

# SDCCAG10

PDB:2HQ6

## Revision

**Revision Type:**created

**Revised by:**created

**Revision Date:**created

**Entry Clone Accession:**

**Entry Clone Source:**MGC

**SGC Clone Accession:**ppi70.008.176:B6; plate SDC066 B6

**Tag:**

**Host:**BL21 (DE3)

## Construct

**Prelude:**

**Sequence:**

mgsshhhhhssglvprgsEPPTNGKVLLKTTAGDIDIELWSKEAPKACRNFIQLCLEAYDNTIFHRVVPGFIVQGGDPTGTGSGG  
ESIYGAPFKDEFHSRLRFNRRGLVAMANAGSHDNGSQFFFTLGRADELNNKHTIFGKVTGDTVYNMLRLSEVDIDDDERPHNPHKIK  
SCEVLFPFDD

**Vector:**

## Growth

**Medium:**

**Antibiotics:**

**Procedure:**Using the LEX bubbling system, the protein was expressed in E. coli BL21 (DE3) grown in Terrific Broth (TB) in the presence of 50 microG/mL of kanamycin at 3deg°C to an OD600 of 7.5. Cells were then induced by isopropyl-1-thio-D-galactopyranoside (IPTG), final concentration 0.05 mM, and incubated overnight at 15degC. The culture was centrifuged and the cell pellets were collected and stored at -80degC.

## Purification

**Procedure**

IMAC purification: 4 microL of clarified supernatant is reserved for later analysis by SDS-PAGE. The rest of the clarified supernatant is then diluted 1:2 in lysis buffer, and loaded at approximately 1mL/min by gravity onto 5 mL of Ni-NTA resin (Qiagen 30450). 5 column volumes of lysis buffer are used to wash the column at approximately 3 mL/min, followed by 5 column volumes of low imidazole buffer (lysis buffer + 10 mM Imidazole (VWR EM-5720) pH 8) at approximately 3 mL/min. A 4 microL sample of the low imidazole wash is saved for later analysis by SDS-PAGE. Samples are eluted from the Ni-NTA resin by exposure to 10 mL elution

buffer (lysis buffer + 250 mM imidazole and 10% glycerol (EMD GX0185-5)) at 1mL/min flow rate. A 10 microL sample of the eluate is saved for SDS-PAGE analysis. 10 microL of each eluate is saved for measurement of protein concentration using Bradford reagent (BioRad 500-0202).

## **Extraction**

### **Procedure**

The cell pellet was resuspended in lysis buffer, inhibitor (0.1 microM phenylmethyl sulfonyl fluoride, PMSF) and lysed using Microfluidizer. The lysate was cleared by centrifugation.

### **Concentration:**

### **Ligand**

### **MassSpec:**

**Crystallization:** Protein concentration: 20 mg/mL; 20% Peg 3350, 0.2M NaI; 50 mM Tris pH 8.0, 500 mM NaCl, 5 mM BME; hanging drop; 1+1; cryo: 20% glycerol; 298.0K

### **NMR Spectroscopy:**

### **Data Collection:**

### **Data Processing:**