

**TRPV4 rabbit pAb****Cat#: orb766645 (Manual)**

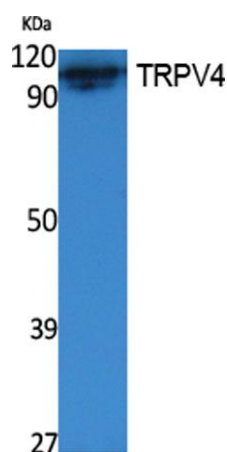
For research use only. Not intended for diagnostic use.

<b>Product Name</b>	TRPV4 rabbit pAb
<b>Host species</b>	Rabbit
<b>Applications</b>	WB;ELISA
<b>Species Cross-Reactivity</b>	Human;Mouse;Rat
<b>Recommended dilutions</b>	Western Blot: 1/500 - 1/2000. ELISA: 1/20000. Not yet tested in other applications.
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human TRPV4. AA range:417-466
<b>Specificity</b>	TRPV4 Polyclonal Antibody detects endogenous levels of TRPV4 protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide..
<b>Storage</b>	Store at -20°C. Avoid repeated freeze-thaw cycles.
<b>Protein Name</b>	Transient receptor potential cation channel subfamily V member 4
<b>Gene Name</b>	TRPV4
<b>Cellular localization</b>	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
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.

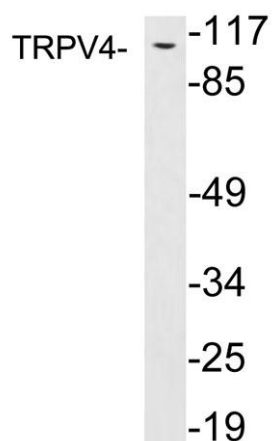
<b>Clonality</b>	Polyclonal
<b>Concentration</b>	1 mg/ml
<b>Observed band</b>	98kD
<b>Human Gene ID</b>	59341
<b>Human Swiss-Prot Number</b>	Q9HBA0
<b>Alternative Names</b>	TRPV4; VRL2; VROAC; Transient receptor potential cation channel subfamily V member 4; TrpV4; Osm-9-like TRP channel 4; OTRPC4; Transient receptor potential protein 12; TRP12; Vanilloid receptor-like channel 2; Vanilloid receptor-like protein 2; VRL-2; Vani

### Background

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 Ca<sup>2+</sup>-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 extracts from K562 cells, using TRPV4 Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000**



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