Three-dimensional Structure of A1A0 ATP Synthase from the Hyperthermophilic Archaeon *Pyrococcus furiosus* by Electron Microscopy

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The archaean ATP synthase is a multisubunit complex that consists of a catalytic A1 part and a transmembrane, ion translocation domain A0. The A1A0 complex from the hyperthermophile *Pyrococcus furiosus* was isolated. Mass analysis of the complex by laser-induced liquid bead ion desorption (LILBID) indicated a size of 730 ± 10 kDa. A three-dimensional map was generated by electron microscopy from negatively stained images. The map at a resolution of 2.3 nm shows the A1 and A0 generated by electron microscopy from negatively stained samples. The A1A0 complex from the hyperthermo-

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[Full text continues here, detailing the structure, subunits, and implications of the research.]
cies have retained an ion-translocating glutamate in every hairpin, *M. jannaschii* has lost it in the first of three and *P. furiosus* has lost it in the first of its two hairpins. Therefore, *P. furiosus* has a typical V-type e-subunit that was always seen as the reason for the inability of the V-type ATPases to synthesize ATP (36–38). However, the A1A0 ATP synthase from *P. furiosus* functions as ATP synthase in vivo (39). The structural basis for its ability to synthesize ATP despite the V-type e-subunit is unknown but might reside in the A0 domain and may be based on a much larger c-ring (40).

We have recently enriched the A-type ATP synthase from the hyperthermophilic archaeon *P. furiosus* and shown that it uses Na+ as a coupling ion (41). Here, we have improved the purification protocol and purified the entire complex. We present the three-dimensional structure of the complex by electron microscopy as well as a mass analysis by laser-induced liquid bead ion desorption mass spectrometry (LILBID-MS). Based on the results of these two methods, we propose a model for the subunit composition and a solution for the enigma of ATP synthesis.

**EXPERIMENTAL PROCEDURES**

*Organism—* *P. furiosus* DSM 3638 was obtained from the Deutsche Sammlung für Mikroorganismen und Zellkulturen, Braunschweig, Germany. For purification of the ATPase, *P. furiosus* was grown in a 300-liter fermentor at 98 °C as described before (41). The pellets were stored at −80 °C.

*Purification of the A1A0 ATP Synthase of *P. furiosus*—*P. furiosus* was grown in a 300-liter fermentor at 98 °C as described before (41). The pellets were stored at −80 °C. The supernatant after this wash step contained negligible ATP hydrolysis activity. The supernatant was collected by centrifugation (120,000 **g**; 1 h). After MgCl2 was added to the supernatant to a final concentration of 50 mM, the ATP synthase was further purified by polyethylene glycol 6000 (4.1%, wt/wt) precipitation, where contaminating proteins were precipitated. The precipitate was removed by centrifugation (120,000 × **g**; 1 h), and the pellet contained negligible ATP hydrolysis activity. The supernatant was applied to a 20–66% sucrose gradient and centrifuged for 19 h in a vertical rotor at 153,000 × g 4 °C (Beckman L100K, VTi 50 rotor). Samples containing the highest ATP hydrolysis activity (determined as described in Pisa *et al.* Ref. 41) were pooled and applied to anion exchange chromatography (DEAE-Sephrose), equilibrated with 100 mM Tris, pH 7.5, 5 mM MgCl2, 10% glycerol (v/v), 0.1% Triton X-100, and 0.1 mM PMSF.

Elution was performed with a salt gradient from 0 to 1 M KCl in 100 mM Tris, pH 7.5, 5 mM MgCl2, 10% glycerol (v/v), 0.1% Triton X-100, and 0.1 mM PMSF. The ATPase-active fractions were pooled and concentrated on Vivaspin 20-m1 (Biofiltronic GmbH, Nörten-Hardenberg, Germany) 100-kDa concentrators. The concentrated sample was loaded on a Superose 6 column (10/30 Amersham Biosciences) and eluted with 50 mM Tris-HCl, pH 7.5, 5 mM MgCl2, 10% glycerol (v/v), 50 mM KCl, 0.1% Triton X-100, 0.1 mM PMSF. The peak fractions were analyzed by SDS-PAGE. All steps were performed at 4 °C.

**Tryptic Digest and MALDI-TOF Analysis—**The products of the A1A0 ATPase were excised from a gel and subjected to in-gel digestion protocols and MALDI-TOF analysis as described (41). The A1A0 ATPase protein was buffer-exchanged and concentrated to a final concentration of 3 μM in 10 mM ammonium acetate with 0.05% dodecylmaltoside (DDM) with Microcon filters (Microcon YM-100, Millipore GmbH Schwalbach/Ts. Germany). LILBID-MS of the sample was performed as described previously (44, 45). Briefly, microdroplets (50-μm diameter; volume, 65 pl) of the sample solution was introduced into vacuum by an on-demand, piezo-driven droplet generator at a frequency of 10 Hz. There the droplets are irradiated one by one by IR laser pulses, tuned to the absorption maximum of water at around 3 μm. The laser energy is transferred into the liquid via excitation of the stretching vibrations of water, inducing an instantaneous transition into the supercritical state of the liquid. There the droplets explode and the charged biomolecules from solution are set free. Because of the loss of solvent screening in this process, most of the ions are neutralized and only 1 in 10,000 escape into vacuum, called “lucky survivors.” These are accelerated and mass-analyzed in a home-built time-of-flight (TOF) reflectron mass spectrometer, equipped with a custom-built high mass ion-detector. The spherical isotropic explosive expansion of the ion cloud (with a velocity of around 800 m/s) limits the present resolving power in the kilodalton mass range to an approximate M/ΔM value of 100 and even lower in the higher mass range.

**Electron Microscopy (EM) and Image Analysis—**The A1A0 sample was applied to glow-discharged carbon films and negatively stained with 1% (w/v) uranyl acetate. Electron micrographs were collected using a Philips CM120 (FEI, Eindhoven, the Netherlands) at an accelerating voltage of 120 kV under low dose conditions. Images were taken at a magnification of ×44,000 on Kodak SO-163 electron image film. The negatives were developed for 12 min in full-strength D-19 developer. Negatives checked by optical diffraction for correct defocus and lack of drift and astigmatism were digitized on a PhotoScan scanner (Z/I Imaging, Aalen, Germany) at a pixel size of 7 μm. Subsequently, adjacent pixels were averaged to yield a pixel size

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The abbreviations used are: LILBID, laser-induced liquid bead ion desorption; EM, electron microscopy; MS, mass spectrometry; MALDI, matrix-assisted laser desorption/ionization; TOF, time-of-flight; PEG, polyethylene glycol; PDB, Protein Data Bank; PMSF, phenylmethylsulfonyl fluoride.
on the specimen of 4.77 Å. 12,912 particle images were selected using the boxer program from EMAN (46). Images were processed using Imagic V (47). Briefly, images were band pass-filtered, aligned translationally and rotationally, and subjected to multivariate statistical analysis and classification. Images assigned to the same class were averaged. As all classes represented views along the long axis of the molecule, for three-dimensional reconstruction an initial model was created by assigning angles to some class averages and creating a volume in Imagic. The model was iteratively refined by the standard refine command of EMAN (46). To estimate the resolution of the final map the Fourier shell correlation between two maps calculated from half of the particle images each was determined by the EMAN routine eotest. Three-dimensional visualization and fitting of x-ray structures into the map was done by UCSF Chimera (48).

RESULTS

Purification and Characterization of the A₁A₀ ATP Synthase of P. furiosus—The A₁A₀ ATP synthase preparation studied previously lacked the subunits F and H and contained an AAA ATPase (thermosome) as a major contaminant (41). The thermosome, which has a calculated molecular mass around 900 kDa, was clearly visible in electron micrographs and easily distinguishable from the A₁A₀ ATP synthase. The new protocol yielded an intact enzyme and removed the thermosome (see “Experimental Procedures”). In short, the membranes were centrifuged at low speed, the enzyme was solubilized, precipitated with PEG, and further purified by sucrose density centrifugation and ion exchange chromatography followed by gel filtration. The purification is summarized in Table 1. The specific activity of various enzyme preparations ranged from 1.9 to 6.5 units/mg protein, and the overall yield was 1.8%.

The ATP synthase preparation in SDS-PAGE contained 14 polypeptides of apparent molecular masses 130, 97, 65, 60, 54, 52, 40, 30, 25, 23, 16, 12, 11, and 9 kDa (Fig. 1). Proteins were identified by MALDI-TOF analysis or MS/MS. The 130-kDa protein was identified as methylmalonyl-CoA decarboxylase subunit α. The 97-kDa protein was identified by MALDI-TOF analysis and MS/MS in duplicate to be subunit C. No other polypeptide was found, indicating that this band represents a multimer of subunit C (42 kDa). Subunit A (65 kDa) matches exactly the mass deduced from the genome sequence. The band at 60 kDa was identified by MS/MS to contain an oligomeric form of subunit c and subunit a. The subunits could not be resolved by SDS-PAGE, neither in 8% nor in 10% gels. Furthermore, the band cross-reacts with the antiserum against subunit a or c of M. jannaschii. The hydrophobic character of this membrane protein complex makes it difficult to estimate its molecular weight on an SDS-PAGE gel. The 54-kDa protein was also identified as methylmalonyl-CoA decarboxylase subunit α (which has a predicted mass of 57.1 kDa). The 52, 40, 25, and 23 kDa proteins were identified as subunits B, C, D, and E. The 30-kDa protein could not be identified, and the 16-kDa protein was identified as the hypothetical protein PF0672 from the P. furiosus genome (49). The 12-, 11-, and 9-kDa proteins were identified as subunits c (monomeric form), H, and F. These data showed the absence of thermosomes in the preparation and demonstrated that all subunits deduced from the genome sequence are indeed present in the preparation.

**TABLE 1**

| Purification step         | Protein | ATPase activity | Specific activity | Enrichment | Yield |
|---------------------------|---------|-----------------|-------------------|------------|-------|
| Membranes                 | 335     | 106             | 300               | 1          | 100   |
| TX-100 supernatant        | 66      | 60              | 900               | 3          | 56.6  |
| PEG 6000 supernatant      | 64      | 67              | 1100              | 4          | 63.4  |
| Sucrose gradient          | 3.1     | 76.8            | 24600             | 78         | 72.4  |
| DEAE column conc.         | 1       | 5.1             | 6000              | 19         | 4.8   |
| Gel filtration            | 0.3     | 2               | 6500              | 20.5       | 1.8   |

FIGURE 1. Subunit composition of the A₁A₀ ATP synthase of P. furiosus. The preparation was subjected to SDS-PAGE on 12.5% gels and silver-stained. A molecular mass marker is shown on the left and subunits on the right. Letters indicate the A₁A₀ subunits, and numbers the impurities: 1, methylmalonyl-CoA decarboxylase subunit α; 2, not identified; 3, hypothetical protein PF0672 (see text).
same mass range (51). The mass spectrum in Fig. 2a shows an m/z distribution of the anionic complex with charges varying from 3 to 9. The peaks are broadened due to an increase in initial energy of the ions with higher m/z. Different from other membrane protein complexes but similar to the F1F0 ATP synthase (50), the detergent is more or less completely lost by the transcriptionally or post-translationally modified. The mass calculated for subunit c is 15.8 kDa, and indeed, a 15.8-kDa protein was found in the complex. There is no mass found for a protein of around 8 kDa, which would have only two transmembrane helices. In addition, oligomers of c appear in the mass spectrum such as c2 and c3, indicating an incomplete fragmentation of the c-ring at this elevated level of laser intensity. However, an ion signal at (c/2)c is not observed, again indicating a c-subunit size of 15.8 kDa.

The spectra contained some additional peaks that could not be assigned to any of the subunits. Most of these are at low m/z, and some may arise from the 16- and 30-kDa impurities (Fig. 1). Interestingly, a series of unexpected ion signals in the higher m/z region corresponds to a mass to charge ratio of 233 kDa with charge states of −3 to −6 (purple bars in Fig. 2b). Such large complexes form broad peaks of which the onsets are measured to determine the theoretical masses. The peak around m/z = 46,000 is overlaid by a peak from c3 and the peak around m/z = 78,000 with the a peak. The a peak is reproducibly very broad, therefore the assumption of an overlay with another peak seems reasonable. Among the various combinations of subunits, only an ac10 (75 + 10 × 15.8 kDa) subcomplex would correspond to a mass of 233 kDa. This gives additional evidence, apart from the SDS-PAGE, for a stable ac10 subcomplex, and indicates that the c-ring consists of 10 subunits.

Electron Microscopy and Image Processing—The P. furiosus A1A0 preparation was monodisperse and showed typical dumbbell-shaped particles with no hint of barrel-shaped thermosomes (Fig. 3a). An initial image analysis by IMAGIC V of a 7500-particle data set resulted in many different class averages with features including a head with one or more spikes, a distinct central stalk, a prominent “collar,” indications of one or two peripheral stalks, and a rather large A0 domain with clear substructure (Fig. 3b). The large number of different views precluded a reconstruction by random conical tilt, for which particles in identical orientations are needed. It suggested that the particles might represent a single-axis tilt series around the long axis of the molecule, which would contain all information necessary for a three-dimensional reconstruction. The complex was reconstructed by EMAN, which can iteratively refine a data

### Table 2: Theoretical and measured masses of the A1A0 ATP synthase subunits

| Subunit | Theoretical mass | LILBID mass |
|---------|------------------|-------------|
| A       | 65,900           | 66.0        |
| B       | 51,785           | 51.9        |
| C       | 42,515           | 42.5        |
| D       | 24,895           | 24.6        |
| E       | 23,502           | 23.0        |
| F       | 11,740           | 11.9        |
| H       | 12,394           | 12.2        |
| a       | 75,466           | 75.5        |
| c       | 15,806           | 15.8        |
set to an initially very crude three-dimensional model (46). An initial model consisting of a large and a small domain was created by assigning azimuthal angles to some class averages. The data set was iteratively refined against this model, resulting in a clearly asymmetric structure with hints of two peripheral stalks. More data were added from particles in slightly different orientations that had initially been excluded. Refinement of this 13,000-particle data set resulted in a model with two continuous peripheral stalks. All class averages from the initial analysis by IMAGIC could be matched to one reprojection of the three-dimensional volume (Fig. 4). The relative angles of the classes that had been used in the starting model had no relation to their original assignment, showing clearly that the reconstruction process by EMAN was robust enough to overcome any bias from the starting model. The final volume was estimated to have a resolution of 23 Å by Fourier Shell Correlation (0.5 cut-off criterion). The map is shown in slices along the long axis (Fig. 5a), and in the perpendicular direction (Fig. 5b) and in a surface representation (Fig. 6). It shows a hollow head displaying six distinct densities, an asymmetric A₀ domain consisting of a hollow ring with a bulge on one side, a central stalk made up of two distinct domains, and a stator with two peripheral stalks, which are both connected to a collar around the central stalk while one of them is connected to the bulge on A₀.

The Head—The A₁ head shows pseudo-3-fold symmetry and has a strong resemblance to the isolated A₁ from Methanosarcina mazei Gö1 (23). The head shows clear asymmetry because of the presence of the central stalk. It displays a central cavity surrounded by six densities representing the three A and three B subunits, as can be seen in the sections 22–31 of Fig. 5a. Three of the densities have a bulge on the top, visible as “spikes” in the class averages (Fig. 4b), and can thus be identified as the catalytic subunits A (equivalent to /H9252 in F₁), which are distinguished by a 90-amino acid insertion forming a β-sandwich (11). X-ray structures of the individual A subunit from Pyrococcus horiko-
shii (PDB 1VDZ) (11) and the B subunit from M. mazei Go"1 (PDB 2C61) (12) have been determined. Because of the low resolution of the model, it was not possible to fit the individual A and B subunits unambiguously. Instead, the bovine F1 head (PDB 1H8E) was fitted to the EM density, and three A and three B subunits were fitted to the bovine F1 subunits at the positions dictated by the presence of the bulges. The absolute handedness of the structure cannot be determined by our reconstruction procedure, so we tried the fit with both the EM structure and its mirror image. Because of the well-defined structure and asymmetry of the head, especially at the lower part where the A subunits protrude much further than the B subunits (see Fig. 7h), the correct handedness could be determined unambiguously, and the model fitted quite well (Fig. 7a, e, h). It is clear that the peripheral stalks connect to the B subunits exclusively (Fig. 7a), as was observed for the T. thermophilus structure (12).

The Central Stalk—The central stalk descends from the head at an acentric position and attaches to the A0 domain inside the ring (Fig. 5a, slices 39–40). Two distinct domains can be distinguished, a smaller, knob-like one above the collar and a larger, funnel-shaped one starting inside the collar and continuing inside the ring. The structure of the C subunit of T. thermophilus ATPase was solved by x-ray crystallography (14), and it was localized in the lower position of the central stalk (19, 52). This position can be clearly confirmed in the P. furiosus map (Fig. 7, c, e, f). The C subunit consists of three bundles of four \( \alpha \) -helices each. Three domains can also be recognized in the EM density (Fig. 5a, slices 45–48). The C subunit fits very well in the stalk density, although it is not possible to distinguish the three domains (Fig. 7, c, e, f). The top surface is at the level of the collar and at the bottom the C subunit connects to A0 inside the c-ring, which is hollow at this level (Fig. 7, d–f).

The other central stalk components are subunits D and F, with molecular masses of 25 kDa and 12 kDa, respectively. Subunit D can be cross-linked to the inside of subunit B (53) and has a high \( \alpha \)-helical content (54). Unlike the C and F subunits, it is protected from trypsin degradation in an A3B3CDF A1 subcomplex (8); it is probably the equivalent of the \( \gamma \) subunit in F-ATP synthase and would thus be located inside the head. Recently an x-ray structure of the P. furiosus F subunit (PDB 2QAI) as well as an NMR structure of subunit F of M. mazei Go"1 were presented (16). Both structures show a globular domain consisting of helices and strands, with the C terminus forming a flexible extension in the NMR structure that was for the most part not visible in the x-ray map. Based on its shape, the globular domain fits well in the central stalk domain above the C subunit (Fig. 7, b, e, f). The C terminus can be cross-linked to B (16), so the F subunit is probably oriented with the C-terminal helix pointing toward the head and the N-terminal globular domain connecting to the C subunit. At the low resolution of the EM map, the orientation of the F subunit in the bulge cannot be determined, and the subunit has been fitted manually (Fig. 7b). The bulge may also be occupied by a part of the D subunit, for which there is no structure.
The A\textsubscript{0} Domain—The \textit{P. furiosus} A\textsubscript{0} domain has an oval shape and displays clear substructure. The cytoplasmic half is hollow, and there is a shallow indentation opposite the hole on the extracellular side (Fig. 5, \textit{a}, slices 52–55 and \textit{b}, slices 37–40). These features are likely to represent the center of the c-ring. As the LILBID analysis suggested that the ring consists of ten 15.8-kDa subunits, the map density was compared with c-rings of known structure. Rings of various sizes ranging from 10 to 15 \(\alpha\)-helical hairpin subunits have been found for the F\textsubscript{1}F\textsubscript{0} type (29, 31–35, 55). For A- and V-type, the only known stoichiometries are for the \textit{Enterococcus hirae} ring consisting of 10 duplicated subunits, comprising 20 hairpins (52) and the \textit{T. thermophilus} ring, which has 12 hairpins (56). Two rings have been solved by x-ray crystallography: the bacterial F\textsubscript{1}F\textsubscript{0} c-ring from \textit{Ilyobacter tartaricus} (57) with 11 hairpins and the 20-hairpin, V/A-type c-ring from \textit{E. hirae} (52). Comparison with the EM structures of the ATP synthases from \textit{T. thermophilus} (19) and from \textit{Bacillus} sp. TA2.A1 (58) shows that the \textit{P. furiosus} membrane domain is larger than that of these species, which were recently shown to contain a c\textsubscript{12}-ring (56) and a c\textsubscript{13}-ring (33), respectively. The \textit{I. tartaricus} c\textsubscript{12}-ring is clearly much too small when compared with the hole in the ring (not shown), whereas the \textit{E. hirae} c-ring fits quite well (Fig. 7, \textit{b} and \textit{h}). The hole seen in the cytoplasmic side of the \textit{P. furiosus} ring is also present in \textit{E. hirae}, where the other side is filled with lipid. A similar situation is likely to exist in \textit{P. furiosus}. The c-subunit sequences are quite similar and like in \textit{E. hirae}, the first and third helices, which form the inner ring, are completely hydrophobic at the N termini (facing away from A\textsubscript{1}), whereas the C-terminal parts (facing A\textsubscript{1}) contain several charged residues (41). This is consistent with the negative stain images, as the lipid-filled hydrophobic end would not be accessible to stain unlike the water-filled top part of the ring. We conclude that the \textit{P. furiosus} ATP synthase has a 20-hairpin c-ring similar to the one in \textit{E. hirae}.

Outside the ring, another mass is visible, which is connected to the collar (Fig. 7, \textit{d–f}). This indicates the presence of the hydrophobic part of the \(\alpha\)-subunit at this location. This subunit is predicted to have 7–8 transmembrane helices (1), which is compatible with the size of the A\textsubscript{0} density. The \(\alpha\)-subunit has a large hydrophilic domain that forms part of the stator, as discussed below.

The Stator—Two peripheral stalks run from prominent knobs on A\textsubscript{1} to a level above A\textsubscript{0}, where they are connected to each other via a collar surrounding the central stalk. The knobs are \(\sim 120^\circ\) apart and connect to two B subunits, as discussed above. Unlike in the \textit{T. thermophilus} reconstruction (19), the two knobs are not connected to each other across the \(\alpha\)- and \(\beta\)-subunits. The collar runs along the upper part of the C subunit parallel to the c-ring, and the collar and C subunit share a flat surface 30 Å above the A\textsubscript{0} domain. Although in the surface representation the collar and central stalk seem to be connected, a separation between these two features is visible in the EM density (Fig. 5, \textit{a}, slices 45–49). This separation is expected considering the rotary mechanism of the ATP synthase. One of the peripheral stalks is connected to A\textsubscript{0}; the other stalk extends laterally outside the A\textsubscript{0} region and has no connection below the level of the collar.

The three subunits forming the stator are E, H, and A. The 12-kDa H subunit is the smallest protein in A\textsubscript{1}A\textsubscript{0}. Recently, solution x-ray scattering experiments have shown that the isolated H subunit forms a parallel dimer and has an extended “boomerang” shape with two arms of 68 and 120 Å length, respectively (17). A secondary structure prediction by PSIPRED (59, 60) for the H subunit shows an N-terminal helix of \(\sim 60\) amino acids separated by a short coil region from a 40-amino acid C-terminal helix (supplemental Fig. S1a), which fits well to the SAXS data. The C terminus of H can be cross-linked to the N terminus of the A subunit (on top of the head) in \textit{M. mazei} G601 (61), suggesting that H is part of the peripheral stalk, oriented with the C terminus on the head and the N terminus near the collar.

The 23-kDa E subunit consists of two domains, a highly helical N-terminal domain (E\textsubscript{N}) and a C-terminal domain (E\textsubscript{C}) with short helices and \(\beta\)-strands (supplemental Fig. S1c). The x-ray structure of the E\textsubscript{C} domain from \textit{P. horikoshii} has recently been determined (13); it has an elongated shape \(\sim 75\) Å long with \(\alpha\)-helical and \(\beta\)-strand elements and forms a tight dimer in the crystal structure, with all the predicted secondary structure elements (supplemental Fig. S1c). The \(\alpha\)-helical N-terminal domain (E\textsubscript{N}) is homologous to H and to the F\textsubscript{1}F\textsubscript{0} b-subunit and interacts with the H subunit (13). The equivalent E subunit in yeast V-type ATPase can be cross-linked to the full-length of the outside of the B subunit (53, 62), placing E in the peripheral stalk. A recent study shows that the E and H subunit from \textit{Thermoplasma acidophilum}, when expressed together or reconstituted, form a stable heterodimer with high affinity (63), suggesting that the H or E homodimers formed in solution or in crystals of the isolated subunit are artifacts. This was confirmed in an EM study of the \textit{T. thermophilus} ATPase, where antibodies against an EG complex (equivalent to EH in archaea) attached to both sides of the head, while not or weakly binding to isolated E or G (64). All this evidence places an EH dimer in

![FIGURE 7. Fitting of x-ray structures in the EM map. a–f, cut-open views of the map. a–d, vertical views of the map, cut at the levels indicated in e. a, fitting of A\textsubscript{1}B\textsubscript{1}C\textsubscript{1}A\textsubscript{2}B\textsubscript{2}C\textsubscript{2}; A from PDB file 1VDZ (11) (blue) and B from 2CG1 (12) (green), b, fitting of the F subunit (1QAI) (purple), c, fitting of the C subunit (turquoise) (1RSZ (14)), d, fitting of the 20-hairpin \textit{E. hirae} A-type c-ring from 2BL2 (52) (gold), e and f, two perpendicular views along the membrane plane. The EC domain is fitted to the knobs from 2DM9 (13) (pink), g, all fitted subunits in the same orientation as f, \(h\), model for the stator with the \(\alpha\)-subunit in yellow and the EH dimers in pink. The molecular graphics images were produced using the UCSF Chimera package from the Resource for Biocomputing, Visualization, and Informatics at the University of California, San Francisco (supported by NIH P41 RR-01081).](image)
each peripheral stalk. The globular $E_c$ domain (13) has the right dimensions for the knobs on the head (Fig. 7, e and f), and the C terminus of the H subunit is also likely to reside there, as shown above. The stalks would thus be formed by heterodimers of the $N$-terminal domains of E and H, both of which have high $\alpha$-helical propensity (supplemental Fig. S1, b and c) and are likely to form coiled-coils.

The 75-kDa $a$-subunit is the largest protein in the $A_1A_0$ complex. It has a hydrophobic C-terminal domain predicted to have $7–8$ $\alpha$-helices, forming the equivalent of $a$ in $F_1F_0$ (1), whereas the $N$-terminal domain is hydrophilic. We conducted a secondary structure prediction with PSIPRED (59, 60) (supplemental Fig. S1b). The C-terminal 300 amino acids form the hydrophobic domain. The 360 amino acid N-terminal domain contains two long $\alpha$-helices for residues 100–140 and 235–290 as well as several shorter helices, stretches of $\beta$ strands and loops. The only possible location for $a_N$ is the collar connecting the two peripheral stalks, which is the only part of the three-dimensional volume not accounted for by other subunits and is also connected to the $A_0$ domain where the $a_C$ domain has been localized. Thus, the two domains of the $a$-subunit are at right angles to each other (Fig. 7h). A structure parallel to the membrane domain has been seen in a three-dimensional EM structure of bovine brain $V_0$ (65), supporting this interpretation.

The two peripheral stalks in this model have an identical composition, and being 120° apart on the 3-fold symmetric head, their interactions with $A_1$ via a $B$ subunit will be identical as well. However, the symmetry breaks at their connection to the collar, which is made up of a single subunit. Close inspection of the $a$-subunit sequence (supplemental Fig. 1a) shows some repeating structural elements. There are two predicted long $\alpha$-helices, one of $\sim$40 amino acids (residues 100–140) and one of $\sim$55 (235–290). At four places, there is a short predicted $\alpha$-helix flanked by two $\beta$-sheets, the first at the N terminus, the second and third immediately adjacent to each other after the first $\alpha$-helical stretch, and the fourth after the longest $\alpha$-helix, just before the membrane domain. The first and third $\beta\alpha\beta$ domain are $\sim$28-amino acids long, and the second and fourth $\sim$33. Interestingly, there is 33% sequence identity and more than 50% similarity between the second and fourth $\beta\alpha\beta$ domain, and the $\alpha$-helices in both domains are amphipathic (Fig. 8), although there is no significant homology between the first and third domain (Fig. 8). We propose that the N-terminal domain of the $a$-subunit is a loop with the first and the fourth $\beta\alpha\beta$ domain near the connection to $A_0$ and the second and third on the other side, leaving the main $\alpha$-helical domains between the $\beta\alpha\beta$ domains forming the collar proper. The interaction with the EH peripheral stalks is mediated by the two similar sets of $\beta\alpha\beta$ domains, which may have arisen by a gene duplication. A recent report (66) shows a 1:1:1 complex between the E, H, and $a_N$ equivalents of $E. hirae$. Small angle x-ray scattering experiments showed this complex to be L-shaped with dimensions which fit our model. Moreover, dimerization of E and H as well as interaction of H with $a_N$ is shown, but no interaction of E with $a_N$ in the absence of H. This implies that the interaction of the peripheral stalk with a is mediated through the N terminus of H. The only discrepancy between this study and our model is the presence of only one

![Structure of Archaeal ATP Synthase](image-url)

**FIGURE 8. Secondary structure elements of the $P. furiosus$ $a$-subunit.** Four regions of the sequence are shown with predicted $\beta$-strands in pink and $\alpha$-helices in yellow. Identical amino acids between the two sets of regions are indicated with capital X and homologous amino acids with lowercase x. Helical wheels are shown for the helices with hydrophobic side chains shown in black, polar in green, positively charged in blue, and negatively charged in red. TEH dimer in the ternary complex. This may be caused by the absence of the $a_C$ domain and resulting conformational differences.

**DISCUSSION**

$A_1A_0$ ATP synthases and $V_1V_0$ ATPases form two distinct groups of enzymes that are related to each other and more distantly to the F-type ATPases. A core group of subunits is highly conserved, whereas others are specific for certain groups. The structures of several subunits have been solved by x-ray crystallography and NMR spectroscopy in the last few years. For the whole enzyme, projections (18, 67, 68) and three-dimensional structures (19–22) by EM are available for several species. Cross-linking experiments have clarified the positions of subunits relative to each other (53, 54, 61, 62). The stoichiometry of the subunits is not yet firmly established. Whereas a consensus is emerging for the structure of the head and the rotor, different models exist for the stator. The eukaryotic $V_1V_0$ ATPases not only have additional subunits and a regulation ($V_1V_0$ dissociation/association) that is not observed in $A_1A_0$, but there are indications that they have a different stator than the prokaryotic enzymes, including possibly a third peripheral stalk.

The $A_1A_0$ ATP synthase from the hyperthermophilic archaeon $P. furiosus$ is very suitable for structural investigation because of its high stability. Using the new technique of LILBID analysis, we were able to determine the total molecular weight
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and identify all subunits. The high stability made it possible to do a three-dimensional reconstruction, which is the first of an archaeal A3,B3,CDE,F2 subunits with 20 hairpins and 10 ion translocation sites. An ac10 subcomplex, and a large c10-ring fits the map well. We conclude that the P.furiosus ATP synthase has a ring of 10 subunits with 20 hairpins and 10 ion translocation sites. An ATP synthase with a stoichiometry of A3B3CDE,F2 subunits would have a molecular mass of 738 kDa, fitting well with the mass of 730 ±10 kDa as determined by LILBID.

The ability of pyrococci to synthesize ATP with a duplicated c-subunit that has lost one ion binding site (39) has been an enigma. F-type ATP synthases have so far been shown to have 10 (29, 32) to 15 (35) ion binding sites in the ring and prokaryotic A-type 10 (52) to 12 (56). The pyrococci would need a c-ring with a considerably larger physical size than the F-type ATP synthases to have a sufficient number of ion binding sites, and we have shown that this is the case. According to \( \Delta G_0 = -nF \Delta \mu_{\text{ion}} \), a phosphorylation potential (\( \Delta G_{\text{phos}} \)) of ~50 to 70 kJ/mol would be sustained by the use of n = 2.9 – 4 ions/ATP at a physiological electrochemical ion potential of ~180 mV (\( \Delta \mu_{\text{ion}} \)) (69). Ten subunits and thus ten ion binding sites in the ATP synthase from P.furiosus would give an ion/ATP stoichiometry of 3.3. This would be sufficient to synthesize ATP, providing an explanation for the enigma of ATP synthesis in P. furiosus.

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