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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

TIMM8A (Human) Recombinant Protein (P01)

Catalog Number: H00001678-P01

Regulation Status: For research use only (RUO)

Product Description: Human TIMM8A full-length ORF (AAH05236, 1 a.a. - 72 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence:

MLLNDKWVNEEIKKKIEKCLETNDNGNTTYQNLWDTA
KAVVRGKFIAISTYIKKEEKLQINNLTMNLIELEN

Host: Wheat Germ (in vitro)

Theoretical MW (kDa): 33.66

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: 1678

Gene Symbol: TIMM8A

Gene Alias: DDP, DDP1, DFN1, MGC12262, MTS

Gene Summary: This translocase is involved in the import and insertion of hydrophobic membrane proteins from the cytoplasm into the mitochondrial inner membrane. The gene is mutated in Mohr-Tranebjaerg syndrome/Deafness Dystonia Syndrome (MTS/DDS) and it is postulated that MTS/DDS is a mitochondrial

disease caused by a defective mitochondrial protein import system. Defects in this gene also cause Jensen syndrome; an X-linked disease with opticoacoustic nerve atrophy and muscle weakness. This protein, along with TIMM13, forms a 70 kDa heterohexamer. Alternative splicing results in multiple transcript variants encoding distinct isoforms]