

Wnt-1 Polyclonal Antibody

Catalog No: YT4907

Reactivity: Human; Mouse

Applications: WB;IHC;IF;ELISA

Target: Wnt-1

Fields: >>mTOR signaling pathway;>>Wnt signaling pathway;>>Hippo signaling

pathway;>>Signaling pathways regulating pluripotency of stem

cells;>>Melanogenesis;>>Cushing syndrome;>>Alzheimer disease;>>Pathways

of neurodegeneration - multiple diseases;>>Human papillomavirus infection;>>Pathways in cancer;>>Proteoglycans in cancer;>>Chemical carcinogenesis - receptor activation;>>Basal cell carcinoma;>>Breast

cancer;>>Hepatocellular carcinoma;>>Gastric cancer

Gene Name: WNT1

Protein Name: Proto-oncogene Wnt-1

P04628

P04426

Human Gene Id: 7471

Human Swiss Prot

No:

Mouse Gene ld: 22408

Mouse Swiss Prot

No:

Immunogen: The antiserum was produced against synthesized peptide derived from human

WNT1. AA range:301-350

Specificity: Wnt-1 Polyclonal Antibody detects endogenous levels of Wnt-1 protein.

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, lgG

Dilution: WB 1:500 - 1:2000. IHC 1:100 - 1:300. IF 1:200 - 1:1000. ELISA: 1:10000. Not

yet tested in other applications.



Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 45kD

Cell Pathway: WNT;WNT-T CELLHedgehog;Melanogenesis;Pathways in cancer;Basal cell

carcinoma;

Background: The WNT gene family consists of structurally related genes which encode

secreted signaling proteins. These proteins have been implicated in oncogenesis and in several developmental processes, including regulation of cell fate and patterning during embryogenesis. This gene is a member of the WNT gene family. It is very conserved in evolution, and the protein encoded by this gene is known to be 98% identical to the mouse Wnt1 protein at the amino acid level. The studies in mouse indicate that the Wnt1 protein functions in the induction of the mesencephalon and cerebellum. This gene was originally considered as a candidate gene for Joubert syndrome, an autosomal recessive disorder with cerebellar hypoplasia as a leading feature. However, further studies suggested

This gene is clustered with another family member, WNT10B, in

Function: function:Ligand for members of the frizzled family of seven transmembrane

receptors. Probable developmental protein. May be a signaling molecule

important in CNS development. Is likely to signal over only few cell

diameters., similarity: Belongs to the Wnt family., subunit: Interacts with PORCN.

that the gene mutations might not have a significant role in Joubert syndrome.

Interacts with RSPO1, RSPO2 and RSPO3.,

Subcellular

Location:

Secreted, extracellular space, extracellular matrix. Secreted.

Expression: Testis,

Tag: orthogonal,hot

Sort:

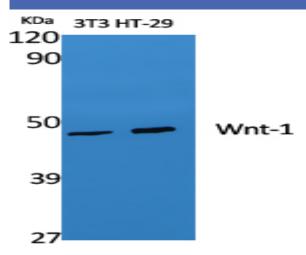
No4:

Host: Rabbit

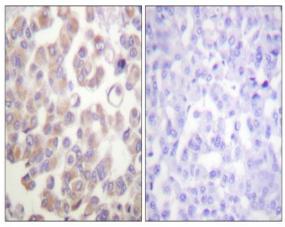
Modifications: Unmodified



Products Images



Western Blot analysis of various cells using Wnt-1 Polyclonal Antibody diluted at 1:1000. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Immunohistochemistry analysis of paraffin-embedded human breast carcinoma tissue, using WNT1 Antibody. The picture on the right is blocked with the synthesized peptide.