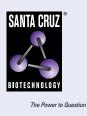
# SANTA CRUZ BIOTECHNOLOGY, INC.

# TCP-1 β (F39 P7 F11): sc-47717



#### BACKGROUND

The protein TCP-1 (t complex polypeptide 1) is a subunit of the hetero-oligomeric complex CCT (chaperonin containing TCP-1) present in the eukaryotic cytosol. The CCT of eukaryotic cytosol is composed of eight different subunit species, TCP-1  $\alpha$ ,  $\beta$ ,  $\gamma$ ,  $\delta$ ,  $\epsilon$ ,  $\zeta$ ,  $\eta$  and  $\theta$ , each encoded by a different gene. Two  $\zeta$  subunits have been described: TCP-1  $\zeta$  (also designated TCP-1  $\zeta$ 1) and TCP-1  $\zeta$ 2. TCP-1 subunits are proposed to have independent functions in folding its *in vivo* substrates, the Actins and Tubulins. TCP-1 was first identified in the mouse as relevant for tail-less and embryonic lethal phenotypes. Sequences homologous to TCP-1 have been isolated in several other species, and the yeast TCP-1 has been shown to encode a molecular chaperone for Actin and Tubulin. TCP-1 found in mammalian cells and yeast plays an important role in the folding of cytosolic proteins.

### **REFERENCES**

- Ahnert, V., et al. 1996. Cucumber T-complex protein. Molecular cloning, bacterial expression and characterization within a 22-S cytosolic complex in cotyledons and hypocotyls. Eur. J. Biochem. 235: 114-119.
- 2. lijima, M., et al. 1998. A *Dictyostelium discoideum* homologue to TCP-1 is essential for growth and development. Gene 213: 101-106.
- Ritco-Vonsovici, M. and Willison, K.R. 2000. Defining the eukaryotic cytosolic chaperonin-binding sites in human Tubulins. J. Mol. Biol. 304: 81-98.

#### **CHROMOSOMAL LOCATION**

Genetic locus: CCT2 (human) mapping to 12q15; Cct2 (mouse) mapping to 10 D2.

## SOURCE

TCP-1  $\beta$  (F39 P7 F11) is a mouse monoclonal antibody raised against ovalbumin-conjugate synthetic peptide RKRVPDHHPC.

## PRODUCT

Each vial contains 200  $\mu g$  IgG\_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

## **APPLICATIONS**

TCP-1  $\beta$  (F39 P7 F11) is recommended for detection of TCP-1  $\beta$  of mouse, rat, human and *Xenopus laevis* origin by Western Blotting (starting dilution 1:100, dilution range ), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

Suitable for use as control antibody for TCP-1  $\beta$  siRNA (h): sc-36622, TCP-1  $\beta$  siRNA (m): sc-36625, TCP-1  $\beta$  shRNA Plasmid (h): sc-36622-SH, TCP-1  $\beta$  shRNA Plasmid (m): sc-36625-SH, TCP-1  $\beta$  shRNA (h) Lentiviral Particles: sc-36622-V and TCP-1  $\beta$  shRNA (m) Lentiviral Particles: sc-36625-V.

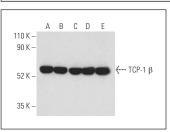
Molecular Weight of TCP-1  $\beta$ : 50 kDa.

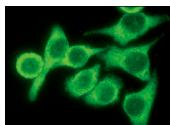
Positive Controls: Caki-1 cell lysate: sc-2224, HEL 92.1.7 cell lysate: sc-2270 or F9 cell lysate: sc-2245.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>™</sup> Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850. 4) Immunohistochemistry: use m-IgGκ BP-HRP: sc-516102 with DAB, 50X: sc-24982 and Immunohistomount: sc-45086, or Organo/Limonene Mount: sc-45087.

#### DATA





TCP-1  $\beta$  (F39 P7 F11): sc-47717. Western blot analysis of TCP-1  $\beta$  expression in HEL 92.1.7 (**A**), Caki-1 (**B**), ZR-75-1 (**C**), F9 (**D**) and NIH/3T3 (**E**) whole cell lysates Detection reagent used: m-lgG<sub>28</sub> BP-HRP: sc-542731

TCP-1  $\beta$  (F39 P7 F11): sc-47717. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

## **SELECT PRODUCT CITATIONS**

- Song, H. and Sokolov, M. 2009. Analysis of protein expression and compartmentalization in retinal neurons using serial tangential sectioning of the retina. J. Proteome Res. 8: 346-351.
- Tashiro, E., et al. 2013. Prefoldin protects neuronal cells from polyglutamine toxicity by preventing aggregation formation. J. Biol. Chem. 288: 19958-19972.
- Gao, X., et al. 2013. Splice isoforms of phosducin-like protein control the expression of heterotrimeric G proteins. J. Biol. Chem. 288: 25760-25768.

## **STORAGE**

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

### PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.