



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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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

### ATP7B (Human) Recombinant Protein (Q01)

**Catalog Number:** H00000540-Q01

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

**Product Description:** Human ATP7B partial ORF ( NP\_000044, 1372 a.a. - 1465 a.a.) recombinant protein with GST-tag at N-terminal.

**Sequence:**

QLKCYKKPDLERYEAQAHGHMKPLTASQVSVHIGMD  
DRWRDSPRATPWDQVSYVSQVLSLSTSDKPSRHSA  
AADDGDKWSLLLNGRDEEQYI

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

**Theoretical MW (kDa):** 36.08

**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:** 540

**Gene Symbol:** ATP7B

**Gene Alias:** PWD, WC1, WD, WND

**Gene Summary:** 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). [provided by RefSeq]