

FZD7 / Frizzled 7 Antibody (C-Terminus) Rabbit Polyclonal Antibody Catalog # ALS10788

Specification

FZD7 / Frizzled 7 Antibody (C-Terminus) - Product Information

Application Primary Accession Reactivity

Host Clonality Calculated MW Dilution IHC-P, E <u>O75084</u> Human, Mouse, Rabbit, Hamster, Monkey, Bovine, Dog Rabbit Polyclonal 64kDa KDa IHC-P~~N/A E~~N/A

FZD7 / Frizzled 7 Antibody (C-Terminus) - Additional Information

Gene ID 8324

Other Names Frizzled-7, Fz-7, hFz7, FzE3, FZD7

Target/Specificity Human FZD7 / Frizzled 7. BLAST analysis of the peptide immunogen showed no homology with other human proteins, except FZD1 (61%), FZD2 (61%).

Reconstitution & Storage Long term: -70°C; Short term: +4°C

Precautions FZD7 / Frizzled 7 Antibody (C-Terminus) is for research use only and not for use in diagnostic or therapeutic procedures.

FZD7 / Frizzled 7 Antibody (C-Terminus) - Protein Information

Name FZD7

Function

Receptor for Wnt proteins. Most frizzled receptors are coupled to the beta-catenin canonical signaling pathway, which leads to the activation of disheveled proteins, inhibition of GSK-3 kinase, nuclear accumulation of beta-catenin and activation of Wnt target genes. A second signaling pathway involving PKC and calcium fluxes has been seen for some family members, but it is not yet clear if it represents a distinct pathway or if it can be integrated in the canonical pathway, as PKC seems to be required for Wnt-mediated inactivation of GSK-3 kinase. Both pathways seem to involve interactions with G-proteins. Activation by WNT8 induces expression of beta-catenin target genes (By similarity). Following ligand activation, binds to CCDC88C/DAPLE which displaces DVL1 from FZD7 and leads to inhibition of canonical Wnt signaling, activation of G-proteins by CCDC88C



and triggering of non-canonical Wnt responses (PubMed:26126266). May be involved in transduction and intercellular transmission of polarity information during tissue morphogenesis and/or in differentiated tissues.

Cellular Location

Cell membrane; Multi-pass membrane protein. Endosome membrane; Multi-pass membrane protein. Note=Associated to the plasma membrane in the presence of FZD7 and phosphatidylinositol 4,5-bisphosphate (PIP2). Localized in recycling endosomes in other conditions

Tissue Location

High expression in adult skeletal muscle and fetal kidney, followed by fetal lung, adult heart, brain, and placenta Specifically expressed in squamous cell esophageal carcinomas

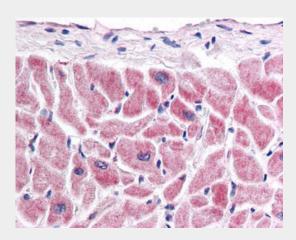
Volume 50 μl

FZD7 / Frizzled 7 Antibody (C-Terminus) - Protocols

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

- <u>Western Blot</u>
- <u>Blocking Peptides</u>
- <u>Dot Blot</u>
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

FZD7 / Frizzled 7 Antibody (C-Terminus) - Images



Anti-FZD7 / Frizzled 7 antibody ALS10788 IHC of human heart. FZD7 / Frizzled 7 Antibody (C-Terminus) - Background

Receptor for Wnt proteins. Most of frizzled receptors are coupled to the beta-catenin canonical signaling pathway, which leads to the activation of disheveled proteins, inhibition of GSK- 3 kinase, nuclear accumulation of beta-catenin and activation of Wnt target genes. A second signaling pathway involving PKC and calcium fluxes has been seen for some family members, but it is not yet clear if it represents a distinct pathway or if it can be integrated in the canonical pathway, as PKC



seems to be required for Wnt-mediated inactivation of GSK-3 kinase. Both pathways seem to involve interactions with G-proteins. May be involved in transduction and intercellular transmission of polarity information during tissue morphogenesis and/or in differentiated tissues.

FZD7 / Frizzled 7 Antibody (C-Terminus) - References

Tanaka S., et al. Proc. Natl. Acad. Sci. U.S.A. 95:10164-10169(1998). Hillier L.W., et al. Nature 434:724-731(2005). Mural R.J., et al. Submitted (SEP-2005) to the EMBL/GenBank/DDBJ databases. Sagara N., et al. Biochem. Biophys. Res. Commun. 252:117-122(1998). Kwon H.S., et al. Mol. Cell. Biol. 29:2139-2154(2009).