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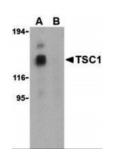
HIGH PERFORMANCE ANTIBODIES ... AND MORE

**ProSci Incorporated** 12170 Flint Place Poway, CA 92064 Toll Free: +1 (888) 513 9525 Local: +1 (858) 513 2638 Fax: +1 (858) 513 2692

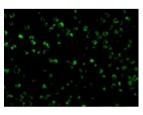
techsupport@prosci-inc.com

## **TSC1 Antibody**

CATALOG NUMBER: 3505



Western blot analysis of TSC1 in EL4 cell lysate with TSC1 antibody at 1 ug/mL in the (A) absence and (B) presence of blocking peptide.



Immunofluorescence of TSC1 in L1210 cells with TSC1 antibody at 10 ug/mL.



Immunocytochemistry of TSC1 in EL4 cells with TSC1 antibody at 2 ug/mL

Specifications	
SPECIES REACTIVITY:	Human, Mouse, Rat
TESTED APPLICATIONS:	ELISA, ICC, IF, WB
APPLICATIONS:	TSC1 antibody can be used for the detection of TSC1 by Western blot at 1 ug/mL. Antibody can also be used for immunocytochemistry starting at 2 ug/mL. For immunofluorescence start at 2 ug/mL.
USER NOTE:	Optimal dilutions for each application to be determined by the researcher.
POSITIVE CONTROL:	1) Cat. No. 1287 - EL4 Cell Lysate
PREDICTED MOLECULAR WEIGHT:	Predicted: 128 kDa
	Observed: 135 kDa
SPECIFICITY:	At least two isoforms of TSC1 are known to exist; this antibody will detect both isoforms. TSC1 antibody is predicted to not cross-react with TSC2.
IMMUNOGEN:	TSC1 antibody was raised against a 15 amino acid synthetic peptide from the middle region of human TSC1.
	The immunogen is located within amino acids 220 - 270 of TSC1.
HOST SPECIES:	Rabbit
Properties	
PURIFICATION:	TSC1 Antibody is affinity chromatography purified via peptide column.
PHYSICAL STATE:	Liquid
BUFFER:	TSC1 Antibody is supplied in PBS containing 0.02% sodium azide.
CONCENTRATION:	1 mg/mL
STORAGE CONDITIONS:	TSC1 antibody can be stored at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high
	temperatures.
CLONALITY:	temperatures. Polyclonal

CONJUGATE:	Unconjugated
Additional Info	
ALTERNATE NAMES:	TSC1 Antibody: LAM, TSC, KIAA0243, Hamartin, Tuberous sclerosis 1 protein
ACCESSION NO.:	AAC51674
PROTEIN GI NO.:	2331281
OFFICIAL SYMBOL:	TSC1
GENE ID:	7248
Background	
BACKGROUND:	TSC1 Antibody: Tuberous sclerosis complex (TSC) is an autosomal dominant tumor syndrome caused by mutations in either of the TSC1 or TSC2 tumor suppressor genes. The products of these genes form a protein complex that indirectly decreases the signaling of the mammalian Target of Rapamycin (TOR), an evolutionarily conserved serine/threonine kinase that regulates cell growth and cell cycle through its ability to integrate signals from nutrient levels and growth factors. TOR activity is stimulated by Rheb, a member of the Ras superfamily of G-proteins, when the GTP/GDP ratio bound to Rheb is high. Immunoprecipitated TSC1/TSC2 has been shown to stimulate TOR activity. This is supported by experiments showing overexpression of TSC1 and TSC2 results in a significant decrease in the GTP/GDP ratio bound to Rheb and the inhibition of cell growth. A shorter 40 kDa isoform of TSC1 has been shown to exist but its function is unknown.
REFERENCES:	1) Shamji AF, Ngheim P, and Schreiber SL. Integration of growth factor and nutrient signaling: implications for cancer biology. Mol. Cell 2003; 12:271-80.
	2) Inoki K, Ouyang H, Li Y, et al. Signaling by target of rapamycin proteins in cell growth control. Microbiol. Mol. Biol. Rev. 2005; 69:79-100.
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	4) Inoki K, Li Y, Xu T, et al. Rheb GTPase is a direct target of TSC2 GAP activity and regulates mTOR signaling. Genes Dev. 2003; 17:1829-34.

FOR RESEARCH USE ONLY

December 12, 2016