SANTA CRUZ BIOTECHNOLOGY, INC.

Shrm (T-17): sc-10309



The Power to Question

BACKGROUND

The gene shrm encodes a PDZ domain protein which regulates aspects of cytoarchitecture required for proper neuralation. PDZ domains mediate protein-protein interactions which facilitate membrane protein localization and signaling complex assembly. Mutation of the mouse Shrm causes neural tube defects (NTDs) attributed to failure of the neural tube to close during development. Targeted mutation studies have identified a number of factors which regulate neural tube morphogenesis. Shrm is strongly expressed in neural epithelium at the time of cranial tube closure. Shrm is a cytoskeletal protein with a size of ~205 kDa which localizes to adherens junctions and directly binds F-Actin. The Shrm protein can exist in a short and long form, ShrmS and ShrmL respectively.

REFERENCES

- Chen, Z.F. and Behringer, R.R. 1995. twist is required in head mesenchyme for cranial neural tube morphogenesis. Genes Dev. 9: 686-699.
- Ponting, C.P., Phillips, C., Davies, K.E., and Blake, D.J. 1997. PDZ domains: targeting signalling molecules to sub-membranous sites. Bioessays 19: 469-479.
- Songyang, Z., Fanning, A.S., Fu, C., Xu, J., Marfatia, S.M., Chishti, A.H., Crompton, A., Chan, A.C., Anderson, J.M., and Cantley, L.C. 1997. Recognition of unique carboxyl-terminal motifs by distinct PDZ domains. Science 275: 73-77.
- Hildebrand, J.D. and Soriano, P. 1999. Shroom, a PDZ domain-containing actin-binding protein, is required for neural tube morphogenesis in mice. Cell 99: 485-497.
- Kuan, C.Y., Yang, D.D., Samanta Roy. D.R., Davis, R.J., Rakic, P., and Flavell, R.A. 1999. The Jnk1 and Jnk2 protein kinases are required for regional specific apoptosis during early brain development. Neuron 22: 667-676.

CHROMOSOMAL LOCATION

Genetic locus: SHROOM3 (human) mapping to 4q21.1; Shrm (mouse) mapping to 5 E3.

SOURCE

Shrm (T-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of Shrm of mouse origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-10309 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

Store at 4° C, **D0 NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

Shrm (T-17) is recommended for detection of Shrm long and short forms 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).

Suitable for use as control antibody for Shrm siRNA (h): sc-42248, Shrm siRNA (m): sc-42249, Shrm shRNA Plasmid (h): sc-42248-SH, Shrm shRNA Plasmid (m): sc-42249-SH, Shrm shRNA (h) Lentiviral Particles: sc-42248-V and Shrm shRNA (m) Lentiviral Particles: sc-42249-V.

Molecular Weight of Shrm: 205 kDa.

Positive Controls: H4 cell lysate: sc-2408, IMR-32 cell lysate: sc-2409 or T98G cell lysate: sc-2294.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 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 donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA





Shrm (T-17): sc-10309. Western blot analysis of Shrm expression in H4 (A) IMR-32 (B) and T98G (C) whole cell lysates.

Shrm (T-17): sc-10309. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

SELECT PRODUCT CITATIONS

 Taylor, J., et al. 2008. The scaffold protein POSH regulates axon outgrowth. Mol. Biol. Cell 19: 5181-5192.

RESEARCH USE

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

PROTOCOLS

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