

# SULT1C2

**PDB:**2AD1

## Revision

**Revision Type:**created

**Revised by:**created

**Revision Date:**created

**Entry Clone Accession:**gi: 28830308

**Entry Clone Source:**MGC

**SGC Clone Accession:**

**Tag:**N-terminal: His-tag with integrated thrombin protease site: MGSSHHHHHSSGLVPRGS

**Host:**E.coli BL21 (DE3) codon plus RIL (Stratagen).

## Construct

**Prelude:**

**Sequence:**

**Vector:**p28a-LIC

## Growth

**Medium:**

**Antibiotics:**

**Procedure:**

## Purification

### Procedure

## Extraction

### Procedure

Cells were harvested by centrifugation at 6,000 rpm. The cell pellets were frozen in liquid nitrogen and stored at -80°C. For the purification the cell paste was thawed and resuspended in lysis buffer (50 mM HEPES-NaOH, pH 7.4, 0.5 M NaCl, 5 mM imidazol, 2 mM  $\beta$ -mercaptoethanol, 5% glycerol) with protease inhibitor (0.1mM phenylmethyl sulfonyl fluoride, PMSF). The cells were lysed by passing through Microfluidizer (Microfluidics Corp.) at 20,000 psi.

**Concentration:**

**Ligand**

**MassSpec:**

**Crystallization:** Purified SULT1C2 was crystallized using the hanging drop method at 20°C by mixing 2  $\mu$ l of the protein solution with 2  $\mu$ l of the reservoir solution containing 14-20% polyethylene glycol 3350, 0.2 M Li-Citrate, 0.1 M Na-Citrate, pH 4.6.

**NMR Spectroscopy:****Data Collection:****Data Processing:**