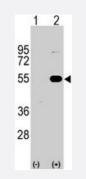
SPRED1 polyclonal antibody

Catalog # PAB3102 Size 400 uL

Applications



Western Blot (Transfected lysate)

Western blot analysis of SPRED1 (arrow) using rabbit SPRED1 polyclonal antibody (Cat # PAB3102). 293 cell lysates (2 ug/lane) either nontransfected (Lane 1) or transiently transfected with the SPRED1 gene (Lane 2) (Origene Technologies).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Formalin-fixed and paraffin-embedded human brain tissue reacted with SPRED1 polyclonal antibody (Cat # PAB3102), which was peroxidaseconjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of SPRED1.
Immunogen	A synthetic peptide (conjugated with KLH) corresponding to N-terminus of human SPRED1.
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Protein A purification



Product Information

Recommend Usage	Western Blot (1:1000) Immunohistochemistry (1:10-50) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Entrez GenelD	<u>161742</u>
Protein Accession#	<u>NP_689807;Q7Z699</u>
Gene Name	SPRED1
Gene Alias	FLJ33903, NFLS
Gene Description	sprouty-related, EVH1 domain containing 1
Omim ID	<u>609291</u> 611431
Gene Ontology	<u>Hyperlink</u>
Gene Summary	The protein encoded by this gene is a member of the Sprouty family of proteins and is phosphoryl ated by tyrosine kinase in response to several growth factors. The encoded protein can act as a h omodimer or as a heterodimer with SPRED2 to regulate activation of the MAP kinase cascade. Defects in this gene are a cause of neurofibromatosis type 1-like syndrome (NFLS). [provided by RefSeq



Product Information

Other Designations

EVH1/Sprouty domain containing protein|sprouty-related protein 1 with EVH-1 domain|suppresso r of Ras/MAPK activation

Publication Reference

Germline loss-of-function mutations in SPRED1 cause a neurofibromatosis 1-like phenotype.

Brems H, Chmara M, Sahbatou M, Denayer E, Taniguchi K, Kato R, Somers R, Messiaen L, De Schepper S, Fryns JP, Cools J, Marynen P, Thomas G, Yoshimura A, Legius E.

Nature Genetics 2007 Sep; 39(9):1120.

Application: WB-Ce, Mouse, MEFs

Violating the splicing rules: TG dinucleotides function as alternative 3' splice sites in U2-dependent introns.

Szafranski K, Schindler S, Taudien S, Hiller M, Huse K, Jahn N, Schreiber S, Backofen R, Platzer M. Genome Biology 2007 Jan; 8(8):R154.

 Spreds, inhibitors of the Ras/ERK signal transduction, are dysregulated in human hepatocellular carcinoma and linked to the malignant phenotype of tumors.

Yoshida T, Hisamoto T, Akiba J, Koga H, Nakamura K, Tokunaga Y, Hanada S, Kumemura H, Maeyama M, Harada M, Ogata H, Yano H, Kojiro M, Ueno T, Yoshimura A, Sata M.

Oncogene 2006 May; 25(45):6056.

Pathway

• Jak-STAT signaling pathway

Disease

- Cafe-au-Lait Spots
- <u>Cardiovascular Diseases</u>
- Diabetes Mellitus
- Edema
- Neurofibromatosis
- Neurofibromatosis 1



Product Information

• <u>Syndrome</u>