



# PJVK rabbit pAb

<b>Catalog No</b>	YP-Ab-07909
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human; Mouse
<b>Applications</b>	WB
<b>Gene Name</b>	PJVK DFNB59
<b>Protein Name</b>	PJVK
<b>Immunogen</b>	Synthesized peptide derived from human PJVK AA range: 69-119
<b>Specificity</b>	This antibody detects endogenous levels of PJVK at Human/Mouse
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.23% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
<b>Dilution</b>	WB 1:500-2000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	Pejvakin (Autosomal recessive deafness type 59 protein)
<b>Observed Band</b>	38kD
<b>Cell Pathway</b>	Peroxisome membrane . Cell projection, cilium . Associates with the peroxisomal membrane; it is unclear whether it is embedded or just associated with the peroxisomal membrane. Localizes to ciliary rootlet. .
<b>Tissue Specificity</b>	
<b>Function</b>	disease:Defects in PJVK are the cause of non-syndromic sensorineural deafness autosomal recessive type 59 (DFNB59) [MIM:610220]. DFNB59 is a form of sensorineural hearing impairment with absent or severely abnormal auditory brainstem response but normal otoacoustic emissions (auditory neuropathy or auditory dys-synchrony). Auditory neuropathies result from a lesion in the area including the inner hair cells, connections between the inner hair cells and the cochlear branch of the auditory nerve, the auditory nerve itself and auditory pathways of the brainstem.,function:Essential in the activity of auditory pathway neurons.,miscellaneous:'Pejvakin' means 'echo' in Persian.,similarity:Belongs to the gasdermin family.,
<b>Background</b>	The protein encoded by this gene is a member of the gasdermin family, a family which is found only in vertebrates. The encoded protein is required for the proper function of auditory pathway neurons. Defects in this gene are a cause of non-syndromic sensorineural deafness autosomal recessive type 59 (DFNB59).



[provided by RefSeq, Dec 2008],

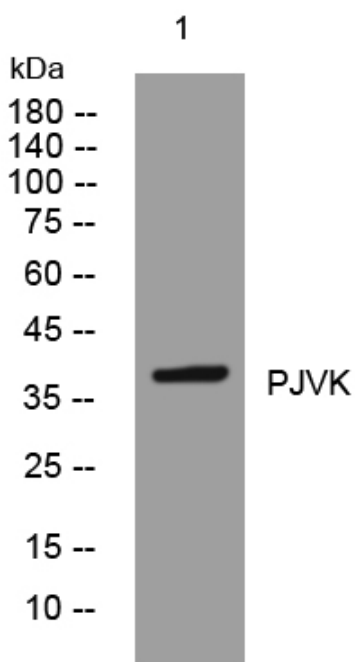
**matters needing attention**

Avoid repeated freezing and thawing!

**Usage suggestions**

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

**Products Images**



Western blot analysis of lysates from HeLa cells, primary antibody was diluted at 1:1000, 4° over night