

ETS2

PDB:4MHV

Revision

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SGC Clone Accession:ETS2A-c023

Tag:Tag TEV-cleavable N-terminal his6 tag.

Host:BL21(DE3)-R3-pRARE2

Construct

Prelude:

Sequence:

mhhhhhhssgvdlgtenlyfq^SMKAVMSQALKATFSGFKKEQRRLGIPKNPWLWSEQQVCQWLLWATNEFSLVNVNLQRFGMNGQM
LCNLGKERFLELAPDFVGDILWEHLEQMIKEN

Vector:pNIC28-Bsa4. Details [[PDF](#)]; Sequence [[FASTA](#)] or [[GenBank](#)].

Growth

Medium:

Antibiotics:

Procedure:The expression plasmid was transformed into the host strain and plated on LB-agar containing 50 µg/ml kanamycin and 34 µg/ml chloramphenicol. Several colonies were combined to inoculate a 1ml culture in TB (+ 50 µg/ml kanamycin, 34 µg/ml chloramphenicol). The culture was grown overnight, glycerol was added to 15% v/v (from a 50% stock), and the resulting glycerol stock was frozen at -80°C in 100 µl aliquots. A loopful of cells from the glycerol stock was inoculated into 10-ml of TB medium containing 50 µg/ml kanamycin and 34 µg/ml chloramphenicol and grown overnight at 37°C. 1L of TB medium (containing 50 µg/ml kanamycin) was inoculated with 10 ml of the overnight culture and grown in 2.5L UltraYield baffled flasks until OD600 of 2.0. Cells were cooled to 18°C, IPTG added to 0.1 mM and growth continued at 18°C overnight. The cells were collected by centrifugation then the pellets were scraped out and transferred to 50-ml Falcon tubes and frozen at -80°C.

Purification

Procedure

Extraction

Procedure

Cell extraction : Frozen cell pellets (approx. 100g) were thawed briefly in a bath of warm water (20 - 37°C) then transferred to ice. Three volumes (i.e. 3 ml for every gram of cells) of lysis buffer was added, and the cells were resuspended by agitating and disrupted by pulsed sonication (10 sec on 20 sec off) for 20 minutes on ice. Polyethylenimine was added to the lysate at a final concentration of 0.15 % and lysates were centrifuged for 60 minutes at 50,000xg to remove cellular debris.

Lysis buffer: 50 mM HEPES, pH 7.5, 500 mM NaCl, 5% glycerol, 1 mM TCEP, 10 mM Imidazole

Column 1 : 3ml bed volume Ni-IDA

Solutions:

Wash buffer: 50 mM HEPES, pH 7.5, 500 mM NaCl, 30 mM imidazole, 5% Glycerol, 1 mM TCEP.

Elution buffer: 50 mM HEPES, pH 7.5, 500 mM NaCl, 300 mM imidazole, 5% Glycerol, 1 mM TCEP.

Procedure: The cell extract was incubated with 5ml of Ni-IDA resin (GE Healthcare) for 1hr at 4°C with gentle agitation. The resin was pelleted by centrifugation (700g for 6 min), and the supernatant was decanted. The resin was washed initially with 10 column volumes of lysis buffer and pelleted by centrifugation. Ten column volumes of wash buffer were added and the resuspended resin was transferred to a gravity flow column and washed with 5 column volumes of wash buffer. Proteins were eluted by addition of 2 x 10 ml of elution buffer and combined with the fractions from the wash buffer.

Column 2 : Gel filtration, Hiload 16/60 Superdex S75 prep grade, 120 ml (GE Healthcare)

GF buffer: 50 mM HEPES, pH 7.5, 500 mM NaCl, 5% Glycerol, 1 mM TCEP.

Procedure: The fractions from Ni-IDA column were concentrated to 2ml using a 3 KDa MWCO centrifugal concentrator and loaded on the gel filtration column in GF buffer at 1.2 ml/min. Eluted proteins were collected in 2-ml fractions and analysed on SDS-PAGE.

Enzymatic treatment and Column 3: The N-terminal His6-tag was cleaved by incubating the protein overnight with tobacco etch virus (TEV) protease (1:42 ratio at 4°C) whilst being dialyzed into dialysis buffer (50 mM HEPES, 500 mM NaCl, 5 % Glycerol, 1 mM TCEP) using a 3,500 MWCO snakeskin dialysis membrane. Cleaved protein was purified by passing over a 5 ml pre-equilibrated 50% Ni-IDA bead solution. Elution was done in GF buffer supplemented with step gradients of 20 mM, 40 mM 100 mM and 300 mM imidazole, with flowthrough and fractions collected.

Column 4 : Gel filtration, Hiload 16/60 Superdex S75 prep grade, 120 ml (GE Healthcare)

GF buffer: 50 mM HEPES, pH 7.5, 500 mM NaCl, 5% Glycerol, 1 mM TCEP.

Procedure: The fractions from Ni-IDA column were concentrated to 2ml using a 3 KDa MWCO centrifugal concentrator and loaded on the gel filtration column in GF buffer at 1.2 ml/min. Eluted proteins were collected in 2-ml fractions and analysed on SDS-PAGE.

Concentration: The cleaved purified protein was concentrated using a 3 KDa MWCO centrifugal concentrator to 24 mg/ml and stored at 4°C. The protein concentration was determined spectrophotometrically using $\epsilon_{280} = 27500$

Ligand

MassSpec: Observed mass of protein was 11373.1 calculated mass is 11373.2

Crystallization: Crystallisation experiments were performed using the sitting drop vapour diffusion technique. Crystals of the Ets-2 PNT domain grew at 4°C from conditions containing

14.4%(w/v) PEG 10000, 20% glycerol, 0.16M calcium acetate, 0.1M cacodylate pH 6.5. Crystals were loop mounted and directly plunged into liquid nitrogen.

NMR Spectroscopy:

Data Collection: The structure was solved by multiple isomorphous replacement using mercury and platinum derivative crystals prepared by soaking crystals for 15 minutes in well solution supplemented with 10 mM thiomersal and 10 mM potassium tetrachloroplatinate respectively. Native and derivative data were collected at Diamond Light Source, beamlines I04 and I04-1, processed using XDS and the structure was solved by multiple isomorphous replacement using the program autoSHARP. Model building was performed using the program COOT and the model refinement was performed using BUSTER to a final Rfactor=23.1 % , Rfree =23.6 %.

Data Processing: