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MRPL32 Protein (Myc-DYKDDDDK Tag)



Image



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Quantity:	20 μg
Target:	MRPL32
Origin:	Human
Source:	HEK-293 Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This MRPL32 protein is labelled with Myc-DYKDDDDK Tag.
Application:	Antibody Production (AbP), Standard (STD)
Product Details	
Characteristics:	 Recombinant human MRPL32 protein expressed in HEK293 cells. Produced with end-sequenced ORF clone
Purity:	> 80 % as determined by SDS-PAGE and Coomassie blue staining
Target Details	
Target:	MRPL32
Alternative Name:	Mrpl32 (MRPL32 Products)
Background:	Mammalian mitochondrial ribosomal proteins are encoded by nuclear genes and help in protein synthesis within the mitochondrion. Mitochondrial ribosomes (mitoribosomes) consist of a small 28S subunit and a large 39S subunit. They have an estimated 75 % protein to rRNA composition compared to prokaryotic ribosomes, where this ratio is reversed. Another difference between mammalian mitoribosomes and prokaryotic ribosomes is that the latter

Target Details

>	Κp.
r	ibosomal protein family. A pseudogene corresponding to this gene is found on chromosome
t	by sequence homology. This gene encodes a 39S subunit protein that belongs to the L32
Ç	greatly in sequence, and sometimes in biochemical properties, which prevents easy recognition
C	contain a 5S rRNA. Among different species, the proteins comprising the mitoribosome differ

Molecular Weight:

21.2 kDa

NCBI Accession:

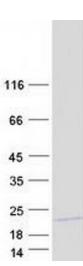
NP_114109

Application Details

Application Notes:	Recombinant human proteins can be used for:
	Native antigens for optimized antibody production
	Positive controls in ELISA and other antibody assays
Comment:	The tag is located at the C-terminal.
Restrictions:	For Research Use only

Handling

Concentration:	50 μg/mL	
Buffer:	25 mM Tris.HCl, pH 7.3, 100 mM glycine, 10 % glycerol.	
Storage:	-80 °C	
Storage Comment: Store at -80°C. Thaw on ice, aliquot to individual single-use tubes, and then re-fi immediately. Only 2-3 freeze thaw cycles are recommended.		



Western Blotting

Image 1. Validation with Western Blot