

**DVL1 Antibody (Center) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP12326c****Specification**

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**DVL1 Antibody (Center) Blocking peptide - Product Information**

Primary Accession [O14640](#)

**DVL1 Antibody (Center) Blocking peptide - Additional Information**

**Gene ID** 1855

**Other Names**

Segment polarity protein dishevelled homolog DVL-1, Dishevelled-1, DSH homolog 1, DVL1

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**DVL1 Antibody (Center) Blocking peptide - Protein Information**

**Name** DVL1

**Function**

Participates in Wnt signaling by binding to the cytoplasmic C-terminus of frizzled family members and transducing the Wnt signal to down-stream effectors. Plays a role both in canonical and non-canonical Wnt signaling. Plays a role in the signal transduction pathways mediated by multiple Wnt genes. Required for LEF1 activation upon WNT1 and WNT3A signaling. DVL1 and PAK1 form a ternary complex with MUSK which is important for MUSK-dependent regulation of AChR clustering during the formation of the neuromuscular junction (NMJ).

**Cellular Location**

Cell membrane; Peripheral membrane protein; Cytoplasmic side. Cytoplasm, cytosol. Cytoplasmic vesicle Note=Localizes at the cell membrane upon interaction with frizzled family members.

**DVL1 Antibody (Center) Blocking peptide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

**DVL1 Antibody (Center) Blocking peptide - Images****DVL1 Antibody (Center) Blocking peptide - Background**

DVL1, the human homolog of the *Drosophila* dishevelled gene (*dsh*) encodes a cytoplasmic phosphoprotein that regulates cell proliferation, acting as a transducer molecule for developmental processes, including segmentation and neuroblast specification. DVL1 is a candidate gene for neuroblastomatous transformation. The Schwartz-Jampel syndrome and Charcot-Marie-Tooth disease type 2A have been mapped to the same region as DVL1. The phenotypes of these diseases may be consistent with defects which might be expected from aberrant expression of a DVL gene during development.

**DVL1 Antibody (Center) Blocking peptide - References**

Metcalfe, C., et al. *J. Cell. Sci.* 123 (PT 9), 1588-1599 (2010) ; Hu, T., et al. *J. Biol. Chem.* 285(18):13561-13568(2010) ; Varelas, X., et al. *Dev. Cell* 18(4):579-591(2010) ; Jugessur, A., et al. *PLoS ONE* 5 (7), E11493 (2010) ; Guo, J., et al. *PLoS ONE* 4 (11), E7982 (2009) :