

## ATP7B

**Reactivity:**Human Mouse Rat

**Tested applications:**WB IHC IF

**Recommended Dilution:**WB 1:500 - 1:2000 IHC 1:50 - 1:200 IF 1:50 - 1:200

**Calculated MW:**157kDa

**Observed MW:**Refer to Figures

**Immunogen:**

Recombinant protein of human ATP7B

**Storage Buffer:**

Store at -20. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Synonym:**

WD; PWD; WC1; WND;

**Catalog #:**A5676

**Antibody Type:**

Polyclonal Antibody

**Species:**Rabbit

**Gene ID:**540

**Isotype:**IgG

**Swiss Prot:**P35670

**Purity:**Affinity purification

For research use only.

**Background:**

This gene is a member of the P-type cation transport ATPase family and encodes a protein with several membrane-spanning domains, an ATPase consensus sequence, a hinge domain, a phosphorylation site, and at least 2 putative copper-binding sites. This protein functions as a monomer, exporting copper out of the cells, such as the efflux of hepatic copper into the bile. Alternate transcriptional splice variants, encoding different isoforms with distinct cellular localizations, have been characterized. Mutations in this gene have been associated with Wilson disease (WD).

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