

METL8 rabbit pAb

Catalog No: YT8123

Reactivity: Human; Mouse

Applications: IHC;WB

Target: METTL8

Gene Name: METTL8

Protein Name: Methyltransferase-like protein 8 (EC 2.1.1.-)

Q9H825

A2AUU0

Human Swiss Prot

No:

Mouse Gene ld: 228019

Mouse Swiss Prot

No:

Immunogen: Synthesized peptide derived from human C-ternal METL8

Specificity: This antibody detects endogenous levels of METL8 at Human, Mouse

Formulation : Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:500-2000 IHC 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 32kD

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Function:

Mitochondrial S-adenosyl-L-methionine-dependent methyltransferase that mediates N(3)-methylcytidine modification of residue 32 of the tRNA anticodon loop of mitochondrial tRNA(Ser)(UCN) and tRNA(Thr) . N(3)-methylcytidine methylation modification regulates mitochondrial translation efficiency and is required for activity of the respiratory chain . N(3)-methylcytidine methylation of mitochondrial tRNA(Ser)(UCN) requires the formation of N(6)-dimethylallyladenosine(37) (i6A37) by TRIT1 as prerequisite . May also mediate N(3)-methylcytidine modification of mRNAs . The existence of N(3)-methylcytidine modification on mRNAs is however unclear, and additional evidences are required to confirm the role of the N(3)-methylcytidine-specific mRNA methyltransferase activity of METTL8 in vivo .

Subcellular Location:

Mitochondrion . Mitochondrial protein: the cytoplasmic or nuclear localization observed by some groups is either the result of an incorrect localization caused by N-terminal tagging that interferes with mitochondrial targeting, or splice isoforms that lack the N-terminal mitochondrial transit sequence.

Sort		999
JUIL	-	333

No4:

Host: Rabbit

Modifications: Unmodified

Products Images

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