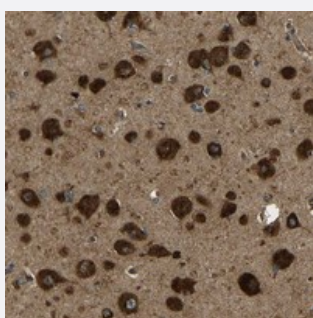


ATP7A polyclonal antibody

Catalog # PAB20632 Size 100 uL

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human hippocampus with ATP7A polyclonal antibody (Cat # PAB20632) shows strong cytoplasmic positivity in neuronal cells and glial cells at 1:10-1:20 dilution.

Specification

Product Description	Rabbit polyclonal antibody raised against recombinant ATP7A.
Immunogen	Recombinant protein corresponding to amino acids of human ATP7A.
Sequence	TETLGTCIDFQVVPGCGISCKVTNIEGLLHKNNWNIEDNNIKNASLVQIDASNEQSSTSSSMIIDAQIS NALNAQQYKVLIGNREWMIRNGLVINNDVNDFMTEHERKGRTAVLVAVDDEL C
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Antigen affinity purification
Isotype	IgG
Recommend Usage	Immunohistochemistry (1:10-1:20) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide)

Storage Instruction

Store at 4°C. For long term storage store at -20°C.
Aliquot to avoid repeated freezing and thawing.

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human hippocampus with ATP7A polyclonal antibody (Cat # PAB20632) shows strong cytoplasmic positivity in neuronal cells and glial cells at 1:10-1:20 dilution.

Gene Info — ATP7A

Entrez GeneID[538](#)**Protein Accession#**[Q04656](#)**Gene Name**

ATP7A

Gene Alias

FLJ17790, MK, MNK

Gene DescriptionATPase, Cu⁺⁺ transporting, alpha polypeptide**Omim ID**[300011](#) [304150](#) [309400](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

This gene encodes a transmembrane protein that functions in copper transport across membranes. The protein localizes to the trans-Golgi network, where it is predicted to supply copper to copper-dependent enzymes in the secretory pathway. The protein relocalizes to the plasma membrane under conditions of elevated extracellular copper and functions in the efflux of copper from cells. Mutations in this gene result in Menkes disease, X-linked cutis laxa, and occipital horn syndrome. [provided by RefSeq]

Other Designations

Cu⁺⁺-transporting P-type ATPase|Menkes disease-associated protein|Menkes syndrome|OTTHUMP00000023593|OTTHUMP00000062077|copper pump 1|copper-transporting ATPase 1