

WFS1 rabbit pAb

Cat No.:ES11949

For research use only

Overview

Product Name	WFS1 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	WFS1 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 $^\circ\!\mathrm{C}$. Avoid repeated freeze-thaw cycles.
Protein Name	Wolframin
Gene Name	WFS1
Cellular localization	Endoplasmic reticulum membrane ; Multi-pass
	membrane protein . Cytoplasmic vesicle, secretory
	vesicle . Co-localizes with ATP6V1A in the secretory
	granules in neuroblastoma cell lines
Purification	The antibody was affinity-purified from rabbit
	antiserum by affinity-chromatography using
	epitope-specific immunogen.
Clonality	Polyclonal
Concentration	1 mg/ml
Observed band	97kD
Human Gene ID	7466
Human Swiss-Prot Number	076024
Alternative Names	
Background	This gene encodes a transmembrane protein, which
	is located primarily in the endoplasmic reticulum
	and ubiquitously expressed with highest levels in
	brain, pancreas, heart, and insulinoma beta-cell
	lines. Mutations in this gene are associated with
	Wolfram syndrome, also called DIDMOAD (Diabetes



+86-27-59760950

ELKbio@ELKbiotech.com

www.elkbiotech.com

23-2, No.388 Gaoxin 2nd Road, Wuhan East Lake Hi-tech Development Zone, Hubei , P.R.C



Insipidus, Diabetes Mellitus, Optic Atrophy, and Deafness), an autosomal recessive disorder. The disease affects the brain and central nervous system. Mutations in this gene can also cause autosomal dominant deafness 6 (DFNA6), also known as DFNA14 or DFNA38. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Mar 2009],



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