

# Cp-CAP: *Cryptosporidium parvum* cyclase-associated protein

PDB:2B0R

## Revision

**Revision Type:**created

**Revised by:**created

**Revision Date:**created

**Entry Clone Accession:**[cgd5\\_440](#)

**Entry Clone Source:***Cryptosporidium parvum* strain Iowa genomic DNA

**SGC Clone Accession:**CP-PFA0260c; plate MR:H4

**Tag:**N-terminal: His-tag with integrated thrombin protease site: MGSSHHHHHHSSGLVPR\*GS

**Host:***E. coli* BL21-(DE3)-CodonPlus-RIL from Stratagene

## Construct

**Prelude:**

**Sequence:**

```
mgsshhhhhssglvprgskSQIYLKKEKKMKAARQVVTNGSPKVELQKDTYLVENHVNCADPITLSEGSIKNKVSVRCSQNSRIIV  
EQKVNSIFIENCVGCIFLVNGVISSIEIVNCDDIKLQMTGIVPTISLDKSNKVNIIYTSKEGKNVEVYSSKSSEMNLFPGEEDWK  
ELAIPEQFVTKYNESKGLKLESMVSPLYG
```

**Vector:**pET28a-LIC

## Growth

**Medium:**Terrific Broth (TB)

**Antibiotics:**50 microG/mL kanamycin and 25 microG/mL chloramphenicol

**Procedure:**A single colony was inoculated into 10 mL of LB with of Antibiotics and incubated with shaking at 250 rpm overnight at 37 degC. The culture was transferred into 50 mL of TB with Antibiotics in a 250 mL shaking flask and incubated at 37 degC for 3 hours. The culture was then transferred into 1.8 L of above-specified growth medium with Antibiotics and 0.3 mL of antifoam (Sigma) in a 2L bottle and cultured using the LEX system to an OD600 of 5, cooled to 15 degC and induced with 0.5 mM isopropyl-1-thio-D-galactopyranoside (IPTG) overnight at 15 degC.

## Purification

**Procedure**

The cleared cell lysate was loaded onto a DE52 (Whatman) column packed with 10 g of resin (previously activated with 2.5 M NaCl and equilibrated with Binding Buffer), and subsequently onto a 2.5 mL Ni-NTA column at approximately 1.5 mL/min. When all the lysate was loaded, 20 mL of Binding Buffer was added to the DE52 column. Then the Ni-NTA column was washed with 200 mL of Wash Buffer at 2.5 mL/min. After washing, the protein was eluted from the

Ni-NTA column with 15 mL of Elution Buffer. EDTA was added immediately to 1 mM. DTT was then added to 1 mM 15 minutes later.

The eluted protein was applied to a Sephadex S200 26/60 gel filtration column pre-equilibrated with Gel Filtration Buffer. The collected fractions corresponding to the eluted protein peak were concentrated using a 15 mL Amicon Ultra centrifugal filter device from Millipore (10 kD cutoff). The concentrated sample was flash frozen in N<sub>2</sub>(l) and stored at -80 degC.

## **Extraction**

### **Procedure**

Cells were resuspended to approximately 40 mL/L of cell culture in Binding Buffer with protease inhibitor (1 mM benzamidine-HCl and 1 mM phenylmethyl sulfonyl fluoride, PMSF).

Resuspended pellets stored at -80 degC were thawed overnight at 4 degC on the day before purification. Prior to mechanical lysis, each pellet from 1 L of culture was pretreated with 0.5% CHAPS and 500 units of benzonase for 40 minutes at room temperature. Cells were mechanically lysed with a microfluidizer (Microfluidizer Processor, M-110EH) at approximately 18,000 psi. The cell lysate was centrifuged at ~75,000 x g for 20 minutes at 10 degC.

**Concentration:** 30 mg/mL for uncleaved protein and 16.9 mg/mL for cleaved protein

### **Ligand**

### **MassSpec:**

**Crystallization:** The protein was crystallized by means of hanging drop vapor diffusion in Linbro plate. The plate was set with 1.5 microL cleaved protein and 1.5 microL buffer in each drop, and 300 microL reservoir volume per well. Crystals grew overnight in 17 % Peg 3350 and 200 mM diammonium tartrate at 18 degC.

### **NMR Spectroscopy:**

### **Data Collection:**

### **Data Processing:**