10320 Camino Santa Fe, Suite G San Diego, CA 92121 Tel: 858.875.1900 Fax: 858.875.1999



# Anti-ATP7A Antibody

Catalog # AP54130

# **Product Information**

Application WB
Primary Accession 004656

**Reactivity** Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 163373

## **Additional Information**

Gene ID 538

Other Names MC1; MNK; Copper-transporting ATPase 1; Copper pump 1; Menkes

disease-associated protein

**Target/Specificity** KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human ATP7A. The exact sequence is proprietary.

**Dilution** WB~~1/500 - 1/1000

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

## **Protein Information**

Name ATP7A {ECO:0000303 | PubMed:28389643, ECO:0000312 | HGNC:HGNC:869}

**Function** ATP-driven copper (Cu(+)) ion pump that plays an important role in

intracellular copper ion homeostasis (PubMed:10419525, PubMed:11092760, PubMed:28389643). Within a catalytic cycle, acquires Cu(+) ion from donor protein on the cytoplasmic side of the membrane and delivers it to acceptor protein on the lumenal side. The transfer of Cu(+) ion across the membrane is coupled to ATP hydrolysis and is associated with a transient phosphorylation that shifts the pump conformation from inward-facing to outward-facing state

(PubMed: 10419525, PubMed: 19453293, PubMed: 19917612,

PubMed:<u>28389643</u>, PubMed:<u>31283225</u>). Under physiological conditions, at low cytosolic copper concentration, it is localized at the trans-Golgi network (TGN) where it transfers Cu(+) ions to cuproenzymes of the secretory pathway (PubMed:<u>11092760</u>, PubMed:<u>28389643</u>). Upon elevated cytosolic copper concentrations, it relocalizes to the plasma membrane where it is responsible for the export of excess Cu(+) ions (PubMed:<u>10419525</u>, PubMed:<u>28389643</u>). May play a dual role in neuron function and survival by regulating cooper

efflux and neuronal transmission at the synapse as well as by supplying Cu(+) ions to enzymes such as PAM, TYR and SOD3 (By similarity) (PubMed:28389643). In the melanosomes of pigmented cells, provides copper cofactor to TYR to form an active TYR holoenzyme for melanin biosynthesis (By similarity).

#### **Cellular Location**

Golgi apparatus, trans-Golgi network membrane; Multi-pass membrane protein. Cell membrane; Multi-pass membrane protein Melanosome membrane {ECO:0000250 | UniProtKB:Q64430}; Multi-pass membrane protein. Early endosome membrane {ECO:0000250 | UniProtKB:Q64430}; Multi-pass membrane protein. Cell projection, axon {ECO:0000250 | UniProtKB:P70705} Cell projection, dendrite {ECO:0000250 | UniProtKB:P70705}. Postsynaptic density {ECO:0000250 | UniProtKB:P70705}. Note=Cycles constitutively between the TGN and the plasma membrane (PubMed:9147644). Predominantly found in the TGN and relocalized to the plasma membrane in response to elevated copper levels. Targeting into melanosomes is regulated by BLOC-1 complex (By similarity). In response to glutamate, translocates to neuron processes with a minor fraction at extrasynaptic sites (By similarity). {ECO:0000250 | UniProtKB:P70705, ECO:0000250 | UniProtKB:Q64430, ECO:0000269 | PubMed:9147644} [Isoform 5]: Endoplasmic reticulum

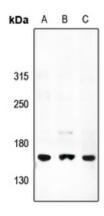
### **Tissue Location**

Widely expressed including in heart, brain, lung, muscle, kidney, pancreas, and to a lesser extent placenta (PubMed:8490646, PubMed:8490659). Expressed in fibroblasts, aortic smooth muscle cells, aortic endothelial cells and umbilical vein endothelial cells (at protein level) (PubMed:16371425)

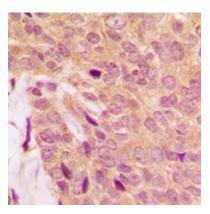
# **Background**

Rabbit polyclonal antibody to ATP7A

# **Images**



Western blot analysis of ATP7A expression in A549 (A), BV2 (B), H9C2 (C) whole cell lysates.



Immunohistochemical analysis of ATP7A staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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