

## ATP7B-Specific Recombinant antibody

**Cat:**B36464S**Company:** HaoKebio**Uniprot ID:**P35670**Applications:** IHC:1:1000-1:4000**Organism:**Rabbit

IHC-Polymer:1:4000-1:16000

**Species reactivity:**Human Mouse Rat

IHC-TSA:1:5000-1:20000

**Molecular Weight Calculation:** 157 kDa

IF:1:50-1:200

**Observed Molecular Weight:** 150 kDa

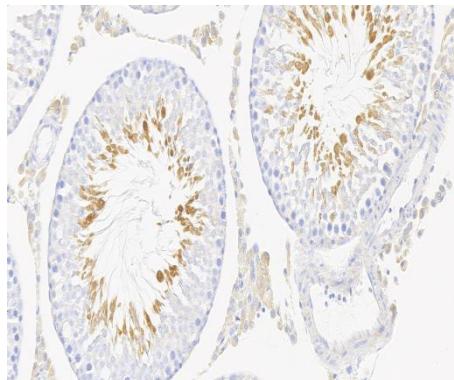
WB:1:5000-1:50000

FC:1:200-1:600

**Background:**

ATP7B, also named as PWD, WC1 and WND, belongs to the cation transport ATPase (P-type) family and Type IB subfamily. ATP7B is involved in the export of copper out of the cells, such as the efflux of hepatic copper into the bile. ATP7B catalyzes the reaction: ATP + H<sub>2</sub>O + Cu<sup>2+</sup>(In) = ADP + phosphate + Cu<sup>2+</sup>(Out). Defects in ATP7B are the cause of Wilson disease (WD). This antibody is specific to ATP7B.

Antigen retrieval: Citrate buffer (pH 9.0) , Medium high heat for 8 minutes, stop for 7 minutes, medium high heat for 8 minutes. Incubate antibody, 4°C overnight. Secondary antibody: Poly-HRP Goat Anti-Rabbit & Mouse Universal Secondary Antibody, RT, 1h.

**Images:**

Sample: Mouse testis, 4% PFA 12-24h

**Source of Reagents:**

发表[中文论文]请标注:ATP7B-Specific(B36464S)由杭州浩克生物技术有限公司提供;

发表[英文论文]请标注:ATP7B-Specific(B36464S) were kindly provided by Hangzhou Haoke Biotechnology Co., Ltd.

**Synonyms:**

ATP7B, 242827B3, ATPase, Cu<sup>++</sup> transporting, beta polypeptide, Copper-transporting ATPase 2, EC:7.2.2.8

**Immunogen:**

Recombinant protein

**Isotype:**

IgG

**Subcellular location:**

Cytoplasm

**Purity:**

Affinity purification

**Form:**

Liquid

**Storage Buffer:**

PBS with 0.02% sodium azide, 100 μg/ml BSA and 50% glycerol.

**Storage:**

Store at -20 °C for one year.

**Experimental procedure:**