

## **Product Datasheet**

# Anti-Copper ATPase 1 (Menke's Disease Protein) Antibody FL490 Conjugate



Overview

Catalog #	75-142-FL490
Conjugate	FL490 Ex: 491 nm, Em: 515 nm
Isotype	lgG2b
Clone Number	L60/4
Size	200 μL
Concentration	0.5 mg/mL
Host Species	Mouse Monoclonal
Format	Purified by Protein A chromatography
Buffer	PBS with 0.09% azide
Applications	ICC, IHC
Species Reactivity	Human, Mouse, and Rat
Immunogen	Synthetic peptide amino acids 42-61 (cytoplasmic N- terminus) of human Copper-transporting ATPase 1 (accession number Q04656)
Molecular Weight	180 kDa in rat brain membrane preparations
Cite this Antibody	Antibodies Inc Cat# 75-142-FL490, RRID: AB_2939524
Details	
Target Description	Copper-transporting ATPase 1 or ATPase Copper Transporting Alpha is encoded by the gene ATP7A and belongs to the cation transport ATPase (P-type) family. Copper-transporting ATPase 1 is a transmembrane enzyme that functions to provide copper to copper requiring proteins in the Golgi network and is also to remove excess copper across plasma membranes out of the cell. Copper-transporting ATPase 1 is found in most tissues except liver. Mutations in this gene are associated with Menkes disease, X-linked distal spinal muscular atrophy, and occipital horn syndrome.
Specificity	No cross-reactivity reported
Purification Method	Produced by in vitro bioreactor culture of hybridoma line followed by Protein A affinity chromatography and conjugation of purified mAb. Purified mAbs are >90% specific antibody.
Quality Control Tests	Each new lot of antibody is quality control tested on cells overexpressing target protein and confirmed to give the expected staining pattern.

### Storage

Aliquot and store at  $\leq$  -20°C for long term storage. For short term storage, store at 2-8°C. For maximum recovery of product, centrifuge the vial prior to removing the cap.

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