

## **F<sub>0</sub>F<sub>1</sub>-ATP synthase subunit $\alpha$ – A tale of two fragments**

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The F<sub>0</sub>F<sub>1</sub>-ATP synthase is a reversible nanomotor synthesizing ATP in bacteria and eukaryotic mitochondria. The core catalytic F<sub>1</sub> moiety of this multisubunit complex is formed by a globular hexamer of alternating subunits  $\alpha$  and  $\beta$  sitting on a central stalk consisting of subunit  $\gamma$  and small subunits  $\delta$  and  $\epsilon$  [1]. The composition and structure of the core F<sub>1</sub>-ATPase is believed to be strictly conserved throughout evolution [1], however this notion is based on the established structures of F<sub>0</sub>F<sub>1</sub>-ATP synthase complexes of bacteria and model eukaryotes and may not reflect full eukaryotic diversity. Several reports have indicated that the Euglenozoa F<sub>1</sub>-ATPase subunit  $\alpha$  is split into two fragments, presumably by proteolytic cleavage [2–5]. Both fragments stay associated with the complex. This feature appears to have no parallel in any other group of organisms. In this project, we are investigating whether  $\alpha$  cleavage results in novel features of this key enzyme, that are important for the structure/function of F<sub>0</sub>F<sub>1</sub>-ATP synthase in trypanosomes. We also aim to identify the protease responsible.

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