Structural Investigations of the Membrane-Embedded Rotor Ring of the F-ATPase from *Clostridium paradoxum*

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The Na+-translocating F-ATPase of the thermoalkaliphilic bacterium *Clostridium paradoxum* harbors an oligomeric ring of c subunits that resists dissociation by sodium dodecyl sulfate. The c ring has been isolated and crystallized in two dimensions. From electron microscopy of these c-ring crystals, a projection map was calculated to 7 Å resolution. In the projection map, each c ring consists of two concentric, slightly staggered, packed rings, each composed of 11 densities representing the α-helices. On the basis of these results, it was determined that the F-ATPase from *C. paradoxum* contains an undecameric c ring.

MATERIALS AND METHODS

Bacterial strain and culture conditions. *Clostridium paradoxum* DSM 7308 was maintained as a spore stock preparation in sterile distilled water at 4°C. Spores were germinated by heat shocking at 75°C for 2 min, followed by rapid cooling on ice for 45 s. *C. paradoxum* was grown anaerobically at 56°C in YT medium, which contained, per liter, 5.3 g of Na2CO3, 0.36 g of Na2HPO4 ·2 H2O, 0.075 g of KCl, 5 g of yeast extract, 10 g of tryptone, 0.2 g of cysteine HCl, 0.2 g of Na2S · 9H2O, and 0.3% (wt/vol) glucose. The pH of the medium was adjusted to 10.5 at 25°C with 5 M NaOH. Bacterial growth was routinely monitored by following the optical density of the culture at 600 nm. For large-scale growth, 200 liters of *C. paradoxum* cells was grown anaerobically in a 300-liter fermentor. The fermentor was inoculated with 5 liters of a preculture, and after 6 h at 55°C, the bacteria were harvested at a final optical density at 600 nm of 0.4 by continuous centrifugation. The cells were frozen in liquid nitrogen and stored at −80°C.

Purification of the c ring from *C. paradoxum*. The F1F0-ATPase of *C. paradoxum* was purified as previously described (7). The ATPase was mixed with 1% (wt/vol) N-lauroylsarcosine and incubated for 10 min at 65°C. After the mixture was cooled to room temperature, ammonium sulfate was added to yield 67.5% (wt/vol) saturation. Spores were germinated by heat shocking at 75°C for 2 min, followed by rapid cooling on ice for 45 s. 

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against 250 ml of 10 mM Tris-HCl (pH 8.0), 200 mM NaCl, and 3 mM NaN3. The mixture was dialyzed in 25-l buttons (Hampton Research, Aliso Viejo, CA) at lipid-to-protein ratios of 0.5, 1.0, and 1.5 (wt/wt). The sn-glycero-3-phosphocholine (POPC; Avanti Polar Lipids 1-palmitoyl-2-oleyl-Zwittergent 3-12 (0.4% [wt/vol])-solubilized c ring (1 mg/ml) was mixed with Zwittergent 3-12. The final concentration of the purified c ring was 4.7 mg/ml. The sample was then washed three times with dialysis buffer containing 0.15% Zwittergent 3-12. To further concentrate the c ring and remove excess salt, the sample was loaded on an Amicon Ultra-4 tube (50,000-Da molecular mass cutoff). The c ring was then adsorbed to 1/50 volume of hydroxyapatite in an Eppendorf tube (Bio-Gel HT; Bio-Rad, Richmond, CA), and after two washes with 1 M potassium phosphate buffer (pH 8.0) containing 0.15% Zwittergent, the c ring was eluted by centrifugation (23,000 × g for 20 min). The supernatant, which contained the c ring, was filtered through a 0.22-μm polyvinylidene fluoride filter unit (Millipore) and dialyzed against 10 mM Tris-HCl buffer (pH 8.0) overnight at 25°C. The c ring was then adsorbed to 1/50 volume of hydroxyapatite in an Eppendorf tube (Bio-Gel HT; Bio-Rad, Richmond, CA), and after two washes with 10 ml of dialysis buffer containing 0.15% (wt/vol) Zwittergent 3-12, the c ring was eluted with 1 M potassium phosphate buffer (pH 8.0) containing 0.15% Zwittergent 3-12. To further concentrate the c ring and remove excess salt, the sample was loaded on an Amicon Ultra-4 tube (50,000-Da molecular mass cutoff). The sample was then washed three times with dialysis buffer containing 0.15% Zwittergent 3-12. The final concentration of the purified c ring was 4.7 mg/ml.

2D crystalization of the c ring. For crystallization in two dimensions (2D), Zwittergent 3-12 (0.4% [wt/vol])-solubilized c ring (1 mg/ml) was mixed with 1-palmitoyl-2-oleyl-sn-glycero-3-phosphocholine (POPC; Avanti Polar Lipids Inc., Alabaster, AL) at lipid-to-protein ratios of 0.5, 1.0, and 1.5 (wt/wt). The mixture was dialyzed in 25-μl buttons (Hampton Research, Aliso Viejo, CA) against 250 ml of 10 mM Tris-HCl (pH 8.0), 200 mM NaCl, and 3 mM NaN3 buffer for 24 h at 25°C and another 24 h at 37°C. The two-dimensional crystals were stored at 4°C until further analysis.

Electron microscopy and image processing. Two-dimensional crystal samples were prepared in 4.5% (wt/vol) trehalose on molybdenum grids (Pacific Grid-Tech, San Diego, CA) (19) by the back-injection method (8). Grids were examined in a FEI 3000 SFF helium-cooled electron microscope at 4 K at an accelerating voltage of 300 kV. Images were recorded by a spot-scanning procedure, with 24 spots by 30 spots per image on Kodak SO-163 film at a magnification of ×70,000 and with an electron dose of 20 to 30 electrons/Å². The films were developed for 12 min in full-strength Kodak D-19 developer. Images selected by optical diffraction were digitized on a Zeiss SCAI scanner using a pixel size of 7 μm, corresponding to 1 Å on the specimen. Images were processed using the MRC (4) and CCP4 (2) program suites. Data were merged to a resolution of 7 Å.

Other methods. SDS-polyacrylamide gel electrophoresis (SDS-PAGE) was performed as described previously (16), and gels were stained with silver (14). ATPase activity was determined by the coupled spectrophotometric assay as described previously (7). The protein concentration was determined according to the bicinechonic acid method (17), with bovine serum albumin as the standard. To dissociate the c ring, c-ring samples (2 to 5 μg of protein) were treated with 5 volumes of 12% (wt/vol) trichloroacetic acid solution and incubated on ice for 5 min. The pellet was collected by centrifugation (11,000 × g for 5 min at 4°C) and dissolved in 1% (wt/vol) SDS-containing loading buffer. Prior to being loaded on an SDS-gel, the sample was neutralized by the addition of 1-μl aliquots of 0.5 M NaOH until the color of the bromphenol blue-stained loading buffer switched from yellow to blue.

FIG. 1. Sequence alignment of the atpE gene products from selected species. The sequences form an α-helical hairpin. A cytoplasmic loop (shown in black) connects both helices. The H⁺- or Na⁺-binding glutamate on the C-terminal helix is shown in red. The residues involved in Na⁺ binding and the hydrogen bridge-providing tyrosine are shown in yellow. At the end of the alignment, the corresponding coupling ion (C) of the corresponding ATP synthase is indicated. The amino acids which are in the area of the 2D crystal contacts are shown in dark green (cytoplasmic side) and blue (periplasmic side). The numbering of the amino acids is according to that of the C. paradoxa sequence (7). The selected species are Clostridium paradoxa (CP), Ilyobacter tartaricus (IT), Bacillus strain PS3 (PS3), Spinacia oleracea (SO), and Escherichia coli (EC).

FIG. 2. (A) SDS-PAGE of the c ring isolated from the F-ATP synthase of Clostridium paradoxa cells. Lane 1, c ring (c11) from Ilyobacter tartaricus; lane 2, c ring isolated from C. paradoxa; lane 3, subunit c monomer (c1) from C. paradoxa, derived from the oligomer by acidification with trichloroacetic acid. The gel was stained with silver and a standard gel marker is indicated on the left side. (B) MS-MALDI analysis of isolated c-ring preparations. The identified mass (Da) is indicated. The samples were prepared as described in Materials and Methods.
RESULTS AND DISCUSSION

Purification of the native c ring from *C. paradoxum* F₁F₀-ATPase. Previous investigations have shown that c-ring assemblies of Na⁺-translocating F₁F₀-ATP synthases are of exceptional stability, even sustaining being boiled in SDS (1, 9, 15). This remarkable stability is shared by the c ring of the Na⁺-translocating F₁F₀-ATPase from the thermoalkaliphilic bacterium *C. paradoxum* (7). A Na⁺-binding motif is also visible in

FIG. 3. Electron microscopy of 2D crystals from *C. paradoxum* c rings. (A) Overview of 2D crystals, negatively stained with uranyl acetate. The white scale bar represents 100 nm. (B) Projection map of six averaged images at a resolution of 7 Å. A unit cell containing two c rings is indicated.
the amino acid sequence of the c subunit of this bacterium (Fig. 1). As a consequence, the oligomeric c ring of *C. paradoxum* could be isolated by a modification of the protocol developed for the purification of the c ring from the *Ilyobacter tartaricus* ATP synthase (11). The results of an analysis of the purified c-ring sample of *C. paradoxum* by SDS-PAGE and mass spectrometry–matrix-assisted laser desorption/ionization (MS-MALDI) are shown in Fig. 2. After SDS-PAGE, the c ring migrated as a distinct band with an apparent molecular mass of 45 kDa (Fig. 2A, lane 2). Another protein band migrating with an apparent molecular mass of <6.5 kDa indicated a partial dissociation of the c ring into its monomeric units. After acidification of the sample with trichloroacetic acid, the c ring was completely dissociated into monomeric c subunits (Fig. 2A, lane 3). The mobility of the c ring from *C. paradoxum* was slightly faster than that of *I. tartaricus,*
and a corresponding difference in the mobilities of the c monomers was also observed. The calculated molecular mass of subunit c from *C. paradoxum* was 8,257 Da (Fig. 2B), 538 Da smaller than that of the c ring from *I. tartaricus* (8,795 Da), which could account for the different mobilities of the c monomers and of the c rings if these contained the same number of subunits (see below).

**Structural investigations of the *C. paradoxum* c ring.** Two-dimensional crystals of the *C. paradoxum* c ring were obtained at a lipid-to-protein ratio of 0.5 to 1.0 with the lipid POPC. Figure 3A shows an overview of a two-dimensional crystalline *C. paradoxum* c-ring sample which consists of vesicles containing crystalline areas with dimensions up to 0.5 μm. The 2D crystals are of plane group p1 (Table 1). A unit cell has the dimensions 99.2 ± 0.9 Å for a, 98.5 ± 0.4 Å for b, and 131.1 ± 0.3° for β and contains two c rings, each with a stoichiometry of 11. Each ring is in contact with three neighboring c rings. Such a packing was also observed in the 2D crystals of the *I. tartaricus* (18, 20) and *Propionigenium modestum* (11) c11 rings. The projection map, calculated to 7 Å resolution (Fig. 3B), shows that each c ring contains two concentric, slightly staggered, packed rings, each composed of 11 densities. Whereas the inner ring of densities is tightly packed, the outer one is more loosely arranged, and, by analogy with the *I. tartaricus* c ring, the densities correspond to the N- and C-terminal α-helices of the c ring, respectively. The inner and outer diameters of the rings, measured at the density borders, are the same as in the *I. tartaricus* c rings, approximately 17 Å and 50 Å.

**Comparison of the crystal packing (*I. tartaricus* and *C. paradoxum*).** The packing of the 2D crystals of c rings from *C. paradoxum* and *I. tartaricus* ATPase appears to have the same unit cell containing two rings, with each ring having three oppositely oriented neighboring rings (Fig. 4). In both crystals, all three crystal contacts in the unit cell are identical. However, the two p1 unit cells differ significantly in their dimensions and angles and also the packing of adjacent rings is different. The crystal contacts in *I. tartaricus* always involve two helices from one of the rings and one helix from the other, in a staggered arrangement, whereas in *C. paradoxum*, two helices from each ring are involved. The shape of the c ring resembles an hourglass, and therefore, the amino acids at the ends of the helices are responsible for the contact sites. In order to determine which amino acids are involved, we have built the high-resolution X-ray structure of the *I. tartaricus* c ring (12) into the electron density obtained from the *I. tartaricus* c-ring 2D crystals (20). According to this model, the crystal contacts consist on one side of the membrane of the side chains between K50 and S55 (ring 1) and between P83 and G86 (ring 2). On the other side, the contact sites are between Y80 and L87 (ring 1) and between K50 and L59 (ring 2) (Fig. 1). Between the *I. tartaricus* and the *C. paradoxum* c subunits, four amino acids differ in the K50-L59 peptides and four amino acids differ in the Y80-L87 peptides. For clarity, we have marked in Fig. 1 the residues in the crystal contact regions with colors. In a *C. paradoxum* homology model that is based on the crystal structure of the *I. tartaricus* c ring, several side chains that differ from those in *I. tartaricus* would cause several steric clashes, involving K50 to Q, I54 to L, S55 to R, V58 to L, and G86 to N (*I. tartaricus* numbering). To avoid these clashes, the rings are therefore reoriented versus each other in the *C. paradoxum* 2D crystal.

**The Na⁺-binding site in the *C. paradoxum* c ring.** The Na⁺-binding site in the c ring is the most notable feature in the structure. The way a Na⁺ ion is coordinated between two subunits has profound consequences in the mechanism of ion translocation and, hence, torque generation. Therefore, we have looked at the binding site in the homology model (Fig. 5). Based on the *I. tartaricus* model, the coordination site is located in the middle of the membrane at the tightest part of the ring and the entire ring is able to bind 11 sodium ions. The Na⁺ ion is bound at the interface of three helices: an N-terminal helix and two C-terminal helices. A putative Na⁺ coordination sphere can be made up by the conserved side chains of Q28, E61, S62, and the carbonyl oxygen of V59. Moreover, one could assume that E61 acts not only as one of the four Na⁺-binding ligands but also as the recipient of hydrogen bonds, one hydrogen bond being provided by Y66. In addition, the homology model shows that all side chains in close proximity to the ion-binding site are conserved as well. Almost all the differences between residues from the *C. paradoxum* and *I. tartaricus* rings are located at the termini or face towards the center of the ring. Based on these observations, we propose that the sodium ion translocation mechanisms are likely to be similar between the two c rings (6).

For an anaerobic bacterium like *C. paradoxum*, the conse-
quence of having an 11-mer oligomeric c ring is that, for every three ATP molecules hydrolyzed by the ATPase during glucose fermentation, 11 Na\(^{+}\) ions are pumped across the membrane to generate the $\Delta \mu_{\text{Na}^{+}}$. In the Na\(^{+}\) F-type ATP synthases that operate primarily in the synthesis direction during succinate or tartrate fermentation (e.g., those in *P. modestum* or *I. tartaricus*, respectively), the c ring also exists as an 11-mer. We conclude, therefore, that an undecameric stoichiometry is suitable for both operational modes in the F-ATP synthases.

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**REFERENCES**

1. Aufurth, S., H. Schägger, and V. Müller. 2000. Identification of subunits a, b, and c\(_1\) from *Acetobacterium woodii* Na\(^{+}\)-F\(_{1}\)-ATPase. Subunits c\(_2\), c\(_3\), and c\(_4\) constitute a mixed c-oligomer. J. Biol. Chem. 275:33329–333301.