Coupled rotation within single F₀F₁ enzyme complexes during ATP synthesis or hydrolysis

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Abstract F₀F₁ ATP synthases are the smallest rotary motors in nature and work as ATP factories in bacteria, plants and animals. Here we report on the first observation of intersubunit rotation in fully coupled single F₀F₁ molecules during ATP synthesis or hydrolysis. We investigate the Na⁺-translocating ATP synthase of Propionigenium modestum specifically labeled by a single fluorophore at one c subunit using polarization-resolved confocal microscopy. Rotation during ATP synthesis was observed with the immobilized enzyme reconstituted into proteoliposomes after applying a diffusion potential, but not with a Na⁺ concentration gradient alone. During ATP hydrolysis, stepwise rotation of the labeled c subunit was found in the presence of 2 mM NaCl, but not without the addition of Na⁺ ions. Moreover, upon the incubation with the F₀-specific inhibitor dicyclohexylcarbodiimide the rotation was severely inhibited. © 2002 Federation of European Biochemical Societies. Published by Elsevier Science B.V. All rights reserved.

Key words: F₀F₁ ATP synthase; Single molecule; Intersubunit rotation; Propionigenium modestum

1. Introduction

F₀F₁ ATP synthases couple H⁺ or Na⁺ transport to the synthesis of ATP from ADP and phosphate using the energy of a transmembrane electrochemical gradient. The multisubunit enzyme complex comprises the membrane-extrinsic F₁ domain and the membrane-intrinsic F₀ domain (Fig. 1A). F₁ consists of five different subunits in an α₃β₅γδε stoichiometry. The F₀ moiety of bacteria is built from three different subunits in an α₂β₃c₈ stoichiometry. Remarkably, the number of c subunits per F₀ subcomplex varies between species, i.e. 10 in yeast mitochondria [1], 11 in the bacterium Hyobacter tartaricus [2], and 14 in spinach chloroplasts [3].

An exceptional reaction mechanism for the synthesis of ATP, first outlined on the basis of kinetic measurements [4], gained dramatic support by the X-ray structure of F₁ [5]. The new concept, designated the ‘binding change mechanism’, involves the intimate coupling of catalysis at F₁ and ion flux across F₀ by rotation of the rotor (γεc₈) against the stator (α₃β₅γδε) (Fig. 1A). In order to achieve ATP synthesis, the transmembrane ion flux generates torque within F₀ that is transmitted by rotation of the rotor against the stator into conformational changes of the catalytic β subunits. Conversely, ATP hydrolysis would generate torque in F₁ that turns the rotor in the opposite direction and causes ion pumping.

During ATP hydrolysis, rotation of the γ subunit within the F₁ subcomplex has indeed been confirmed by biochemical [6–9] and spectroscopic techniques [10,11]. Most convincingly, the rotation of a micrometer-sized fluorescent actin filament attached to subunit γ has been directly visualized by video microscopy of single F₁ molecules [12] and rotation of γ could also be detected by single-fluorophore imaging [13]. It has been demonstrated that the ε subunit rotates together with γ [14,15] and that γ rotation proceeds in three discrete 120° steps [16]. Recently this approach has also been extended to the c subunit [17,18]. High-speed imaging revealed that a single 120° step is composed of two substeps of 90° and 30°, corresponding to ATP binding and ADP release, respectively [19]. One ATP was hydrolyzed per 120° motion and the torque of about 40 pN nm remains constant during rotation over a broad range of speed, load and ATP concentration [19].

Here, we use polarization-resolved confocal microscopy to study the rotation within the entire, functionally coupled F₀F₁ enzyme. Our study object is the Na⁺-translocating ATP synthase from Propionigenium modestum and we investigate the Na⁺ dependence of the rotation during ATP hydrolysis. Interestingly, this rotation could be blocked by the F₀-specific inhibitor N,N′-dicyclohexylcarbodiimide (DCCD). More important, by labeling of one c subunit with a single fluorophore and incorporation of F₀F₁ into liposomes, rotation during ATP synthesis could be observed. Rotation is initiated by the establishment of an electro-chemical Na⁺ gradient across the liposome membrane, thus demonstrating the structural integrity of the F₀F₁ enzyme complex. These findings prove the tight chemo-mechanical coupling between the F₀ and the F₁ subcomplex.

2. Materials and methods

2.1. Synthesis of the F₀ subunits a, b and c from P. modestum in Escherichia coli and purification of the proteins

As described previously [20], E. coli PEF42 (DE3) containing plasmids pMaHisN or pPhbHisC were used for the synthesis of subunits

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Abbreviations: DCCD, N,N′-dicyclohexylcarbodiimide; NTA, nitri- triacetic acid; ACF, autocorrelation function
a or b of the P. modestum ATP synthase, respectively. The recombi-
nant a subunit contains an N-terminal fusion of 10 histidine residues and six histidines are genetically engineered to the C-terminal end of subunit b. After expression and solubilization from the membranes, both subunits were purified by Ni\(^{2+}\) affinity chromatography. The P. modestum subunit c was overexpressed and purified as described [21].

2.2. Mutagenesis and labeling of P. modestum subunit c

To obtain fluorophore-conjugated P. modestum subunit c, the Asp at position 2 was replaced by a Cys using the ‘QuikChange’ site-
directed mutagenesis kit (Stratagene, La Jolla, CA, USA). Positive mutants were confirmed by DNA sequencing of both strands. The cyanine dye Cy3 was used as a label and Cy3-maleimide was produced from Cy3-NHS (Amersham Biosciences, Freiburg, Germany) as de-
dcribed [22]. Following established procedures [13], the genetically introduced single Cys residue was labeled with Cy3-maleimide at a labeling ratio of 0.93 mol Cy3 per mol c subunit.

2.3. Reconstitution of functional F\(_{0}\)F\(_{1}\) complexes

A functional F\(_{0}\)F\(_{1}\) sector was reconstituted into liposomes from the purified a, b and subunits of the Na\(^{+}\)-translocating ATP synthase of P. modestum [20]. During the reconstitution, the Cy3-labeled mutant c subunits were mixed with unlabeled wildtype c subunits at 1/15 to 1/25 ratios to minimize the occurrence of more than one fluorophore per F\(_{0}\)F\(_{1}\). Under these conditions, the probability of one incorporated fluorophore is five times higher than the incorporation of two fluo-
rophores with circularly polarized light. The fluorophore-conjugated liposomes were subjected to double-labeled complexes by detecting one bleaching event of Cy3 instead of two. Only data derived from singly labeled enzymes were used for further analysis.

Since the dye-labeled c subunit was separately modified, unspecific labeling of other F\(_{0}\)F\(_{1}\) subunits can be excluded. After incubation of the F\(_{0}\) liposomes with purified F\(_{1}\) containing subunits a, b (with a-Ni\(^{2+}\)-tag), and c (with a-E. coli and untagged subunit d from P. modestum [20] and removal of residual F\(_{1}\) by washing and centrif-
ugation, the F\(_{0}\)F\(_{1}\) holoenzyme was formed [20]. F\(_{0}\)F\(_{1}\) complexes were solubilized by treatment with 0.5-1% Triton X-100, purified by Ni\(^{2+}\)- affinity chromatography and concentrated by precipitation with 12% polyethyleneglycol 6000. The pellet was dissolved in PGM buffer (5 mM potassium phosphate buffer, pH 7.5, 0.2 mM MgCl\(_{2}\), 10% glycerol and 0.1% 2-mercaptoethanol) containing 0.05% Triton X-100 or K\(^{+}\)-cholate typically yielding an F\(_{0}\)F\(_{1}\) concentration of about 2 \(\mu\)M. The modification of single c subunits was found to be without effect on the Na\(^{+}\)-dependent ATP hydrolysis activity in comparison to the non-modified enzyme.

2.4. Reconstitution of in vitro assembled F\(_{0}\)F\(_{1}\) into liposomes

Liposomes were generated by extruding a suspension in PGM buñer equilibrated with an oxygen scavenging system consisting of 30 mM glucose, 0.3 mg/ml catalase and 0.2 mg/ml glucose oxidase and an ATP regenerating system consisting of 2.5 mM creatine phosphate and 0.2 mM creatine kinase. The reconstitution was performed at room temperature for at least 2 h. After each coating step, the cover glasses were rinsed with ultrapure water. For immobilization, solubilized, in vitro assembled F\(_{0}\)F\(_{1}\) was di-

2.5. Ni-NTA coating of cover glasses

The glass surface of cover glasses (24 \(\times\) 24 mm, Menzel, Braunschweig, Germany) was coated with Ni-NTA according to established procedures [13]. After pre-modifications, by baking at 500°C for 2 h, the cover glasses were incubated in sealing solution (2% (v/v) 3-glycidoxypropyl-trimethoxysilane (Fluka, Buchs, Switzerland), 0.01% (v/v) acetic acid) at 90°C for 3 h, coating solution (2% (v/v) N,N'-bis(carboxymethyl)lysine (Fluka, Buchs, Switzerland), 2 mM KHCO\(_{3}\), pH 10.0) at 60°C for 16 h and Ni\(^{2+}\)-solution (10 mM NiSO\(_{4}\), 5 mM glucose, pH 8.0) at room temperature for at least 2 h. After each coating step, the cover glasses were rinsed with ultrapure water.

2.6. Immunobilization of solubilized F\(_{0}\)F\(_{1}\) and F\(_{0}\)F\(_{1}\)-containing liposomes

For immobilization, solubilized, in vitro assembled F\(_{0}\)F\(_{1}\) was di-

2.7. Polarization-resolved confocal microscopy

The measurements were performed using a home-built scanning confocal fluorescence microscope [23]. A pulsed frequency-doubled Nd:YLF laser (Antelion) was used for excitation (532 nm). The fluorescence emission was collected with a dichroic mirror (560 nm) and a 620 nm longpass filter. The emission light was detected with a cooled charge-coupled device (CCD) camera (Roper Scientific, West  

3. Results and discussion

To observe ATP-driven rotation of the c ring in a function-
ally coupled Na\(^{+}\)-translocating F\(_{0}\)F\(_{1}\) ATP synthase, the holo-
enzyme was assembled from the individual a, b and c subunits of P. modestum and a hybrid F\(_{1}\) complex containing subunit d from P. modestum and the remaining subunits including a Hiss\(_{10}\) tag-containing b subunit from E. coli [20]. Subunit c carrying a D2C substitution was labeled with the fluorophore
Cy3-maleimide and added to a 15–25-fold molar excess of wildtype subunit c during reconstitution to favor the incorporation of a single fluorescent c subunit into the ATP synthase complex. Confocal images of single immobilized F₀F₁ molecules were taken using circularly polarized laser excitation (Fig. 1B). Subsequently, individual molecules were selected and upon continuous excitation the fluorescence intensity was recorded until photo-bleaching occurred.

To study rotation during ATP hydrolysis we observed individual F₀F₁ proteins immobilized via their His tags on NTA-functionalized coverslips and covered with buffer (Fig. 2A). The constant polarization shown in Fig. 2B indicates a stable steady-state orientation and low rotational mobility of the fluorophore as well as the protein complex, both important preconditions for the ability to detect the rotation of the c subunit. The two traces were recorded in buffer containing only contaminating Na⁺ concentrations (50 μM) and 0.5 μM ATP. Upon addition of 2 mM NaCl, stepwise changes of the polarization occurred and three discrete polarization levels could be detected (Fig. 2C). The dwell times in the polarization levels varied from 0.1 to 0.5 s. We assign this behavior to a complete stepwise rotation of the c₁₁ rotor part of the enzyme during ATP hydrolysis. The corresponding ATP hydrolysis rates ranged from 2 to 10 s⁻¹, consistent with previous studies on the F₁ part of E. coli with similar low ATP concentrations [13–15,24]. The observation that Na⁺ ions are obligatory for rotation of the c subunits proves the tight chemomechanical coupling of the F₀ part and the F₁ part in our Na⁺-translocating ATPase. These data are in accordance with crosslinking experiments performed with the ATP synthase of E. coli: after bridging the rotor subunits γ, ε, and c, coupled H⁺ translocation and ATP synthesis activities were retained [25]. Direct visualization of c subunit rotation with the aid of attached actin filaments was also reported for the E. coli enzyme [17,18]. However, the rotation was not sensitive to DCCD and persisted in the cD61N mutant, which is known to lack any H⁺-coupled ATP synthesis or hydrolysis activity [26]. These enzyme specimens were concluded to be decoupled at its ion motive portion [26], which is consistent with the observation that at the detergent concentrations used in rotation experiments, coupled H⁺ translocation no longer exists [27].

In our experiments, uncoupling was avoided by removing the detergent via extensive washing of the sample with 3×500 μl PGM buffer after immobilization to the cover glass. Under these conditions, 72% of the single molecules rotate in the presence of 0.5 μM ATP and 2 mM NaCl. However, upon modification with 20 μM DCCD only 33% of the molecules show significant fluctuations of the fluorescent dye indicating the specific inhibition of intersubunit rotation within coupled F₀F₁ ATPases.

At higher ATP concentrations (2.5 mM), discrete steps could no longer be resolved in detail (Fig. 3A–C, left column). For an improved time resolution, a shorter integration time was required which led to a decrease in the signal-to-noise ratio. Therefore, autocorrelation techniques were used for the analysis of fast rotational events. The ACFs derived from the polarization time series are depicted in the right column of Fig. 3, together with confidence intervals. They exhibit an exponential decay for short time lags and significant long-ranging oscillations with a periodicity of 10 ms (Fig. 3A) to 20 ms (Fig. 3C). The oscillations result from random stepping rotations, which are observed only for a relatively short time interval. They are more pronounced than expected for an ACF obtained during an infinite detection period (for a more detailed discussion see below). Rotational frequencies of 25–50 Hz correspond to ATP hydrolysis rates of 75–150 ATP s⁻¹ for different F₀F₁ enzymes with almost zero load. These data are consistent with ATP hydrolysis rates measured with the same F₀F₁ ensembles (100–120 s⁻¹).

The yield of rotating single molecules observed during ATP hydrolysis can be summarized as follows. About 90% of all ATPases analyzed exhibited a constant polarization in the absence or presence of ATP if no Na⁺ was added. From the selected molecules 72% rotated after the addition of 2 mM NaCl. This high yield as compared to other investigations (0.4–5% of rotating actin filaments) [17,18] could be explained by the much smaller size of the single dye label,
and therefore the minimal sterical interference between the label and the F0 part.

In vivo, the F0F1 holoenzyme is embedded in a phospholipid membrane and usually works in the ATP synthesis mode. To mimic physiological conditions and observe rotation during ATP synthesis, the Na\(^+\)-ATP synthase was incorporated in monodisperse liposomes (diameter 100 nm) and subsequently immobilized to Ni\(^2+\)-coated glass coverslips via the His\(^{10}\) tags on the \(\text{L}\) subunits (Fig. 4A). To start the ATP synthase activity, an electrochemical gradient has to be generated. In a first step, proteoliposomes loaded with 2 mM NaCl and potassium phosphate buffer were diluted into the same buffer containing in addition 2.5 mM ADP but no NaCl. At this stage, the electric potential is zero but a chemical Na\(^+\) gradient equivalent to approximately 60 mV is present. Under these conditions a constant non-zero polarization was observed that exhibited no correlation (Fig. 4B), suggesting that the proteoliposomes were immobilized on the glass surface via the His tags of the protein and that rotation coupled to ATP synthesis did not occur. In ensemble measurements using the same proteoliposome preparation, no ATP synthesis could be detected even at Na\(^+\) gradients of 120 mV and above. The addition of 200 mM KCl and 2 nM valinomycin triggers the electrogenic influx of potassium ions into the liposomes and establishes a diffusion potential of 95 mV. Under these conditions, large and fast fluctuations of the polarization signal were observed and the corresponding ACF exhibited periodic oscillations (Fig. 4C-E). These data indicate a rotation during ATP synthesis driven by a membrane potential but not by a Na\(^+\) concentration gradient only. They are therefore in complete accord with our previous observation of an obligatory requirement of the electrical component of the driving force for ATP synthesis [28-30]. The ACFs of the three sequentially recorded polarization traces show oscillations with increasing period length probably due to the continuous consumption of the sodium motive force. The ACFs showed periodicities between 20 and 40 ms indicating rotational frequencies of 12.5–25 Hz (Fig. 4C–E). This is consistent with experimentally obtained values from identical F\(_0\)F\(_1\) proteoliposomes indicating that ATP is indeed synthesized with rates of 45–60 ATP s\(^{-1}\) corresponding to rotational frequencies of 15–20 Hz. Furthermore, upon pre-incubation of the proteoliposomes with 20 mM DCCD, only small uncorrelated fluctuations were observed suggesting that rotation is again blocked by this F\(_0\)-specific inhibitor (Fig. 4F).

For a further interpretation of our observations we compared them to Monte Carlo simulations based on three different models suggested in the literature to describe the dynamics of this rotational motor. In the first single-molecule experiments on the F\(_1\) part three steps per full rotation were observed [16]. More recently Yashuda and coworkers succeeded in resolving each of these three steps as a sequence of one 90\(^\circ\) and one 30\(^\circ\) step following nucleotide binding and release [19]. Furthermore, it has been proposed that the rotation of the holoenzyme during ATP synthesis should reveal 11 steps per revolution [31] because of the 11-fold symmetry of the rotor within the F\(_0\) part of the enzyme [2].

Stepping rotation was simulated assuming an exponential distribution of dwell times in each step. The step width varies according to the model (\(3\times2\pi/3, 3\times(\pi/2+\pi/6), 11\times2\pi/11\)). From the stepping rotation the polarization and then the ACF were computed. The number of simulated data points was comparable to the number of experimental data points per polarization time series. All models lead to an ACF with a decay at short time lags and long-ranging oscillations. With
increasing number of steps per revolution the ACF exhibits more pronounced and more regular oscillations. The variety within one model from run to run is significant (see two examples as dashed lines in Fig. 5A), similar as in the experimental data. By visual inspection it was found that the simulation based on the three-double-step model is most consistent with our data. This finding suggests that the ATP/ADP turnover is rate-limiting under Na\(^{+}\) and nucleotide saturation conditions in both modes, ATP hydrolysis and synthesis.

The dynamics of a stochastic stepping rotor can also be described analytically using a Master equation approach [32]. The calculated ACF (Fig. 5A, solid line) comprises a modulated exponential decay, one pronounced minimum, and vanishing oscillatory contributions. The discrepancy between this analytic function and the simulations can be resolved by increasing the number of data points (Fig. 5A, dashed-dotted line). Moreover, also the experimental ACF eventually reproduces the analytic ACF when averaged over several individual molecules with similar stepping rates (Fig. 5B). Again, the three-double-step model fits slightly better to the data than the 11-step model. The reason for the appearance of more pronounced oscillations in the ACF of experimental and short simulated data sets is the limited number of observed revolutions. Therefore, from the expected bell-shaped distribution of rotation periods of a multi-step rotor [13] only a small subset of all possible periods is observed. With a high probability this undersampling leads to polarization traces that contain a narrow distribution of only a few frequencies (Fig. 5A, inset). Hence, the obtained ACFs comprise periodic correlations more long-ranging than the analytic solution would suggest.

In this single-fluorophore polarization study we have shown step-wise rotation of the rotor versus the stator in a functionally coupled Na\(^{+}\)-translocating F\(_{0}\)F\(_{1}\) complex. The presence of sodium ions is an obligatory precondition for ATP hydrolysis [33] and thus rotation. Although the single-chromophore approach contains several experimental difficulties, e.g. low number of detected photons and ambiguity about the direction of rotation, it enabled us to detect rotation in single F\(_{0}\)F\(_{1}\) molecules during ATP synthesis for the first time. Enzymes...
incorporated in liposomes showed no rotation in the presence of a Na\textsuperscript{+} concentration gradient only. However, after applying an electric potential difference, the rotor assembly turned with 12.5–25 Hz during ATP synthesis. This rotation was specifically inhibited by DCCD giving additional evidence that the enzyme used in this study was structurally and functionally the same as F\textsubscript{0}F\textsubscript{1} complexes assembled in vivo.

References