



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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## Datasheet

### ATP7A (Human) Recombinant Protein (Q01)

**Catalog Number:** H00000538-Q01

**Regulation Status:** For research use only (RUO)

**Product Description:** Human ATP7A partial ORF ( NP\_000043, 1406 a.a. - 1500 a.a.) recombinant protein with GST-tag at N-terminal.

**Sequence:**

FLKLYRKPTYESYELPARSQIGQKSPSEISVHVGIDDT  
RNSPKLGLLDRIVNYSRASINSLSDKRSLNSVVTSEP  
DKHSLLVGDFREDDDTAL

**Host:** Wheat Germ (in vitro)

**Theoretical MW (kDa):** 36.19

**Applications:** AP, Array, ELISA, WB-Re  
(See our web site product page for detailed applications information)

**Protocols:** See our web site at  
<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Preparation Method:** [in vitro wheat germ expression system](#)

**Purification:** Glutathione Sepharose 4 Fast Flow

**Storage Buffer:** 50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

**Storage Instruction:** Store at -80°C. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 538

**Gene Symbol:** ATP7A

**Gene Alias:** FLJ17790, MK, MNK

**Gene Summary:** This gene encodes a transmembrane protein that functions in copper transport across membranes. The protein localizes to the trans-Golgi network, where it is predicted to supply copper to copper-dependent enzymes in the secretory pathway.

The protein relocalizes to the plasma membrane under conditions of elevated extracellular copper and functions in the efflux of copper from cells. Mutations in this gene result in Menkes disease, X-linked cutis laxa, and occipital horn syndrome. [provided by RefSeq]