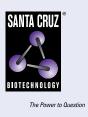
SANTA CRUZ BIOTECHNOLOGY, INC.

Sprouty 1 (H-2): sc-365520



BACKGROUND

Members of the sprouty family (Sprouty 1-4) are inducible negative regulators of growth factors that act through tyrosine kinase receptors. Mammalian Sprouty homologs share a well-conserved cysteine-rich C-terminal domain with their Drosophila counterparts. Both Sprouty 1 and 2 are anchored to membranes by palmitoylation, associate with caveolin-1 in perinuclear and vesicular structures, and are phosphorylated on serine residues. Upon stimulation, a subset is recruited to the leading edge of the plasma membrane. Sprouty 2 can associate with c-Cbl, a downregulator of RTK signaling, and inhibits the activities of several growth factors. Sprouty 2 also functions as a negative regulator of embryonic lung morphogenesis and growth. The wellconserved C-terminus of sprouty contains two domains which are necessary for Sprouty 2 co-localization with microtubules and translocation to membrane ruffles. In addition, the C-terminus is required for the inhibition of cell migration and proliferation. In conclusion, members of sprouty inhibit FGF and VEGF-mediated cell proliferation, suggesting that they may regulate angiogenesis in normal and disease processes.

REFERENCES

- Lim, J., et al. 2000. Sprouty proteins are targeted to membrane ruffles upon growth factor receptor tyrosine kinase activation. Identification of a novel translocation domain. J. Biol. Chem. 275: 32837-32845.
- Impagnatiello, M.A., et al. 2001. Mammalian Sprouty 1 and 2 are membrane-anchored phosphoprotein inhibitors of growth factor signaling in endothelial cells. J. Cell Biol. 152: 1087-1098.
- Ozaki, K., et al. 2001. Erk pathway positively regulates the expression of sprouty genes. Biochem. Biophys. Res. Commun. 285: 1084-1088.

CHROMOSOMAL LOCATION

Genetic locus: SPRY1 (human) mapping to 4q28.1.

SOURCE

Sprouty 1 (H-2) is a mouse monoclonal antibody raised against amino acids 61-180 mapping within an internal region of Sprouty 1 of human origin.

PRODUCT

Each vial contains 200 μg IgG1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Sprouty 1 (H-2) is available conjugated to agarose (sc-365520 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-365520 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-365520 PE), fluorescein (sc-365520 FITC), Alexa Fluor[®] 488 (sc-365520 AF488), Alexa Fluor[®] 546 (sc-365520 AF546), Alexa Fluor[®] 594 (sc-365520 AF594) or Alexa Fluor[®] 647 (sc-365520 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-365520 AF680) or Alexa Fluor[®] 790 (sc-365520 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

RESEARCH USE

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

APPLICATIONS

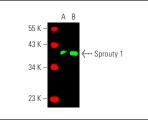
Sprouty 1 (H-2) is recommended for detection of Sprouty 1 of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

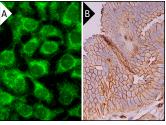
Suitable for use as control antibody for Sprouty 1 siRNA (h): sc-41035, Sprouty 1 shRNA Plasmid (h): sc-41035-SH and Sprouty 1 shRNA (h) Lentiviral Particles: sc-41035-V.

Molecular Weight of Sprouty 1: 35 kDa.

Positive Controls: Hep G2 cell lysate: sc-2227 or AN3 CA cell lysate: sc-24662.

DATA





Sprouty 1 (H-2) Alexa Fluor[®] 680: sc-365520 AF680. Direct near-infrared western blot analysis of Sprouty 1 expression in Hep G2 (**A**) and AN3 CA (**B**) whole cell lysates. Blocked with UltraCruz[®] Blocking Reagent: sc-516214. Cruz Marker[™] Molecular Weight Standards detected with Cruz Marker[™] MW Tag-Alexa Fluor[®] 790 sc-51671. Sprouty 1 (H-2): sc-365520. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization (**A**). Immunoperoxidase staining of formalin fixed, paraffin-embedded human gall bladder tissue showing membrane staining of glandular cells (**B**).

SELECT PRODUCT CITATIONS

- He, Q., et al. 2016. Suppression of Spry1 inhibits triple-negative breast cancer malignancy by decreasing EGF/EGFR mediated mesenchymal phenotype. Sci. Rep. 6: 23216.
- He, Q., et al. 2017. Corrigendum: suppression of Spry1 inhibits triplenegative breast cancer malignancy by decreasing EGF/EGFR mediated mesenchymal phenotype. Sci. Rep. 7: 46791.
- Rosso, V., et al. 2019. Reduced expression of Sprouty 1 contributes to the aberrant proliferation and impaired apoptosis of acute myeloid leukemia cells. J. Clin. Med. 8: 972.
- 4. Tay, J.K., et al. 2022. The microdissected gene expression landscape of nasopharyngeal cancer reveals vulnerabilities in FGF and noncanonical $NF\kappa B$ signaling. Sci. Adv. 8: eabh2445.
- 5. Wang, J., et al. 2023. Circ_0099630 participates in SPRY1-mediated repression in periodontitis. Int. Dent. J. 73: 136-143.

STORAGE

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