

MT2A Rabbit pAb

MT2A Rabbit pAb **Catalog # AP93951**

Specification

MT2A Rabbit pAb - Product Information

Application **Primary Accession**

Reactivity Host Clonality Calculated MW Physical State Immunogen

Epitope Specificity

Isotype **Purity**

affinity purified by Protein A

IHC-P, IHC-F, IF

063871 Mouse **Rabbit Polyclonal** 6 KDa Liquid

KLH conjugated synthetic peptide derived

from mouse MT2A

11-61/61 laG

Buffer

0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. **SIMILARITY**

Belongs to the metallothionein superfamily. Type 1 family.

Important Note

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

Background Descriptions

Metallothioneins (MTs) are a family of low molecular weight, heavy metal-binding proteins characterized by a high cysteine content and lack of aromatic amino acids. MTs bind 7 to 12 heavy metal atoms per molecule of protein. They are ubiquitous in the animal and plant kingdoms and are also found in prokaryotes. In mammals, the cysteine residues are absolutely conserved and serve to coordinate heavy metal atoms such as zinc, cadmium, and copper via mercaptide linkages.

MT2A Rabbit pAb - Additional Information

Gene ID 17749

Other Names

DNA-directed RNA polymerases I, II, and III subunit RPABC4, RNA polymerases I, II, and III subunit ABC4, DNA-directed RNA polymerase II subunit K, Metallothionein-I gene transcription activator, RPB10alpha, Polr2k, Mt1a

Dilution

IHC-P~~N/A<br \> <span class</pre> ="dilution IHC-F">IHC-F~~N/A
class ="dilution IF">IF~~1:50~200

Storage



Store at -20 $^{\circ}$ C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 $^{\circ}$ C.

MT2A Rabbit pAb - Protein Information

Name Polr2k

Synonyms Mt1a

Function

DNA-dependent RNA polymerase catalyzes the transcription of DNA into RNA using the four ribonucleoside triphosphates as substrates. Common component of RNA polymerases I, II and III which synthesize ribosomal RNA precursors, mRNA precursors and many functional non-coding RNAs, and a small RNAs, such as 5S rRNA and tRNAs, respectively.

Cellular Location

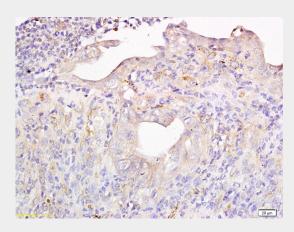
Nucleus {ECO:0000250|UniProtKB:P53803}. Nucleus, nucleolus {ECO:0000250|UniProtKB:P53803}

MT2A Rabbit pAb - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

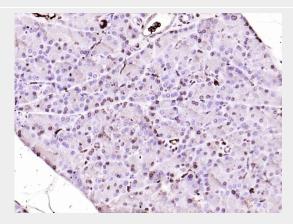
MT2A Rabbit pAb - Images



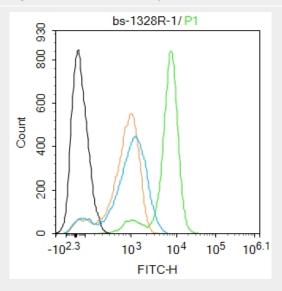
Tissue/cell: mouse colon carcinoma; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-Metallothionein Polyclonal Antibody, Unconjugated(AP93951) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010)



staining



Paraformaldehyde-fixed, paraffin embedded (mouse pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Metallothionein) Polyclonal Antibody, Unconjugated (AP93951) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control:U937. Primary Antibody (green line): Rabbit Anti-Metallothionein antibody (AP93951) Dilution: 1ug/Test; Secondary Antibody: Goat anti-rabbit IgG-FITC Dilution: 0.5ug/Test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

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