

# **Product datasheet**

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# ARG58502 anti-DVL1 / Dishevelled 1 antibody

Package: 100 μl Store at: -20°C

#### **Summary**

Product Description Rabbit Polyclonal antibody recognizes DVL1 / Dishevelled 1

Tested Reactivity Hu, Ms, Rat

Tested Application ICC/IF, IHC-P, WB

Host Rabbit

**Clonality** Polyclonal

Isotype IgG

Target Name DVL1 / Dishevelled 1

Species Human

Immunogen Recombinant fusion protein corresponding to aa. 510-640 of Human DVL1 / Dishevelled 1

(NP\_004412.2).

Conjugation Un-conjugated

Alternate Names DVL; DSH homolog 1; DVL1P1; DVL1L1; DRS2; Segment polarity protein dishevelled homolog DVL-1;

Dishevelled-1

## **Application Instructions**

Application table	Application	Dilution
	ICC/IF	1:50 - 1:200
	IHC-P	1:50 - 1:200
	WB	1:500 - 1:2000
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	
Positive Control	Mouse heart	
Observed Size	90 kDa	

#### **Properties**

Form	Liquid	
Purification	Affinity purified.	
Buffer	PBS (pH 7.3), 0.02% Sodium azide and 50% Glycerol.	
Preservative	0.02% Sodium azide	
Stabilizer	50% Glycerol	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use.	

#### Bioinformation

Gene Symbol DVL1

Gene Full Name dishevelled segment polarity protein 1

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. [provided by RefSeq, Jul 2008]

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). [UniProt]

Calculated Mw 75 kDa

PTM Ubiquitinated; undergoes both 'Lys-48'-linked ubiquitination, leading to its subsequent degradation by

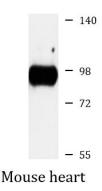
the ubiquitin-proteasome pathway, and 'Lys-63'-linked ubiquitination. The interaction with INVS is required for ubiquitination. Deubiquitinated by CYLD, which acts on 'Lys-63'-linked ubiquitin chains (By

similarity). [UniProt]

Cellular Localization Cell membrane, Peripheral membrane protein, Cytoplasmic side, Cytoplasm, cytosol, Cytoplasmic

vesicle. [UniProt]

#### **Images**



### ARG58502 anti-DVL1 / Dishevelled 1 antibody WB image

Western blot: 25  $\mu g$  of Mouse heart lysate stained with ARG58502 anti-DVL1 / Dishevelled 1 antibody at 1:1000 dilution.