

MRP-L32 Polyclonal Antibody

Description

Product type	Primary Antibody
Code	BT-AP05582
Host	Rabbit
Isotype	IgG
Size	20ul, 50ul, 100ul
Immunogen	The antiserum was produced against synthesized peptide derived from human MRPL32. AA range:101-150
Mol wt	21405
Species reactivity	Human
Clonality	Polyclonal
Recommended application	WB, IHC-p, IF, ELISA
Concentration	1 mg/ml
Full name	MRP-L32 Antibody
Synonyms	MRPL32; HSPC283; 39S ribosomal protein L32; mitochondrial; L32mt; MRP-L32

This product is for research use only, not for use in human, therapeutic or diagnostic procedure.

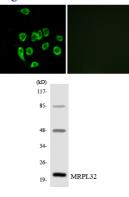
Background

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 contain a 5S rRNA. Among different species, the proteins comprising the mitoribosome differ greatly in sequence, and sometimes in biochemical properties, which prevents easy recognition by sequence homology. MRPL32 (mitochondrial ribosomal protein L32) encodes a 39S subunit protein that belongs to the L32 ribosomal protein family. A pseudogene corresponding to MRPL32 is found on chromosome Xp.

Recommended Dilution

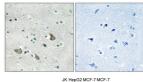
WB: 1: 500 - 1: 2000 IHC: 1: 100 - 1: 300 IF: 1: 200 - 1: 1000 ELISA: 1: 40000 Not yet tested in other applications.

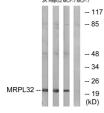
Images



Immunofluorescence analysis of A549 cells, using MRPL32 Antibody. The picture on the right is blocked with the synthesized peptide.

Western blot analysis of the lysates from HeLa cells using MRPL32 antibody.





Storage -20°C for one year

Immunohistochemical analysis of paraffin-embedded Human brain. Antibody was diluted at 1:100(4° overnight). High-pressure and temperature Tris-EDTA,pH8.0 was used for antigen retrieval. Negetive contrl (right) obtaned from antibody was pre-absorbed by immunogen peptide.

Western blot analysis of lysates from MCF-7, HepG2, and Jurkat cells, using MRPL32 Antibody. The lane on the right is blocked with the synthesized peptide.

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