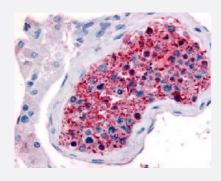


FZD9 polyclonal antibody

Catalog # PAB16383 Size 50 ug

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemical (Formalin/PFA-fixed paraffin-embedded sections) staining in human testis with FZD9 polyclonal antibody (Cat # PAB16383).

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of FZD9.
Immunogen	A synthetic peptide (conjugated with KLH) corresponding to human FZD9.
Host	Rabbit
Reactivity	Bovine, Dog, Human, Mouse, Rabbit, Rat
Specificity	N-terminal extracellular domain of human.
Form	Liquid
Purification	Immunoaffinity purification
Recommend Usage	Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (10 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -80°C. Aliquot to avoid repeated freezing and thawing.

😵 Abnova

Product Information

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical (Formalin/PFA-fixed paraffin-embedded sections) staining in human testis with FZD9 polyclonal antibody (Cat # PAB16383).

Gene Info — FZD9

Protein Accession# Q00144 Gene Name FZD9 Gene Alias CD349, FZD3 Gene Description frizzled homolog 9 (Drosophila) Omim ID 601766 Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq		
Gene Name FZD9 Gene Alias CD349, FZD3 Gene Description frizzled homolog 9 (Drosophila) Omim ID 601766 Gene Ontology Hyperlink Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq	Entrez GenelD	<u>8326</u>
Gene Alias CD349, FZD3 Gene Description frizzled homolog 9 (Drosophila) Omim ID 601766 Gene Ontology Hyperlink Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq	Protein Accession#	<u>000144</u>
Gene Description frizzled homolog 9 (Drosophila) Omim ID 601766 Gene Ontology Hyperlink Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq	Gene Name	FZD9
Omim ID 601766 Gene Ontology Hyperlink Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq	Gene Alias	CD349, FZD3
Gene Ontology Hyperlink Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal muscle, and kidney. [provided by RefSeq	Gene Description	frizzled homolog 9 (Drosophila)
Gene Summary Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal muscle, and kidney. [provided by RefSeq	Omim ID	<u>601766</u>
s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq	Gene Ontology	<u>Hyperlink</u>
Other Designations frizzled 9	Gene Summary	Members of the 'frizzled' gene family encode 7-transmembrane domain proteins that are receptor s for Wnt signaling proteins. The FZD9 gene is located within the Williams syndrome common del etion region of chromosome 7, and heterozygous deletion of the FZD9 gene may contribute to the Williams syndrome phenotype. FZD9 is expressed predominantly in brain, testis, eye, skeletal mu scle, and kidney. [provided by RefSeq
	Other Designations	frizzled 9

Pathway

- Basal cell carcinoma
- Colorectal cancer
- Melanogenesis
- Pathways in cancer
- Wnt signaling pathway