

GNB1L rabbit pAb

Cat No.:ES11904

For research use only

Overview

Product Name	GNB1L rabbit pAb	
Host species	Rabbit	
Applications	WB;ELISA	
Species Cross-Reactivity	Human;Mouse	
Recommended dilutions	WB 1:500-2000 ELISA 1:5000-20000	
Immunogen	Synthesized peptide derived from part region of	
	human protein	
Specificity	GNB1L Polyclonal Antibody detects endogenous	
	levels of protein.	
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and	
	0.02% sodium azide.	
Storage	Store at -20°C. Avoid repeated freeze-thaw cycles.	
Protein Name	Guanine nucleotide-binding protein subunit	
	beta-like protein 1 (G protein subunit beta-like	
	protein 1) (DGCRK3) (WD repeat-containing protein	
	14) (WD40 repeat-containing protein deleted in	
	VCFS) (WDVC	
Gene Name	GNB1L GY2 KIAA1645 WDR14 FKSG1	
Cellular localization	cytoplasm,cytoplasmic side of plasma membrane,	
Purification	The antibody was affinity-purified from rabbit	
	antiserum by affinity-chromatography using	
	epitope-specific immunogen.	
Clonality	Polyclonal	
Concentration	1 mg/ml	
Observed band	35kD	
Human Gene ID	54584	
Human Swiss-Prot Number	Q9BYB4	
Alternative Names		
Background	This gene encodes a G-protein beta-subunit-like	111.
	polypeptide which is a member of the WD repeat	
	protein family. WD repeats are minimally conserved	
	regions of approximately 40 amino acids typically	
	bracketed by gly-his and trp-asp (GH-WD), which	



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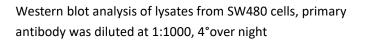
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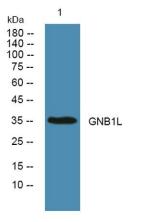
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may facilitate formation of heterotrimeric or multiprotein complexes. Members of this family are involved in a variety of cellular processes, including cell cycle progression, signal transduction, apoptosis, and gene regulation. This protein contains 6 WD repeats and is highly expressed in the heart. The gene maps to the region on chromosome 22q11, which is deleted in DiGeorge syndrome, trisomic in derivative 22 syndrome and tetrasomic in cat-eye syndrome. Therefore, this gene may contribute to the etiology of those disorders. Transcripts from this gene share exons with some transcripts from the C22orf29 gene. [provided by RefSeq, Jul 2008],







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