



# Wnt-1 rabbit pAb

Cat No.:ES3706

For research use only

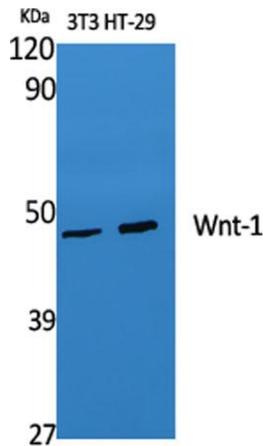
## Overview

<b>Product Name</b>	Wnt-1 rabbit pAb
<b>Host species</b>	Rabbit
<b>Applications</b>	WB;IHC;IF;ELISA
<b>Species Cross-Reactivity</b>	Human;Mouse
<b>Recommended dilutions</b>	Western Blot: 1/500 - 1/2000. Immunohistochemistry: 1/100 - 1/300. Immunofluorescence: 1/200 - 1/1000. ELISA: 1/10000. Not yet tested in other applications.
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human WNT1. AA range:301-350
<b>Specificity</b>	Wnt-1 Polyclonal Antibody detects endogenous levels of Wnt-1 protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Storage</b>	Store at -20°C. Avoid repeated freeze-thaw cycles.
<b>Protein Name</b>	Proto-oncogene Wnt-1
<b>Gene Name</b>	WNT1
<b>Cellular localization</b>	Secreted, extracellular space, extracellular matrix . Secreted .
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Clonality</b>	Polyclonal
<b>Concentration</b>	1 mg/ml
<b>Observed band</b>	45kD
<b>Human Gene ID</b>	7471
<b>Human Swiss-Prot Number</b>	P04628
<b>Alternative Names</b>	WNT1; INT1; Proto-oncogene Wnt-1; Proto-oncogene Int-1 homolog
<b>Background</b>	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 that the gene mutations might not have a significant role in Joubert syndrome. This gene is clustered with another family member, WNT10B, in



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



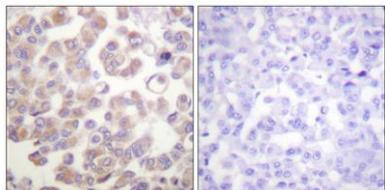
Western Blot analysis of HT29 cells using Wnt-1 Polyclonal Antibody diluted at 1:1000. Secondary antibody(catalog#:RS0002) was diluted at 1:20000





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



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