

**Experiment title:**

STRUCTURE DETERMINATION OF FTSY

Experiment number:

LS324

Beamline:

BL19

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3

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Report:

background:

FtsY is the receptor (or docking protein) for the signal recognition particle in *E. coli*. It has a molecular weight of about 54 kDa, but due to an unusual high content of glutamic acids in the N-terminal region it migrates with an apparent molecular weight of about 92 kDa on an SDS-PAGE. We obtained small crystals of the full length protein which we could improve only by modifying the construct, so that we expressed only a 40 kDa fragment containing the G-domain. This of course is homologous to other GTPases like p21 ras and eIF2. However, the SRP related GTPases form a distinct subgroup of GTPases. No structure of any member of that family has been solved so far.

report:

The crystals of FtsY diffracted to beyond 2Å, space group P21.

Cell dimensions: a=30, b=80, c=60 Å, beta=95.5°

We collected a native and several putative derivative data sets using 100 K, lambda=1Å. The native data were collected to 1.8Å resolution, but diffraction was seen to at least 1.5Å (couldn't collect high resolution due to time limitation). We also collected a IrCl₄,

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0s04, PtCl₄ and a AuCl data set. The completeness of the derivatives is about 91-96%, the native about 88% complete. The problem turned out to be non-isomorphism, especially a change in the beta-angle to 1° and more. The Au and the Pt data could be used together with two other derivatives (collected at EMBL-Heidelberg). However, the phases were not good enough to solve the structure.

We have now replaced the Met by Se-Met and we will perform a MAD experiment - which should not suffer from the non-isomorphism.