



	Experiment title: FRANKFURT BAG: ATOMIC MECHANISMS OF MEMBRANE PROTEINS	Experiment number: MX-135
Beamline: ID14-EH1	Date of experiment: from: 07-MAY-2004 8:30 to: 08-MAY-2004 8:00	Date of report: 15-Feb-2005
Shifts: 3	Local contact(s): Dr. Elena MICOSSI	<i>Received at ESRF:</i>

Names and affiliations of applicants (* indicates experimentalists):

Mauro Mileni*, Hanno Juhnke*, M. Gregor Madej*, C. Roy D. Lancaster
Max-Planck-Institute of Biophysics,
Department of Molecular Membrane Biology,
Max-von-Laue-Str. 3,
D-60438 Frankfurt am Main
e-mail: Roy.Lancaster@mpibp-frankfurt.mpg.de

Report:

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 $a = 130.1 \text{ \AA}$, $b = 130.9 \text{ \AA}$, $c = 164.2 \text{ \AA}$, and $\alpha = 108.6^\circ$, $\beta = 90.6^\circ$, and $\gamma = 118.5^\circ$ and complete diffraction data to 3.9 \AA has been collected in February 2003 at ID14 EH1 (see earlier report). During the two shifts available for this subproject, diffraction data on a new crystal form of space group $P2_1$ with the unit cell dimensions $a = 117.2 \text{ \AA}$, $b = 130.7 \text{ \AA}$, $c = 132.9 \text{ \AA}$, $\beta = 108.0^\circ$ was collected to 3.24 \AA resolution (Table 1) at $T = 4^\circ\text{C}$ from just one crystal. The phase problem has been solved by molecular replacement using the coordinates of the QFR from *Wolinella succinogenes* as a search model. The resulting structure is currently undergoing refinement.

Table 1. Diffraction data collected at ESRF ID14-EH1 on a crystal of *C. jejuni* QFR.

	resol. range [Å]	measured reflections	unique reflections	complete [%]	R _{sym} [%]
32_w2_0	70.0-3.24	241,397	61,387	99.2	8.2
	3.36-3.24	23,433	6,090	98.8	34.3

Other projects (H. Juhnke*, M.G. Madej*, C.R.D. Lancaster)

The remaining shift of beam time was devoted to (ultimately preliminary) attempts to record a diffraction data set of a variant photosynthetic reaction center from *Rhodospseudomonas viridis*.

Table 2. Preliminary diffraction data 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 [Å]	measured reflections	unique reflections	complete [%]	R _{sym} [%]
pH7_AS205_1	50.0-3.20	478,663	43,633	90.6	9.6
	3.27-3.20	11,301	2,380	75.0	19.6

References

[1] CRD. Lancaster, A Kröger, M Auer, H Michel (1999) *Nature* **402**, 377-385.