecular mechanisms of eukaryotic life under extreme conditions. In addition, it will provide a rich source of genes with high potential for bioengineering, novel metabolites, stress resistance, and bioremediation.

1.5 Comparative genomics of Cyanidiales

Comparative genomics (i.e., the comparison of genomes of different but related organisms) is a powerful tool to unravel the molecular foundations of observable traits and phenotypes that cannot easily be deduced from the analysis of individual genome sequences. Its core hypothesis is that conserved regions of DNA between two species often encode for the common features of the organisms, while different traits will appear as differences in the genetic makeup between the two species 40. Similar genomes, such as those of Galdieria and Cyanidioschyzon, are particularly useful for elucidating key differences that account for structural and metabolic differences in these organisms. Using the recently published genome sequence of C. merolae 41 and a G. sulphuraria genomic dataset consisting of an EST-collection 42 and 8 Mbp of non-redundant genomic sequence (approximately 40-50% genome coverage; http://genomics.msu.edu/galdieria), we have recently conducted a comparative genomics pilot study to identify those genes of G. sulphuraria that might be crucial to its extraordinary metabolic flexibility 43. To this end, we identified Galdieria sequences that did not match any of the C. merolae genes and that displayed similarity to genes of known function from other organisms. Although only 50% of the G. sulphuraria genome sequence had been finished at the time of the study, a number of important conclusions could be drawn: (i) Despite their evolutionary distance, the Cyanidiales have retained a high level of overall-similarity in their genomes, (ii) Galdieria genes contain more introns than those of Cyanidioschyzon, (iii) only Galdieria is capable of metabolizing complex cell wall polysaccharides, (iv) the lack of heterotrophy in Cyanidioschyzon is not accompanied by a strong reduction in its carbohydrate metabolism enzymatic make-up, and (v) the Galdieria genome encodes many more membrane transporters than that of Cyanidioschyzon 43.

2. METHODOLOGY

2.1 Isolation of genomic DNA from G. sulphuraria

G. sulphuraria cells were harvested from heterotrophic cultures 16 by centrifugation, frozen in liquid nitrogen, and ground to a fine powder by mortar and pestle. Total nucleic acids were extracted by incubating the ground tissue overnight in 50 mM Tris-Cl pH7.5, 5 mM EDTA, and 1% (w/v) SDS, followed by extraction of proteins with phenol:chloroform:isoamylalcohol (24:24:1), and precipitation of DNA from the aqueous phase by ethanol. The pellet was dissolved in 10 mM Tris-Cl pH 7.5, 1 mM EDTA, and RNA was removed by incubation with DNAsafe RNase, followed by de-proteination with phenol:chloroform:isoamylalcohol, and DNA-precipitation by ethanol. Genomic DNA was further purified by CsCl-density gradient centrifugation of bis-benzamide-treated total DNA as described previously 44.

2.2 Construction of genomic DNA libraries

Small insert plasmid libraries: Small-insert shotgun sequencing plasmid libraries were constructed in pSMART-HC Kan (Lucigen Corp., Middleton, WI; www.lucigene.com). Genomic DNA was randomly fragmented by shearing using a HydroShear device 45 and end repaired to generate blunt ends. Fragments of approximately 2 kbp (isolated...
by preparative agarose gel electrophoresis) were ligated into the pSMART vector and plasmids were transformed into *E. coli* cells (Lucigen).

**Fosmid library:** A fosmid library containing 40 kbp inserts was constructed by physically shearing genomic DNA and ligation of end repaired, size fractionated DNA into the Fosmid vector pCC1FOS, followed by packing into Lambda phages.

**BAC library:** Two BAC libraries (BamHI, EcoRI) containing 100 kbp inserts were constructed using the CopyControl BAC Cloning Kit (Epicentre Technologies, Madison, WI) according to instructions by the manufacturer. High molecular weight genomic DNA embedded in LMP agarose plugs was partially digested by EcoRI or BamHI, respectively. Fragments were size fractionated by pulsed-field gel electrophoresis and isolated by electro-elution. Fragments were ligated into predigested and dephosphorylated pCCBAC vectors.

### 2.3 Plasmid isolation and sequencing
A GeneMachines Mantis colony picker was used to pick bacterial colonies into 96-well plates. Qiagen 3000 robots were used for liquid handling and plasmid purification, in addition to a GeneMachines RevPrep Orbit plasmid purification system. Sequencing reactions were performed on a PE9700 thermocycler and an ABI 3730xl high-throughput capillary DNA sequencing systems was used to acquire sequence information. All sequence data and chromatograms were stored on a Geospiza Finch server (Geospiza, Seattle, WA).

### 2.4 Sequence data and similarity searches
*Cyanidioschyzon merolae* predicted peptide sequences were obtained from [http://merolae.biol.s.u-tokyo.ac.jp](http://merolae.biol.s.u-tokyo.ac.jp) (Matsuzaki et al., 2004), data released March 16, 2004. Genomic sequences of *G. sulphuraria* were screened against a database consisting of *Porphyra purpurea* chloroplast, *Cyanidium caldarium* str. RK1 chloroplast, and *Cyanidioschyzon merolae* plastid and mitochondrion sequences as published in Genbank. EST libraries and sequences were reported previously 42. Similarity searching was done with the BLAST 2.2.6 program (NCBI) 46. Data were parsed and analyzed using software implementations developed by the Genomic Technology Support Facility (GTSF) at Michigan State University (http://genomics.msu.edu).

### 3. RESULTS AND DISCUSSION

#### 3.1 Current status of the genome-sequencing program
Literature data on the genome size of *Galdieria spec.* vary between 10.8 Mb and 14.2 Mb: Using microspectrophotometry, a genome size of 10.8 Mbp was estimated for *G. sulphuraria* 074W 47. Genome sizes ranging between 9.8 Mbp (*G. sulphuraria* 19.71) and 14.2 Mbp (*Galdieria* spec. isolates, Rio Tinto, Spain) were determined using pulsed-field gel electrophoresis 48. The genome of the related species *C. merolae* was recently finished and its 16.5 Mb encode 5,331 genes 41. At the time of writing (June 2005), we have obtained 153,814 random sequence reads from small-insert plasmid libraries, and from cosmid and BAC ends. These equal 94.3 Mb of Q20 bases and 15.5 Mb of non-redundant sequence (5.1-fold coverage), indicating that the *Galdieria* genome is larger than expected from literature data. Based on the scaffold derived from BAC and cosmid end sequencing, we estimate a final genome size between 18 and 20 Mb, similar to the genome of *C. merolae*. Our preliminary results (pulsed-field gels; genome sequence contigs) indicate that *G. sulphuraria* has several very small chromosomes (< 200 kB) and we believe this is one of the reasons why the genome size was underestimated previously. Monthly updates on the sequencing statistics can be found at: [http://galdi.bch.msu.edu/cgi-bin/galdieria/galdi2.pl](http://galdi.bch.msu.edu/cgi-bin/galdieria/galdi2.pl).

#### 3.2 Results from EST-sequencing and analysis
Two directionally cloned cDNA libraries were generated from photoautotrophically and heterotrophically grown *G. sulphuraria* cells, respectively, and 5,915 cDNA inserts were sequenced from their 5′-ends. 3,323 represented the autotrophic library, and 2,592 represented the heterotrophic library. Base calls were screened for vector and *E. coli* contamination, and quality values were assigned to each base using the phred algorithm 49,50. 5,270 sequence reads (89% of the total) passed the filter, and the final dataset consisted of 3,024 and 2,246 sequences generated from the autotrophic and heterotrophic libraries, respectively. The average phred Q20 40 read length was 563 bases. The 5,270 passing sequences were clustered using the stackPACK software package (South African National Bioinformatics Institute, University of the Witwatersrand).
Institute, University of the Western Cape, Republic of South Africa), resulting in 3,047 contigs (1.7 Mbp) of unique, non-redundant sequence.

Of these 3,047 contigs, 1,746 and 1,276 consisted of ESTs from only the autotrophic or heterotrophic libraries, respectively. Little overlap was found between both libraries, only 34 contigs contained sequence information derived from both libraries. This is most likely due to: (i) significantly different gene expression levels under both conditions, hence each library represents a snapshot of the expression profile under a specific culture condition rather than a comprehensive collection of cDNA clones. (ii) a relatively high “novelty rate” that is usually obtained during the early phases of random sequencing projects such as EST sequencing. 2,431 ESTs (80%) were unique sequences, whereas only 20% of the 3,047 contigs contained more than one EST sequence. This indicates that we did not reach saturation; hence only little overlap between both libraries is to be expected. For initial analysis of EST-sequences, we placed emphasis on the reconstruction of metabolic pathways. We found evidence for (i) a complete pathway for lipid A biosynthesis; (ii) export of triose-phosphates from rhodoplasts; (iii) and absence of eukaryotic hexokinases. A detailed description of the results was recently published.

### 3.3 Intron structure

A comparison of the *G. sulphuraria* genomic contigs to the EST dataset showed that more than 50% of all tested *Galdieria* genes contain introns. Based on 10 randomly selected genes containing one to three introns, intron lengths of 45 to 65 bases were found. The borders of the introns displayed typical spliceosomal features as previously described for a LHC gene from *G. sulphuraria*; in 80% of the cases, the splice donor site started with the sequence GU and the same percentage was observed for the splice acceptor site, which consisted of AG. In only 50% of the cases the branching site was composed of the consensus sequence CUPuAPy located 15 to 30 bases upstream of the acceptor site.

![Typical intron structure in G. sulphuraria](image)

**Figure 3:** Typical intron structure in G. sulphuraria. The splice donor and acceptor sites and the consensus branching site are indicated.

### 3.4 Current bioinformatics tools do not correctly predict gene models in *G. sulphuraria*

The *Galdieria* genome has several unusual features, such as very small introns (45 to 60 bp) and short intergenic regions. These specific features of the *Galdieria* genome complicate *ab initio* gene predictions using programs such as GenScan or GeneMark.hmm. In a pilot study using 10 experimentally verified full-length cDNA sequences, we found that none of the tested programs (GenScan, GlimmerR, FGENESH, GeneMark.hmm) precisely predicted more than 60% of gene structures. The best results (60% correct models) were obtained using GeneMark.hmm (trained on Arabidopsis, which has a relatively similar G + C contents). In particular the small introns (< 100 bp) proved to be problematic and frequently adjacent genes were sliced together, leading to gene fusions. Although the accuracy of gene predictions can be improved by combining several programs, or by training predictors with genomes showing similar characteristics, it becomes clear that reliable gene models require experimental support from cDNAs. In addition, cDNA sequences are required to build training sets for *ab initio* gene predictors to improve the quality of the computational predictions.

### 3.5 Mining the *G. sulphuraria* genome for membrane proteins

*G. sulphuraria* represents a particular interesting species for structural genomics and proteomics projects owing to its extraordinary metabolic versatility such as heterotrophic and mixotrophic growth on more than 50 different carbon sources. Moreover, it also tolerates high concentrations of toxic metals such as cadmium, mercury, aluminum.
or nickel. Hence, the genome of *G. sulphuraria* may provide membrane protein targets that are particularly interesting from a biotechnology standpoint, because of their thermostability. Eukaryotic cells are highly compartmentalized, allowing the parallel occurrence and efficient spatial separation of metabolic pathways that are competing for mutual precursors and metabolic intermediates. Examples are the tricarboxylic acid cycle in mitochondrial matrix, the glycosylation and sulphation of proteins in the endomembrane system, and the Calvin cycle in the chloroplasts stroma of plant cells. Many of the products that are synthesized in one compartment are further metabolized in other cellular compartments. Therefore, the functioning of eukaryotic cells strongly relies on a complex metabolic network that intertwines reactions in between intracellular compartments by the exchange of metabolic intermediates. The organellar membranes represent the interface between cellular compartments and contain numerous transport proteins that mediate the flux of metabolites and ions across these membrane(s).

Many intracellular transporters do not have related proteins in prokaryotic organisms. In particular, solute transporters of the mitochondrial membrane, the Golgi/ER-endomembrane system, and of the plastid inner envelope membrane are confined to eukaryotic organisms. In contrast to intracellular transport processes that are exclusive to eukaryotic cells, the transport of solutes across the plasma membrane is crucial for both eukaryotic and prokaryotic organisms and many of the corresponding transport systems can be identified in both kingdoms.

The ongoing genome project for *G. sulphuraria* ([http://genomics.msu.edu/galdieria](http://genomics.msu.edu/galdieria)) and the completely sequenced genome of *C. merolae* ([http://merolae.biol.s.u-tokyo.ac.jp/](http://merolae.biol.s.u-tokyo.ac.jp/)) offer for the first time unique possibilities for a global analysis of the *in silicio* predicted proteomes of thermophilic eukaryotes as a whole or as a subset of proteins, e.g., membrane proteins. Transmembrane proteins contain in general from one to over 14 transmembrane spans in the form of α-helices. There are also β-barrel proteins with an even number of β-strands found in the outer membrane of bacteria, mitochondria, and plastids.

The predicted number of transmembrane proteins present in the genome of *C. merolae* amounts to approximately 1,250 proteins containing one ore more transmembrane spans, corresponding to 22% of the 6,390 proteins encoded by its genome. Based on a preliminary genome annotation for *G. sulphuraria*, the genome of this alga encodes a higher portion of membrane proteins, as is to be expected because of its ability to take up external carbon sources.

Many membrane proteins from Cyanidiales can be classified by the transporter classification (TC) system, a functional/phylogenetic system designed for the classification of all transmembrane transport proteins down to the transporter family level (third digit of the TC system). With few exceptions, at least one member of a particular TC-family can be identified in either *G. sulphuraria* or *C. merolae*. In an initial survey, 122 *C. merolae* membrane proteins and 133 *G. sulphuraria* membrane proteins have been classified by the TC system (data not shown), clearly demonstrating that members of many TC-families are represented in thermophilic Cyanidiales genomes. These unicellular microalgae hence represent a rich source of genes encoding potentially thermo-stable membrane transporters that are highly interesting targets for structural studies. Of particular interest are the many members of the mitochondrial transporter superfamily (57 genes), of the organic cation transporter family (16 genes), sugar porter family (25 genes), the ATP-binding cassette (ABC) transporter family (32 genes), and the drug-metabolite transporter (DMT) family (17 genes).

Below we provide a brief summary of our initial survey of the *Galdieria* genome for membrane proteins that are highly homologous to mammalian and bacterial membrane proteins and that we consider interesting targets for functional and structural studies.

### 3.5.1 Membrane-bound glycosyltransferases

Many of the important glycosyltransferases involved in membrane biogenesis are integral membrane proteins or interfacial proteins. Several important families of glycosyltransferases catalyze the formation of monoglycosyl-sterols, monoglycosyl-ceramide, and glycosylated diacylglycerol, a variety of glycolipids and second messengers. Several genes of *Galdieria* encode proteins that are highly homologous to mammalian (dolichyl-phosphate mannosyltransferase and O-linked N-acetylglucosamine transferase), plant (1,2-diaclyglycerol 3-beta-galactosyltransferase) and bacterial (lipid A disaccharide synthase) enzymes. This work has implications for the development of antibacterial and antifungal agents.

### 3.5.2 Transporters (active, symport, antiport, etc)

We have identified many genes encoding solute transporters in *Galdieria* and several of these targets are being cloned for expression studies. One particular interesting target is...
the plastidic ADP/ATP translocator. It is unrelated to the mitochondrial ADP/ATP carrier, but exhibits significant homology to the plastid nucleotide translocators from higher plants and to the plastidic ADP/ATP translocator from Chlamydia, a major human and animal pathogen.

In addition, we have identified several putative E1-E2 ATPases in Galdieria that are homologous to chromaffin granule ATPase II, which is involved in phospholipid transport. From the very large family of E1-E2 ATPases, only the structure of Ca\(^{2+}\)-ATPase has been determined and there are still many unanswered questions about the structure and function of this critically important family of pumps.

3.5.3 Membrane Proteins involved in Organellar Biogenesis

Within the genome of Galdieria, several genes have been identified that are homologous to mammalian genes whose products are involved in cell and organelle division. Genetic defects of cell and organelle division are quite deleterious and defects in mitochondrial division and fusion is associated with a diverse set of human diseases. We have already identified a gene that is closely homologous to the mitochondrial fission protein Dlp1. The existence of the dynamin-related protein Dlp1/Dnm1 suggests that Galdieria abandoned a bacterial-like mitochondrial division process involving FtsZ shortly after the symbiotic fusion that gave rise to early eukaryotes like Galdieria. We have also identified another gene that is closely homologous to a mitochondrial FtsH-like AAA metalloprotease; the gene in Galdieria also displays surprisingly high sequence homology (E-value = 1e\(^{-107}\)) to the transitional endoplasmic reticulum ATPase from humans. The structural analysis of these proteins from humans, plants, and an early eukaryote like Galdieria would not only shed light on the evolution of organelle biogenesis in eukaryotes, but also provide a better understanding of the impact of organelle defects on cell physiology.

3.5.4 Phosphate translocators and other members of the Drug/Metabolite Transporter (DMT) Superfamily

Plastidic phosphate translocators (PTs) form a sub-group of the DMT-family that cannot be assigned with the typical six or twelve helix topologies that are found in many other membrane transporter families. To date, only one structure of a member of the DMT-family (EmrE) has been solved at 3.8A resolution and this protein has four transmembrane domains. The by far best-studied members of the DMT-family are the phosphate translocators of plant cell plastids. Similar to many other proteins of the DMT-family, these proteins have most likely 8 to 10 transmembrane domains. Plant phosphate translocators are related to a large family of membrane proteins in plants, animals, and fungi that all belong to the DMT-family. The structure of plant phosphate translocators might thus be considered representative for a large number of intracellular transport proteins. In contrast to most other members of the DMT superfamily, pPTs are functionally characterized, and, importantly, heterologous expression in yeast, affinity purification of recombinant transporter protein to apparent homogeneity by metal-affinity chromatography, and functional assay of transporter activity after reconstitution into liposomes has been established for all members of this protein family.

Similar expression and reconstitution systems have been used for structurally and phylogenetically unrelated dicarboxylate transporters of the plastid envelope membrane, indicating that this experimental strategy is potentially applicable to plastid envelope membrane proteins in general. The triose-phosphate/phosphate translocator is the dominant protein of the chloroplast membrane, representing up to 15% of the total membrane protein, suggesting that high amounts of these transporter proteins can accumulate in the native membrane environment. A highly specific inhibitor for phosphate translocators is available that binds in a 1:1 stoichiometry to a lysyl-residue in the active site of native protein, indicating this inhibitor could be used to rigidify the structure of the transport protein during crystallization trials.

It has been previously demonstrated that UDP-galactose-4-epimerase from G. sulphuraria is an unusually thermostable enzyme that was not inactivated even after incubation at 46°C for several hours. It is tempting to hypothesize that this also applies to membrane proteins of these thermophilic organisms. A survey of the genome sequences of G. sulphuraria and C. merolae revealed the presence of small gene families encoding PT-related proteins in both thermophilic algae.

In a preliminary study, we have tested whether the expression and reconstitution systems established for phosphate translocators from mesophilic plants can be applied to orthologous proteins from G. sulphuraria. The recombinant protein could be expressed at very high yields (not shown). In addition, the recombinant protein was active after reconstitution into liposomes, i.e., we were able to detect high rates of phosphate uptake into liposomes reconstituted with recombinant translocator protein. Figure 4 shows that recombinant, reconstituted phosphate translocator of the thermophilic algae G. sulphuraria 4-fold higher activity at 40°C in comparison to 20°C, indicating reduced conformational flexibility at room temperature and higher thermal stability of the transport protein, a desirable feature for structural studies.
3.6 Pathway of carbon export from red algal chloroplasts

Chloroplasts are the sites of photosynthetic carbon assimilation in plant cells. The net product of carbon assimilation by the reductive pentose phosphate pathway, triose phosphate, serves as principle precursor for all biosynthetic reactions in plants. In green plants, recently assimilated carbon is shuttled into starch and sucrose biosynthesis, nitrogen and sulfur metabolism, fatty acid biosynthesis, cell wall biosynthesis, secondary metabolism, and a puzzling variety of other metabolic routes. In red algae, the predominant soluble carbohydrate is floridoside (α-D-galactopyranosyl-(1,2)-glycerol), not sucrose, and in contrast to green plants, red algae store starch in the cytosol, not inside the chloroplast. A key question in plant physiology is how the allocation of carbon to the individual pathways is regulated. The initial step in the allocation of recently assimilated carbon to different metabolic routes is the partitioning of triose phosphates between the plastid stroma and the plant cell cytosol. In addition, reduced carbon can be withdrawn from the regenerative phase of the Calvin Cycle, for example to fuel the plastid-localized shikimic acid pathway. In green plants, carbohydrates that are synthesized in chloroplasts by the reductive pentose phosphate pathway during the day are mainly exported to the cytosol as triose phosphate by the triose phosphate/phosphate translocator (TPT)\textsuperscript{85-87}. The predominant fate of triose phosphate in the cytosol is the conversion to sucrose, which serves as the transport form of photoassimilates that are allocated to sink tissues. We asked the question as to whether a similar pathway for the export of carbohydrates from chloroplasts exists in red algae. A survey of the \textit{G. sulphuraria} and \textit{C. merolae} genomes indicated that the genomes of both red algae harbor genes that encode proteins that show significant homologies to plastidic phosphate translocators from green plants\textsuperscript{42}. Liposomes reconstituted with total \textit{G. sulphuraria} membrane proteins showed the signature phosphate/phosphate antiport activity characteristic for plastidic phosphate translocators and we also could detect triosephosphate/phosphate counter-exchange\textsuperscript{42}. We thus propose that the export of recently assimilated carbon from red algal chloroplasts occurs by a similar counter-exchange mechanism as in green plant chloroplasts and that a protein similar to plastidic triosephosphate/phosphate translocators from green plants catalyzes this counter-exchange (see Figure 5).

Figure 4: Temperature dependency of phosphate uptake into liposomes that had been reconstituted with a plastidic phosphate translocators from \textit{G. sulphuraria} and preloaded with 20 mM unlabeled ortho-phosphate. The rate of phosphate uptake was found to be 4-fold higher at 40°C as compared to 20°C.

Figure 5: Proposed model for carbon export from red algal chloroplasts. CO\textsubscript{2} is assimilated by the Calvin Cycle in the red algal chloroplast (Rhodoplast), yielding reduced organic carbon in the form of triose-phosphate (Triose-P). Triose-P is exported to the cytosol in strict 1:1 counter-exchange with inorganic phosphate (Pi), thus maintaining the phosphate homeostasis of the plastid stroma. Once in the cytosol, Triose-P can be converted to Floridean Starch via UDP-Glucose (UDP-Glc) or to the soluble galactosyl-glycerol Floridoside via Glycerol-3-P and UDP-galactose (UDP-Gal). For clarity, the biosynthetic pathways have been simplified and abbreviated.
The genomes of *G. sulphuraria* and *C. merolae* do not encode plastidic maltose or glucose transporters similar to those found in green plants \(^{43,88-91}\). In green plants, the glucose and the maltose transporter are involved in the export of starch breakdown products from the chloroplast to the cytosol at night \(^{90,91}\). Since starch is stored in the cytosol and not inside the plastid stroma in red algae, these transporters have been either lost early during the evolution of the red algae, or they have been acquired by green plants after the split of the plant lineage into chlorophytes and rhodophytes.

### 3.7 Concluding remarks

Due to space constraints, this article provides only a brief survey of the most recent findings obtained in the *G. sulphuraria* Genome Sequencing Program. The preliminary analysis of the genome sequence indicates that membrane transporters are likely essential for the metabolic versatility and the stress resistance of *G. sulphuraria*. Functional genomics studies such as genome-wide transcript profiling and the establishment of mutants and gene transfer protocols will be required to further our understanding of the biology of this amazing organism. To better understand the metabolism of this extremophilic alga, detailed analyses of the metabolome need to be conducted under various culture conditions. For functional studies of *G. sulphuraria* genes and to establish reliable gene models, we will need a comprehensive collection of full-length cDNAs, representing most, if not all *Galdieria* genes.

### 4. ACKNOWLEDGEMENTS

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### 5. REFERENCES


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