ESRF	Experiment title: FRANKFURT BAG: ATOMIC MECHANISMS OF MEMBRANE PROTEINS	Experiment number : MX-135		
Beamline:	Date of experiment:	Date of report:		
ID14-EH2	I cool contect(s):	Descined at ESDE:		
6	Dr. Elena MICOSSI	Kecelvea al ESKF:		
Names and affiliations of applicants (* indicates experimentalists): Mauro Mileni ^{*1} , Hanno Juhnke ^{*1} , C. Roy D. Lancaster ¹ Guohong Peng ^{*1} , Jürgen Koepke ¹ , Verena Linhard ¹ , Ulrike Wedemeyer ¹ , Vasundara Srinivasan ^{*1} , Hartmut Michel ¹ *Volker Zickermann ² , Carola Hunte ¹ ¹ Max Planck Institute of Biophysics, Department of Molecular Membrane Biology, Max-von-Laue-Str. 3, D-60438 Frankfurt am Main ² Universität Frankfurt, Fachbereich Medizin, Institut für Biochemie I, D-60590 Frankfurt/M. Germany				

Report:

Variant Photosynthetic Reaction Center from *Rhodopseudomonas viridis* (H. Juhnke*, C.R.D. Lancaster)

Two shifts of beam time were devoted to recording (ultimately) four data sets of a variant photosynthetic reaction center from *Rhodopseudomonas viridis*. The best data set is summarized in Table 1. However, the resolution of the resulting electron density maps was still insufficient to reliable discuss any differences to the wild-type structure, so higher resolution data are required.

Table 1. Best of four diffraction data sets collected at ESRF ID14-EH1 on a crystal of a variant *Rp. viridis* RC (P4₃2₁2, a = b = 223.5 Å, c = 113.6 Å)

	resol. range	measured	unique	complete	R _{sym}
	[Å]	reflections	reflections	[%]	[%]
pH7_AS195_1	50.0-2.50	478,663	99,099	98.5	8.8
	2.56-2.50	<i>27,170</i>	<i>6,456</i>	98.0	47.6

Quinol:Fumarate Reductase from Campylobacter jejuni (M. Mileni*, C.R.D. Lancaster)

Quinol:fumarate reductase (QFR), couples the reduction of fumarate to succinate to the oxidation of quinol to quinone, in a reaction opposite to that catalysed by mitochondrial complex II (succinate dehydrogenase). QFR from the anaerobic bacterium *Campylobacter jejuni* consists of three protein subunits, FrdA, FrdB, and FrdC. Crystals of this bioenergetically important membrane protein complex have previously been obtained in space group P1 with unit cell dimensions of a = 130.1 Å, b = 130.9 Å, c = 164.2 Å, and α = 108.6°, β = 90.6°, and γ = 118.5° and complete diffraction data to 3.9 Å has been collected in February 2003 at ID14 EH1 (see earlier report). More recently, a new crystal form of space group P2₁ with the unit cell dimensions a = 117.2 Å, b = 130.7 Å, c = 132.9 Å, β =108.0° has been obtained. The remaining shift of beamtime was devoted to (ultimately unsuccesful) attempts to improve the previously recorded data set of this crystal form at 3.24 Å resolution (see May 2004 EH1 report).

Outer Membrane Protein Complex Aq1862 from the Hyperthermophilic Eubacterium *Aquifex aeolicus*

(Guohong Peng*, Jürgen Koepke, Verena Linhard, Ulrike Wedemeyer, Hartmut Michel)

Aq1862 crystals were screened for lower mosaicity and higher resolution, three native datasets at 2.0 Å were collected.

Complex I from Yarrowia lipolytica

(Volker Zickermann*, Carola Hunte)

Complex I is the largest and least understood enzyme of the respiratory chain. Structural information is limited to low resolution and is based on electron microscopy of single particles and 2 D crystals. We have obtained crystals of complex I from the strictly aerobic yeast *Yarrowia lipolytica* with a monoclonal antibody fragment. Crystals are small and diffraction cannot be screened at the home source. 20 crystals were tested for diffraction. A few crystals diffracted up to 18-12 Å resolution. The pattern indicated a stability and/or a freezing problem. Optimization of crystallization conditions is in progress.

One (night) shift was lost due to ProDC interface problems (Vasundara Srinivasan*)