Structural, Biochemical and Genetic Characterization of Dissimilatory ATP Sulfurylase from Allochromatium vinosum

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Abstract

ATP sulfurylase (ATPS) catalyzes a key reaction in the global sulfur cycle by reversibly converting inorganic sulfate (SO₄²⁻) with ATP to adenosine 5'-phosphosulfate (APS) and pyrophosphate (PPi). In this work we report on the sat encoded dissimilatory ATP sulfurylase from the sulfur-oxidizing purple sulfur bacterium Allochromatium vinosum. In this organism, the sat gene is located in one operon and co-transcribed with the aprMBA genes for membrane-bound APS reductase. Like APS reductase, Sat is dispensable for growth on reduced sulfur compounds due to the presence of an alternate, so far unidentified sulfite-oxidizing pathway in A. vinosum. Sulfate assimilation also proceeds independently of Sat by a separate pathway involving a cysDN-encoded assimilatory ATP sulfurylase. We produced the purple bacterial sat-encoded ATP sulfurylase as a recombinant protein in E. coli, determined crucial kinetic parameters and obtained a crystal structure in an open state with a ligand-free active site. By comparison with several known structures of the ATPS-APS complex in the closed state a scenario about substrate-induced conformational changes was worked out. Despite different kinetic properties ATPS involved in sulfur-oxidizing and sulfate-reducing processes are not distinguishable on a structural level presumably due to the interference between functional and evolutionary processes.

Introduction

Sulfur compounds are used by a huge variety of organisms for the biosynthesis of sulfur-containing amino acids, cofactors and metabolites [1] and for energy conservation serving either as electron donors or acceptors [2]. Sulfur occurs in the biosphere in oxidation states -II to +VI, mostly in form of hydrogen sulfide, elemental sulfur and sulfate. The dissimilatory sulfur-oxidizing and sulfate-reducing processes share several key enzymes, namely ATP sulfurylase (ATPS), adenosine-5-phosphosulfate reductase (APSR) and sulfate reductase (Sir) [2,3].

In the sulfur-oxidizing pathway ATP sulfurylase (adenylsulfurylase/ATPS:sulfate adenylyltransferase; E.C. 2.7.7.4) catalyzes the final reaction of the oxidation of reduced sulfur compounds by reversibly transferring pyrophosphate (PPi) onto adenosine-5'-phosphosulfate (APS) to form inorganic sulfate (SO₄²⁻) and ATP [4,5,6]: MgPPi+APS ⇌ MgATP+SO₄²⁻. In addition, the ATPS reaction provides a major route for recycling PPI produced by biosynthetic reactions.

In the sulfate-reducing pathway ATPS catalyzes the adenylation of SO₄²⁻ with ATP to APS and PPi. APS is used as the activated form of sulfate in the dissimilatory process and in the assimilatory process of plants, algae and most bacteria whereas phosphoadenosine-3'-phosphosulfate (PAPS) generated from APS by phosphorylation exerts this function in the assimilatory pathway of some organisms such as fungi and some bacteria including most cyanobacteria [2,7,8]. In addition, PAPS serves as the sulfuryl donor for the formation of sulfate esters by sulfotransferases.

Except for a number of bacteria containing a ATPS consisting of four heterodimers (CysDN) for assimilatory sulfate reduction [8,9], a common fold for catalyzing the ATPS reaction occurs for the remaining organisms. In yeast [10], filamentous fungi [11], and bacterial species the assimilatory sulfate-reducing ATPS have been described as homohexamers [12] and in plants as homotetramers [13] of 41–69 kDa-subunits. In some enzymes an APS kinase domain is C-terminally fused to ATPS [8,14]. The APS kinase domain can be enzymatically active as in Aquifex aeolicus [15], possess a regulatory function by binding the allosteric inhibitor PAPS as in filamentous fungi [11] or have no defined function as in yeast [10]. In higher eukaryotes, like in mammalian species [16] or in metazoan organisms, APS kinase is N-terminally fused to ATPS. Sulfate-reducing dissimilatory ATPS is present as a homotrimer with one Zn ion bound to each ATPS subunit [17]. Dissimilatory sulfur-oxidizing ATPS was found to exist as a homodimer [12].

The dissimilatory oxidation of reduced inorganic sulfur compounds is linked to energy transformations via photosynthesis or respiratory processes. Sulfur oxidizers are found among the Archaea and the Bacteria and comprise photo- and chemolithotrophs. Dissimilatory sulfur oxidation in Eukarya is mediated by...
lithotrophic bacterial endosymbionts. In the phototrophic purple sulfur bacterium *Allochromatium vinosum* two types of ATPs where identified on the basis of DNA sequence analysis and by inspection of the complete genome sequence [18]. The genes *cysDV* (Alvin_2448 and Alvin_2449) encode for ATPS in an unusual sulfate assimilation pathway [19] and the gene *sat* (Alvin_1118) for an ATPS involved in the dissimilatory sulfate oxidation pathway, respectively. We have overproduced ATPS of *A. vinosum*, characterized the enzyme kinetically, determined its structure of the *A. vinosum* enzyme in an open state and analyzed the conformational rearrangement upon APS and PPi binding.

**Results and Discussion**

The *sat-aprMBA* Gene Locus in *A. vinosum* and other Phototrophic Members of the Family Chromatiaceae

In *A. vinosum* the *sat* gene encoding ATP sulfurylase (Alvin_1118) is located immediately upstream of the *aprMBA* genes encoding membrane-bound APS reductase (Alvin_1119–1121) [3,18]. AprM is predicted to contain five transmembrane helices with no sequence similarity to any currently known conserved domain or cofactor binding site in the databases. An essential function of AprM as a membrane anchor that allows spatial and functional association of this type of oxidative APS reductase with the membrane has been postulated and it has been suggested that AprM serves as an entry point into the membrane for the electrons released during formation of APS from sulfite and AMP [20]. In the currently available complete genome sequences of phototrophic members of the family Chromatiaceae, the same gene arrangement is present in *Thiorhodobacter* sp. 970. In *Thiothrix marina* DSM 5811 (DSM 56555), *Thiorhodobacter disorderi* AZ1 (DSM 150065) and *Thiolfiavivococcus mobilis* DSM 8321T sat and aprMBA are not linked on the chromosome. The occurrence of *aprMBA* has also been reported for *Thiococcus* pfennigi 4520 [21] while *Thiocytes violacea* DSM 198T encodes Sat, AprBA and QmoABC, each in separate loci. The QmoABC complex was first identified in the dissimilatory sulfate reducing bacterium *Desulfobium desulfuricans* [22]. The complex consists of one membrane (QmoC) and two cytoplasmic subunits (QmoAB). The two QmoC hemes b are reduced by quinols and experimental evidence strongly indicates that the Qmo complex participates in electron flow between the quinone pool and the cytoplasm, i.e. that it acts as the electron-donating unit for APS reductase in sulfate reducers [23,24]. The qmoABC genes are not only present in sulfate-reducing prokaryotes [23] but occur also in many chemotrophic sulfur-oxidizing bacteria as well as in green sulfur bacteria [24,25] and in one further purple sulfur bacterium (*Thiodictyon* sp. Cad16 [21]). In sulfur oxidizers, QmoABC is thought to act as electron acceptor for the electrons released during formation of APS and would thus have a function analogous to that of AprM. It should be noted that purple sulfur bacteria do not use a single mechanism to oxidize sulfite which is apparent from our former findings that APS reductase is not essential for sulfite oxidation in *Allochromatium vinosum* [26,27] and also from the observation that the complete genome of *Marichromatium purpuratum* 984 (DSM 15914T) neither contains sat nor *aprMBA* genes and *apr* genes are also not present in *Isochromatium* [20].

Insertional Inactivation of the *sat* Gene in *A. vinosum*

Experimental evidence for the *sat-aprMBA* genes forming a transcriptional unit was obtained by introducing a kanamycin-*Ω* cassette into the *sat* gene resulting not only in a nearly complete loss of ATP sulfurylase activity (in extracts of cells grown photolithothoautotrophically on sulfide for the wildtype 0.3 U (mg protein)⁻¹ and for the mutant 0.001 U (mg protein)⁻¹; activities were determined by using the assay in the APS synthesis direction as outlined in the “Material and methods” section) but also APS reductase activity in crude extracts of the purple sulfur bacterium. The *A. vinosum* sat::*km* mutant strain was still able to grow photolithothoautotrophically on sulfide and also photoorganoheterotrophically on sulfate as the sole sulfur source. These findings imply that the *sat*-encoded enzyme is neither essential for sulfate assimilation nor for dissipatory sulfate oxidation in this organism. These results are corroborated by an earlier report that the ability to indirectly oxidize sulfite via APS is not required for *A. vinosum* [26] and that a dedicated pathway exists for sulfate assimilation [19].

Kinetic Characterization of Recombinant *A. vinosum* ATP sulfurylase

The recombinant *A. vinosum* ATP sulfurylase produced in *E. coli* was tested for enzymatic activity in vitro and some crucial kinetic parameters were determined (Fig. 1). The enzyme displayed hyperbolic *v* versus *p* plots (Fig. 1) and normal (linear) reciprocal plots. Our measurements showed a *V*ₘₐₓ in the direction of ATP synthesis of 433±18 U (mg protein)⁻¹ at 30°C, pH 8.0 and saturating substrate concentrations (1 mM PPi, 0.2 mM APS). The observed specific activity is in the range of that reported for “*Candidatus Endorhiza Persephone*” [12], the sulfur oxidizing symbiont of *Riftia pachyptila*. *Kₐ* values for APS and pyrophosphate were determined to be 9.5 μM and 47.6 μM, respectively. In the molybdoysis assay *V*ₘₐₓ was 72.3±4.1 U (mg protein)⁻¹, *Kₐ* values for ATP and molybdate were determined to be 0.8 mM for ATP and 2.5 mM for molybdate. A much higher specific activity in the direction of ATP production from APS and PPi, than in the molybdoysis assay has also been observed for “*Candidatus E. persephone*” ATPS while this ratio is generally smaller in assimilatory ATPS indicating a special adaption of the enzymes from sulfur oxidizers for working effectively in the ATP synthesis direction.

Structure Overview

The X-ray structure of ATPS from *A. vinosum* overproduced in *E. coli* was refined to R/R free factors of 17.9%/20.2%. The diffraction data were incompletely collected to 1.6 Å (38% completeness in the highest resolution shell), but at 1.8 Å resolution the local completeness is 92.3% and the accumulated completeness is 94.7% (see Table 1 for further data statistics). However, the inclusion of the incomplete higher-resolution data in the refinement improved the quality of the electron density maps and was essential to our structure analysis. The asymmetric unit contains one homodimer of ATPS with rms differences between the monomers being 1.4 Å. No significant differences were detectable between them except for a rigid-body movement of the C-terminal relative to the other domains in the range of 4 Å and of segment 233:239 which contacts the C-terminal domain. As described previously for ATPS structures from other organisms [28] each subunit of *A. vinosum* ATPS is subdivided into three domains (Fig. 2). The N-terminal domain I (residues 1–172) and the small C-terminal domain III (residues 332–396) of a small three-strand β-sheet and two α-helices. As mentioned, ATPS structures of various assimilatory sulfate reducing organisms such as fungi, yeast and *Aquifex aeolicus* [14,28,29] contain a fourth APS kinase-like domain. The structure of ATPS of *A. vinosum* was determined without any ligand in the active site. Two MES (2N-morpholino-ethansulfonic acid)
acid) molecules (present in the crystallization solution) could be, however, identified close to the monomer-monomer interface far away from the active site (Fig. 2). They are primarily linked to the polypeptide by hydrogen bonds between the oxygen of its morpholino ring and a guanidino group of Arg13 and between two sulfate oxygens and the amide nitrogen of Asp17.

The Substrate Binding Process

The substrate binding and active sites are positioned in a deep groove above the C-terminal ends of the central \( \beta \)-sheet of domain II; its walls are composed of the loops following strands 8 (195:199), 9 (224:230), 10 (260:266), 11 (291:295), and 12 (326:329) as well as of the N-terminal side of helix 12 (361:370) (Fig. 3). The characteristic RNP (199QXRN202) and GRD (295GRD297) loops follow the straddled strands 8 (195:199) and 11 (291:295). ATPS has been structurally analyzed from several organisms in various active site ligation states. In the \textit{A. vinosum} structure the active site is only occupied by water molecules; more than ten of them are visible in the electron density map. Substrate-free \textit{Saccharomyces cerevisiae}, \textit{''Candidatus E. persephone''} and human ATPS structures contained phosphate, sulfate or chloride molecules attracted by patches of positively charged residues of the active site groove (Fig. S2) [28,30,31]. The \textit{S. cerevisiae}, \textit{Penicillium chrysogenum}, \textit{Aquifex aeolicus} and \textit{Thermus thermophilus} ATPS structures were determined in complex with APS [14,28,29,32] and yeast ATPS with APS (or ATP analogues), pyrophosphate and chromate [33]. On this structural basis a scenario for the induced-fit process as a result of substrate binding is proposed. The catalytic process starts by binding APS into the empty active site groove best expressed by the \textit{A. vinosum} ATPS structure present in an open form (Fig. 3). Docking experiments using the program AutoDock [34] revealed that the substrate APS also binds with a significant affinity to the enzyme conformation of the substrate-free state (Fig. S1). Accordingly, APS is attached along the C-terminal end of the central \( \beta \)-sheet of domain II in an L-shaped conformation guided by the positive surface potential.

Initial APS binding activates an induced-fit process resulting in a net shrinkage of the groove from an open to a more closed form observable when comparing ligand-free and APS bound ATPS structures [30,28] (Fig. 3). The interactions between the polypeptide and APS are significantly increased during the induced-fit process which might be necessary to fix APS in the found compressed L-shaped conformation. Attracted by APS, the RNP and GRD loops move 2–3 Å and 1–2 Å, respectively, to form hydrogen bonds between the sulfate oxygens of APS and Gln199-Ne2H, Arg201-NH2 and Ala299-NH, between the \( \alpha \)-phosphate oxygens and Thr200-Oc1, Arg201-NH, Asn202-NH and Asn202-Nd2H2 and between the adenosine ribose hydroxo groups and Gly295-NH, Arg296-O and His298-Nd1. The induced-fit movement is propagated to segments contacting the RNP and GRD loops including the loops that follows strands 9 (224:230) and 10 (260:266), helix 9 (240:254) and the preceding 231GxxKxxD237 loop, helix 13 (381:393) and finally to the entire C-terminal domain (Fig. 3).

The conserved GxxKxD loop after strand 9 (224:230) is in an “up” position and highly mobile in the open \textit{A. vinosum} and “\textit{Candidatus E. persephone}” ATPS. Its mobility is expressed by its increased temperature factor (32.4 Å2; overall 14.2 Å2) and high conformational flexibility of up to 2.5 Å calculated between the two open ATPS structures (Fig. 4). Upon APS binding this loop is shifted 4–9 Å down towards the groove and forms together with the adjacent GRD loop above the phosphosulfate binding site a

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**Figure 1. Kinetics of recombinant ATP sulfurylase from \textit{A. vinosum}.** (A) \( v \) versus [APS] in the ATP synthesis reaction at 1 mM pyrophosphate. (B) \( v \) versus [PPi] at 0.2 mM APS. (C) \( v \) versus [MgATP] in the molybdolysis reaction at 50 mM MoO\(_4^{2-}\). (D) \( v \) versus [molybdate] in the molybdolysis reaction at 10 mM MgATP. Each data point is the mean ± standard deviation of three assays on the same batch of protein but in some cases the error bars are too small to be seen. The solid lines through the data points are the fits to the Michaelis-Menten equation using non-linear regression as described in Materials and Methods.

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kinetic data that indicated PPi binding only in the presence of APS side of helix 12 (361:370) [28]. This finding can be correlated with upon APS binding in the closed form in front of the N-terminal interferes with the pyrophosphate binding site which is created and thus fixes a more closed form. sulfate group of APS, is sandwiched between Gln199 and Ala299 prevented presumably because a sulfate, bound instead of the ATPS structure the described close/open transition is largely induced conformational changes of three of these residues which reflect to a certain extent the induced-fit process upon pyrophosphate binding. His205 is hydrogen-bonded in the A. vinose ATPS with Ser379-Oy1H and via solvent molecules with Tyr391-OyH and swings towards the active site upon sulfate and presumably also upon APS/PP binding (Fig. 4). Likewise, Asn202 is hydrogen-bonded with Asp237 in the A. vinose ATPS structure and moves towards the substrates upon their binding. The guanidinium group of Arg365 [part of the conserved 99SGTxxRX365 motif] present in two conformations dominantly points towards Glu366 in A. vinose ATPS and swings more than 6 Å towards the sulfate in the “Candidatus E. persephone” ATPS structure (Fig. 4).

Structural and Sequence Comparisons within the ATP Sulfurylase Family

Sequence comparison studies revealed a high degree of similarity among the ATPS family members (Fig. 5), in particular, for the catalytic domain II which is confirmed by the X-ray structures. The rms deviation between A. vinose, “Candidatus E. persephone” [30], yeast [28], fungus [14], human [31], T. thermophilus [32], and Aquifex aeolicus ATPS [29] ranges between 0.8–2.9% (more than 87% of Cα positions used) [38]. Considering that the structures are determined in different ligation states that are accompanied by the described large-scale conformational changes the calculated rms values are strongly underestimated. Equivalent enzymatic states would result in rms deviations below 1 Å for residues of domain II, in particular, for those regions involved in the catalytic reaction.

ATPS functions in three different metabolic contexts. In the widely distributed assimilatory sulfate reduction pathway sulfate is recruited for amino acid and cofactor biosynthesis whereas in the dissimilatory sulfate reduction and sulfide oxidation pathways a limited number of microorganisms consume sulfate and reduced sulfur compounds, respectively, in large amounts because these compounds act as electron donor/acceptors in energy conversion processes. A careful sequence and structural analysis indicates that the RNP and GRD loops and other crucial residues of the groove are strictly conserved except for the six residues longer GRD loop in some assimilatory ATPS that, however, does not directly participate in phosphosulfate binding (Fig. 5). Nevertheless, kinetic data suggest distinct substrate binding and catalysis: APS and PP, synthesis is favoured in sulfate-reducing ATPS sulfonylases while the enzymes from sulfur oxidizers appear more adapted to catalyzing ATP and SO42− synthesis [15]. The structural basis for these differences is, however, intricate. Most attractive locations for differences include the irregular region preceding helix 12 (361:370) of domain III and the GxxKxxD region, both being only moderately conserved (Figs. 2+3). Minor perturbations of their conformations modify the size of the APS and PP, binding sites and the dynamics of the induced-fit process.

Table 1. Data collection and refinement statistics.

| Data collection | | |
|----------------|----------------|
| Detector type  | ADSC Quantum 4 ccd |
| X-ray wavelength (Å) | 0.9393 |
| Space group | P21 |
| Unit cell parameters (Å) | a = 73.3, b = 97.0, c = 73.5 (°) |
| Resolution range (Å) | 1.6–1.98 (1.66–1.60) |
| No. of observed reflections | 98813 (4546) |
| Completeness (%) | 83.4 (38.6) |
| Multiplicity | 2.2 (1.4) |
| Rmerge (%) | 15.8 (2.1) |
| Rfree (%) | 6.0 (29.9) |

Model refinement

| Resolution range (Å) | 1.6–1.98 (1.65–1.60) |
| Rfactor/Refe (%) | 17.9/20.2 (31.8/36.6) |
| Overall B factor (Å²) | 35.3 |
| Protein | 35.3 |
| Water | 3.0 |
| R.m.s deviations from ideal geometry | |
| Bond lengths (Å) | 0.017 |
| Bond angles (°) | 1.65 |
| Ramachandran plot | |
| Most favorable (%) | 99.0 |
| Allowed (%) | 0.75 |
| Generously allowed (%) | 0.25 |
| Disallowed (%) | 0.0 |
| PDB code | 4DNX |

Due to the low completeness of the data in the highest resolution shell, the effective resolution is lower than 1.6 Å.

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shield that is absent in the open form (Fig. 3). The up-to-down movement of the GxxKxxD loop is a consequence of the APS induced conformational change of the RNP loop allowing the formation of a hydrogen bond interaction between Asp237 and Arg201 and between Ile238, a solvent molecule and Arg201 and Asn202. Thus, the GxxKxxD loop indirectly participates in APS binding, as already recognized by analyzing the “Candidatus E. persephone” ATPS functions in three different metabolic contexts. In the widely distributed assimilatory sulfate reduction pathway sulfur is recruited for amino acid and cofactor biosynthesis whereas in the dissimilatory sulfate reduction and sulfide oxidation pathways a limited number of microorganisms consume sulfate and reduced sulfur compounds, respectively, in large amounts because these compounds act as electron donor/acceptors in energy conversion processes. A careful sequence and structural analysis indicates that the RNP and GRD loops and other crucial residues of the groove are strictly conserved except for the six residues longer GRD loop in some assimilatory ATPS that, however, does not directly participate in phosphosulfate binding (Fig. 5). Nevertheless, kinetic data suggest distinct substrate binding and catalysis: APS and PP, synthesis is favoured in sulfate-reducing ATPS sulfonylases while the enzymes from sulfur oxidizers appear more adapted to catalyzing ATP and SO42− synthesis [15]. The structural basis for these differences is, however, intricate. Most attractive locations for differences include the irregular region preceding helix 12 (361:370) of domain III and the GxxKxxD region, both being only moderately conserved (Figs. 2+3). Minor perturbations of their conformations modify the size of the APS and PP, binding sites and the dynamics of the induced-fit process.
Besides the described common factors concerning substrate binding ATPS also reveals differences that more indirectly influence the active site groove but may fine-tune the biochemical...
reaction. First, sulfur-oxidizing ATPS are present as homodimers, dissimilatory sulfate-reducing ATPS as homotrimers and assimilatory sulfate-reducing ATPS frequently as homotetramers or -hexamers implicating a specific rigidification and stabilization of regions of the active-site groove. First, in *A. vinosum* ATPS the GxxKxxD loop is in direct contact to the partner monomer. Second, assimilatory ATP sulfurylases frequently contain a fourth APS-kinase like domain [28,14,29] that represents an essential component of the stable hexamer and influences, in parallel, the ATPS reaction by contacting helix 12 (361:370) involved in PPi binding. Third, plant ATPS (see Fig. 5 *Arabidopsis thaliana* and *Zea mays* domain I) possesses a more than 50 amino acids longer N-terminal arm that might be involved in oligomeric interactions. Fourth, domain III of ATPS from dissimilatory sulfate reducing microbes and from various sulfur-oxidizing and assimilatory sulfate-reducing organisms contains a characteristic zinc-binding site [32,17]. The zinc ion is tetrahedrally coordinated by three cysteines and one histidine and is centrally positioned in the solvent-exposed segment that links strand 12 (326:329) and helix 12 (361:370) involved in adenine and PPi binding, respectively. The zinc-binding motif preferably found in ATPS of thermophilic organisms appears to be important for active site stabilization

![Figure 5. Sequence alignment of ATP sulfurylases from sulfur oxidizing and sulfate assimilating and dissimilating organisms.](image)

Residues of the highly conserved RNP and GRD motif and the mobile loop are indicated by asterisks and a blue stripe, respectively. The three cysteines and one histidine of the conserved zinc-binding motif are marked with red boxes. The secondary structure symbols are illustrated according the ATPS structure of *A. vinosum*. The alignment figure was made using the programs ClustalX [50] and ESPript [51]. Species: *Allochromatium vinosum* DSM 180 T (A_vinosum, Alvin_1117, ADC62057), “*Candidatus Endoriftia Persephone*” (R_symbiont, PDB: 1JHD_A), *Aquifex aeolicus* VF5 (A_aeolicus, PDB: 2GKS_A), *Saccharomyces cerevisiae* (S_cerevisiae, Met3p, AAU09752), *Penicillium chrysogenum* (P_chrysogenum, MET3_PENCH, Q12650), *Arabidopsis thaliana* (Arabidopsis_thaliana, AAA21570), *Zea mays* (Z_mays, NP_001104877), *Homo sapiens* (H_sapiens, PDB: 2QJF_A), *Thermus thermophilus* (T_thermophilus, YP_004282.1), *Archaeoglobus fulgidus* (A_fulgidus, SAT_ARCFU, SP: O28606.2), *Desulfovibrio desulfuricans* ATCC 27774 (D_desulfuricans, YP_002479044.1). [doi:10.1371/journal.pone.0074707.g005](#)
when the lack of a APS kinase domain does not allow stabilization by homoxamer formation.

The virtually identical substrate binding site of ATPS - independent of their metabolic functions – arises from its origin very early in evolution prior to the divergence of the three domains of life and from the complex chemical reaction that did not allow substantial variations. Separable classes of ATPS according to their metabolic function are not definable despite the described structural differences and distinguishable kinetic data for the forward and backward reaction. Obviously, subtle adjustments are superimposed by the normal phylogeny of the organisms including the adaptation to specific environmental conditions. One apparent example represents the characteristic zinc-binding motif preferably found in thermophilic microorganisms (Fig. 5). However, a more profound sequence analysis that integrates gene duplication and lateral gene transfer events and a broader functional characterization of individual ATPS is required to explain observations as for example that Aquifex aeolicus ATPS is more related to S. cerevisiae and P. yezuyamae ATPS than to most bacterial ATPS or that the plant enzyme is more similar to human ATPS than to the respective protein from green algae [8].

Materials and Methods

Construction of sat-deficient Allochromatium vinosum

A clone carrying a 3-kb BglII restriction fragment with the complete sat gene (pAW307) was isolated from a library of 2.5- to 3.5-kb BglII restriction fragments of total A. vinosum DNA in the pGEM-3Zf(+) vector using a PCR generated 300 bp sat probe (primers used: aw1 CAGAC/G/TCCG/C/TAC/G/CATGCA and aw2 TCGA/G/ACC/G/ACC/G/CATGCA). The plasmid was digested with EcoRI, blunt-ended and digested with SfiI. This led to plasmid pAW201 which was used to introduce the blunt-ended, 2.3-kb interposon from plasmid pHPI45Ω [40] into a single SfiI site residing in the center of the sat gene, resulting in plasmid pAWP202. This plasmid was transferred to A. vinosum SM50 [26] by conjugation from E. coli SM10 [39] according to [41]. The genotype of the resulting double-cross over mutants was verified by Southern hybridization.

Cloning, Protein Expression and Purification

The sat gene from A. vinosum was amplified from chromosomal DNA by PCR using primers aw9 (5’-AGGAGGTTCCATATGATCGACCCAG9-3’ and aw10 (5’-GGTCAACTCAGGATCCCTC TAGACAA999-3’) and subsequently cloned between the NdeI and BamHI sites of expression vector PET-11a (Novagen, Darmstadt). The protein was overexpressed in E. coli BL21(DE3) pLysS cells (Novagen, Darmstadt) using LB medium at 303 K and 225 rpm agitation. Expression of ATPS was induced at 303 K and 225 rpm agitation. Expression of ATPS was induced at 303 K and 225 rpm agitation. The protein was overexpressed in E. coli buffer (Fig. S3).

For crystalization ATPS was purified as follows: thawed cells were suspended in 20 mM Tris-HCl pH 8.0 and disrupted by sonication. Insoluble components were removed by centrifugation at 171500 g and 4°C. The supernatant was adjusted to an ammonium sulfate concentration of 0.75 M and loaded onto a Phenyl-Sepharose column (1.6 cm x 10 cm; GE Healthcare, Munich). ATPS eluted at three peaks; fractions from the second and third peak were concentrated by ultrafiltration (Amicon Ultra-4, 10 kDa cutoff; Millipore, Eschborn) and dialyzed against 50 mM, Tris-HCl pH 8.0 (Membrane “Visking”, Roth, type 20/32). The protein solution was loaded onto a Resource Q column (0.5 cm x 10 cm; GE Healthcare, Munich) and eluted by a linear gradient (0–1 M NaCl) at about 0.16 M NaCl. After concentration by ultrafiltration purification was completed by using a Superdex 200 HR 10/30 gel filtration column (GE Healthcare, Munich) equilibrated with 50 mM Tris-HCl pH 8.0 and 150 mM NaCl. Protein was concentrated to 10 mg/ml and stored at 100 K in 50 mM Tris-HCl pH 8.0.

Enzyme Assays

ATP sulfurylase was routinely measured in the thermodynamically favoured direction of ATP generation from APS and PPi using the coupled spectrophotometric assay described by [4] with slight modifications. In a total volume of 1.0 ml the reaction mixtures contained 100 mM Tris-HCl, pH 8.0, 20 mM β-D-glucose, 4 mM MgCl2, 0.5 mM Na-NADP, 1 mM Na-PPi, 10 units glucose-6-phosphate dehydrogenase, 7.5 units hexokinase, 0.2 mM APS and enzyme extract. Reduction of NADP was followed at 340 nm (ε = 6.22 mM⁻¹ cm⁻¹) and 50°C.

The continuous spectrophotometric molybdolysis (AMP release) assay [42] was used for characterizing the activity of the enzyme on the “short-circuiting” inorganic substrate molybdate instead of sulfite. The molybdate- and ATP-sulfurylase-dependent formation of AMP was monitored in a reaction mixture (1 ml) containing the following final concentrations of reagents: 50 mM Tris-HCl, pH 8.0, 50 mM MgCl2, 4 mM phosphoenolpyruvate, 0.3 mM Na-NADH, 1 mM KCl, 50 mM NaMoO4, 10 mM ATP, 20 units pyruvate kinase, 22 units lactate dehydrogenase, 14 units adenylate kinase and enzyme extract. Oxidation of NADH was followed at 340 nm and 30°C.

Primary plots of initial rate against substrate concentration fit to the Michealis-Menten equation were created and analyzed by non-linear regression using Graph Pad Prism (version 6.0; Graph Pad).

Presence of APS reductase activity in A. vinosum wild type and mutant strains was assessed via thin-layer chromatography of APS reductase reactions as described in [26].

Crystallization and X-ray Data Collection

Initial crystals were obtained at a temperature of 291 K within a vapor diffusion experiment using the crystallization kits Classic from Jena Bioscience and MDL I-H of Molecular Dimensions. Optimization led to a drop content of 2 µl enzyme solution (10 mg ml⁻¹) in a buffer consisting of 50 mM Tris-HCl pH 8.0 and 2 µl reservoir solution (1.5 M potassium sodium tartrate and 100 mM MES pH 6.5). Crystals grew in the space group P2₁ with unit cell parameters of a = 73.3 Å, b = 97.0 Å, c = 73.5 Å and β = 117.6° and two subunits in the asymmetric unit (Vₐ = 24 Å³Da⁻¹, solvent content 49.2%). For cryoprotection, crystals were placed for several minutes into a buffer containing 1.5 M potassium sodium tartrate, MES pH 6.5 and 15% glycerol. Data were collected at 100 K up to a resolution of 1.6 Å at the ESRF in Grenoble, France and processed with the HKL program suite.
The substrate binding pocket is designed to perfectly position APS and PP, in adequate proximity. The calculation of the electrostatic potential of the surface using the program APBS (Baker, N. A., Sept., D., et al. 2001), “Electrostatics of nanosystems: application to microtubules and the ribosome.” Proc Natl Acad Sci U S A 98:10357-41.) revealed an overall negative potential on the surface (red) and around the substrate binding pocket a positive charge (blue).

**Figure S3 SDS–PAGE (10%) of recombinant *A. vinosum* ATP sulfurylase.** The gel was stained with Coomassie brilliant blue. Protein purity was assessed after Nickel-chelate affinity chromatography (lane 1) and subsequent gel filtration chromatography (lane 2). Protein loaded: lane 1, 2 μg; lane 2, 1 μg.

(PDF)

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**Author Contributions**

Conceived and designed the experiments: KP AW UE CD. Performed the experiments: KP EW AW UD. Analyzed the data: KP EW AW UD. Contributed reagents/materials/analysis tools: UE CD. Wrote the paper: KP UE CD.

**References**


