

TRPV4 rabbit pAb

Cat No.:ES3770

For research use only

Overview

Product Name	TRPV4 rabbit pAb	
Host species	Rabbit	
Applications	WB;ELISA	
Species Cross-Reactivity	Human;Mouse;Rat	
Recommended dilutions	Western Blot: 1/500 - 1/2000. ELISA: 1/20000. Not	
	yet tested in other applications.	
Immunogen	The antiserum was produced against synthesized	
	peptide derived from human TRPV4. AA	
	range:417-466	
Specificity	TRPV4 Polyclonal Antibody detects endogenous	
	levels of TRPV4 protein.	
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and	
	0.02% sodium azide.	
Storage	Store at -20°C. Avoid repeated freeze-thaw cycles.	
Protein Name	Transient receptor potential cation channel	
	subfamily V member 4	
Gene Name	TRPV4	
Cellular localization	Cell membrane . Apical cell membrane ; Multi-pass	
	membrane protein . Cell junction, adherens	
	junction . Cell projection, cilium . Assembly of the	
	putative homotetramer occurs primarily in the	
	endoplasmic reticulum; [Isoform 1]: Cell	
	membrane .; [Isofor	
Purification	The antibody was affinity-purified from rabbit	
	antiserum by affinity-chromatography using	
	epitope-specific immunogen.	
Clonality	Polyclonal	
Concentration	1 mg/ml	
Observed band	98kD	
Human Gene ID	59341	التر
Human Swiss-Prot Number	Q9HBA0	
Alternative Names	TRPV4; VRL2; VROAC; Transient receptor potential	
	cation channel subfamily V member 4: TrpV4:	



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Background

Osm-9-like TRP channel 4; OTRPC4; Transient receptor potential protein 12; TRP12; Vanilloid receptor-like channel 2; Vanilloid receptor-like protein 2; VRL-2; Vani

transient receptor potential cation channel subfamily V member 4(TRPV4) Homo sapiens This gene encodes a member of the OSM9-like transient receptor potential channel (OTRPC) subfamily in the transient receptor potential (TRP) superfamily of ion channels. The encoded protein is a Ca2+-permeable, nonselective cation channel that is thought to be involved in the regulation of systemic osmotic pressure. Mutations in this gene are the cause of spondylometaphyseal and metatropic dysplasia and hereditary motor and sensory neuropathy type IIC. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Apr 2010],





Western blot analysis of lysates from PC12 cells, using TRPV4 antibody.

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