



PNPase Monoclonal Antibody

Catalog No	YP-mAb-00714
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	PNPT1
Immunogen	The antiserum was produced against synthesized peptide derived from human PNPT1. AA range:570-619
Specificity	PNPase Monoclonal Antibody detects endogenous levels of PNPase protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	3'-5' RNA exonuclease OLD35;EC:2.7.7.8;old 35;OLD35;PNPase 1;PNPase; PNPT1;Polyribonucleotide nucleotidyltransferase 1 mitochondrial; Polyribonucleotide nucleotidyltransferase 1;mitochondrial;PNPase old-35; Polynucleotide phosphorylase 1;Polynucleotide phosphorylase-like protein
Observed Band	85kD
Calculated Molecular Weight	85kD
Cell Pathway	Cytoplasm . Mitochondrion matrix . Mitochondrion intermembrane space ; Peripheral membrane protein .
Tissue Specificity	Cervix,Epithelium,Melanoma,Skin,Teratocarcinoma,Urinary bla
Function	catalytic activity:RNA(n+1) + phosphate = RNA(n) + a nucleoside diphosphate., function:Involved in mRNA degradation. Hydrolyzes single-stranded polyribonucleotides processively in the 3'- to 5'-direction.,induction:Up-regulated in cells upon senescence and terminal differentiation. Up-regulated after treatment with interferon beta (IFN-beta).,PTM:Phosphorylated upon DNA damage, probably by ATM or ATR.,similarity:Belongs to the polyribonucleotide nucleotidyltransferase family.,similarity:Contains 1 KH domain., similarity:Contains 1 S1 motif domain.,subunit:Homotrimer (Potential). Interacts



with TCL1A; the interaction has no effect on PNPT1 exonuclease activity.,

Background

The protein encoded by this gene belongs to the evolutionary conserved polynucleotide phosphorylase family comprised of phosphate dependent 3'-to-5' exonucleases implicated in RNA processing and degradation. This enzyme is predominantly localized in the mitochondrial intermembrane space and is involved in import of RNA to mitochondria. Mutations in this gene have been associated with combined oxidative phosphorylation deficiency-13 and autosomal recessive nonsyndromic deafness-70. Related pseudogenes are found on chromosomes 3 and 7. [provided by RefSeq, Dec 2012],

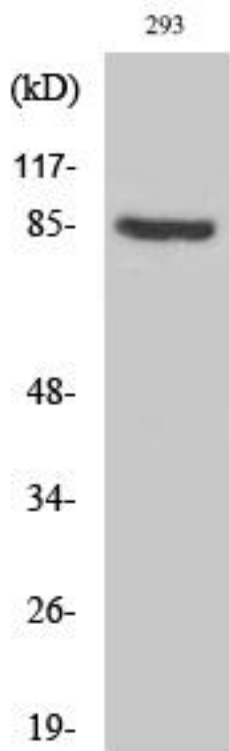
matters needing attention

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 various cells using PNPase Monoclonal Antibody