# SANTA CRUZ BIOTECHNOLOGY, INC.

# TIRAP (B-3): sc-166151



## BACKGROUND

Mammalian Toll-like receptors (TLRs) recognize conserved products of microbial metabolism and activate NF $\kappa$ B and other signaling pathways through the adapter protein MyD88. MyD88 functions as an adapter protein in the association of IL-1 receptor associated kinase (IRAK) with the IL-1 receptor. MyD88 contains a characteristic N-terminal death domain, which is essential for NF $\kappa$ B activation and an adjacent Toll/II-1R homology domain (TIR domain), which is responsible for signal transduction. TIR domain-containing adapter protein (TIRAP), also designated MAL (MyD88 adapter-like), wyatt or TLR-4 adaptor protein, is a cytoplasmic TIR-domain-containing protein that activates NF $\kappa$ B, Jun N-terminal kinase and extracellular signal-regulated kinase-1 and -2. TIRAP forms homodimers and heterodimers with MyD88. IRAK-2, but not IRAK, is required for the activation of NF $\kappa$ B by TIRAP which associates with IRAK-2 through the TIR domain. In addition, TIRAP associates with TLR-4, suggesting that it plays a role in TLR-4 signal transduction.

#### REFERENCES

- Medzhitov, R., et al. 1998. MyD88 is an adaptor protein in the hToll/IL-1 receptor family signaling pathways. Mol. Cell 2: 253-258.
- Burns, K., et al. 1998. MyD88, an adaptor protein involved in interleukin-1 signaling. J. Biol. Chem. 273: 12203-12209.
- Chow, J.C., et al. 1999. Toll-like receptor-4 mediates lipopolysaccharideinduced signal transduction. J. Biol. Chem. 274: 10689-10692.
- 4. Means, T.K., et al. 2000. The biology of Toll-like receptors. Cytokine Growth Factor Rev. 11: 219-232.
- Horng, T., et al. 2001. TIRAP: an adapter molecule in the Toll signaling pathway. Nat. Immunol. 2: 835-841.
- Fitzgerald, K.A., et al. 2001. Mal (MyD88-adapter-like) is required for Tolllike receptor-4 signal transduction. Nature 413: 78-83.

#### CHROMOSOMAL LOCATION

Genetic locus: TIRAP (human) mapping to 11q24.2.

## SOURCE

TIRAP (B-3) is a mouse monoclonal antibody raised against amino acids 1-235 representing full length TIRAP of human origin.

# PRODUCT

Each vial contains 200  $\mu$ g lgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

# STORAGE

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

## PROTOCOLS

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

#### **APPLICATIONS**

TIRAP (B-3) is recommended for detection of TIRAP of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), 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 solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for TIRAP siRNA (h): sc-42932, TIRAP shRNA Plasmid (h): sc-42932-SH and TIRAP shRNA (h) Lentiviral Particles: sc-42932-V.

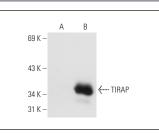
Molecular Weight of TIRAP: 36 kDa.

Positive Controls: Caki-1 cell lysate: sc-2224, ES-2 cell lysate: sc-24674 or TIRAP (h): 293T Lysate: sc-114950.

### **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.

## DATA



TIRAP (B-3): sc-166151. Western blot analysis of TIRAP expression in non-transfected: sc-117752 (A) and human TIRAP transfected: sc-114950 (B) 293T whole cell lysates.

## SELECT PRODUCT CITATIONS

 Israel, L., et al. 2017. Human adaptive immunity rescues an inborn error of innate immunity. Cell 168: 789-800.

#### **RESEARCH USE**

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