

# TRRAP (T-17): sc-5405

## BACKGROUND

The transcription factors c-Myc and E2F are involved in regulating cell cycle progression. Overexpression of c-Myc in certain cell types induces non-cycling cells to enter the cell cycle via a mechanism involving E2F-1. E2F-1 is thought to regulate c-Myc expression via interactions with the retinoblastoma protein. TRRAP (for transformation/transcription domain-associated protein) interacts specifically with both c-Myc and E2F-1. Expression of *trans*-activated mutant TRRAP inhibits the oncogenic transformation of both c-Myc and E2F-1, suggesting that TRRAP is required for these oncogenic transcription factor pathways. TRRAP shares homology with the ATM/PI 3-kinase family, and it is highly conserved in evolution.

## CHROMOSOMAL LOCATION

Genetic locus: TRRAP (human) mapping to 7q22.1; Trrap (mouse) mapping to 5 G2.

## SOURCE

TRRAP (T-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of TRRAP of human origin.

## PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-5405 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## APPLICATIONS

TRRAP (T-17) is recommended for detection of TRRAP of mouse, rat and human origin by Western Blotting (starting dilution 1:200, 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).

TRRAP (T-17) is also recommended for detection of TRRAP in additional species, including canine, bovine, porcine and avian.

Suitable for use as control antibody for TRRAP siRNA (h): sc-36746, TRRAP siRNA (m): sc-36747, TRRAP shRNA Plasmid (h): sc-36746-SH, TRRAP shRNA Plasmid (m): sc-36747-SH, TRRAP shRNA (h) Lentiviral Particles: sc-36746-V and TRRAP shRNA (m) Lentiviral Particles: sc-36747-V.

Molecular Weight of TRRAP: 434 kDa.

Positive Controls: HeLa nuclear extract: sc-2120, K-562 nuclear extract: sc-2130 or Jurkat nuclear extract: sc-2132.

## RESEARCH USE

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

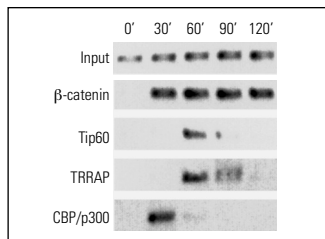
## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.

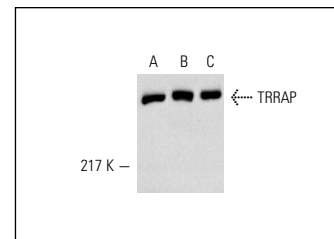
## STORAGE

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

## DATA



ChIP analysis of coactivator recruitment on Cyclin D2 promoter in C2C12 cells treated with LiCl and serum. Antibodies tested include β-catenin (H-102): sc-7199, β-catenin (C-18): sc-1496, β-catenin (E-5): sc-7963, Tip60 (N-17): sc-5725, TRRAP (T-17): sc-5405, TRRAP (Y-18): sc-12375, TRRAP (F-20): sc-12376, TRRAP (H-300): sc-11411, CBP (A-22): sc-369, CBP (C-20): sc-583, CBP (451): sc-1211, CPB (C-1): sc-7300, p300 (H-272): sc-8981, p300 (N-15): sc-584 and p300 (C-20): sc-585. Data kindly provided by M.G. Rosenfeld and reproduced with permission from Kioussi et al., Cell 2002, 111: 673-685.



TRRAP (T-17): sc-5405. Western blot analysis of TRRAP expression in HeLa (A), K-562 (B) and Jurkat (C) nuclear extracts.

## SELECT PRODUCT CITATIONS

- Lang, S., et al. 2001. E2F transcriptional activation requires TRRAP and GCN5 cofactors. J. Biol. Chem. 276: 32627-32634.
- Nikiforov, M.A., et al. 2002. TRRAP-dependent and TRRAP-independent transcriptional activation by Myc family oncoproteins. Mol. Cell. Biol. 22: 5054-5063.
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- Dai, M.S., et al. 2010. Ribosomal protein L11 associates with c-Myc at 5 S rRNA and tRNA genes and regulates their expression. J. Biol. Chem. 285: 12587-12594.
- Huang, L., et al. 2011. Prevention of transcriptional silencing by a replicator-binding complex consisting of SWI/SNF, MeCP1, and hnRNP C1/C2. Mol. Cell. Biol. 31: 3472-3484.