

Product datasheet for TA334643

ATP7A Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: WB

Recommended Dilution: WB

Reactivity: Human

Host: Rabbit

Isotype: IgG

Clonality: Polyclonal

Immunogen: The immunogen for anti-ATP7A antibody is: synthetic peptide directed towards the middle

region of Human ATP7A. Synthetic peptide located within the following region:

MGSAAMAASSVSVVLSSLFLKLYRKPTYESYELPARSQIGQKSPSEISVH

Formulation: Liquid. Purified antibody supplied in 1x PBS buffer with 0.09% (w/v) sodium azide and 2%

sucrose.

Note that this product is shipped as lyophilized powder to China customers.

Purification: Affinity Purified

Conjugation: Unconjugated

Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 55 kDa

Gene Name: ATPase copper transporting alpha

Database Link: NP 000043

Entrez Gene 538 Human

Q04656

Background: The ATP7A gene encodes the Menkes copper-translocating P-type ATPase, a ubiquitous

protein that regulates the absorption of copper in the gastrointestinal tract. Inside cells, this protein has a dual function: it delivers copper to cuproenzymes in the Golgi compartment and effluxes excess copper. The trafficking mechanism and catalytic activity combine to facilitate

absorption and intercellular transport of copper. Menkes disease, a systemic copper

deficiency disorder, is caused by mutations in the ATP7A gene.



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



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Synonyms: DSMAX; MK; MNK; SMAX3

Note: Immunogen Sequence Homology: Dog: 100%; Pig: 100%; Rat: 100%; Human: 100%; Mouse:

100%; Sheep: 100%; Bovine: 100%; Rabbit: 100%; Guinea pig: 100%; Horse: 93%; Yeast: 93%;

Zebrafish: 92%

Protein Families: Druggable Genome, Transmembrane

Product images:



Host: Rabbit; Target Name: ATP7A; Sample Tissue: Fetal Liver lysates; Antibody Dilution: 1.0 ug/ml