

# Product Information

## MemDX™ Membrane Protein Human ABCD3 (ATP binding cassette subfamily D member 3) for Antibody Discovery

Cat. No.: **MP0009X**

This product is for research use only and is not intended for diagnostic use.

This product is a 51.7 kDa Human ABCD3 membrane protein expressed in *in vitro* wheat germ expression system. The protein is for research use only and is not approved for use in humans or in clinical diagnosis.

### Product Specifications

#### Host Species

Human

#### Target Protein

ABCD3

#### Protein Length

Full-length

#### Molecular Weight

51.7 kDa

#### TMD

4

#### Sequence

MAAFSKYLTARNSSLAGAAFLLLCLLHKRRRALGLHGKKSQKPPPLQNNKEGKKERAVVDKVFFSRLLQILKIMVPRTFCKETGYLV

### Product Description

#### Application

Enzyme-linked Immunoabsorbent Assay, Western Blot (Recombinant protein), Antibody Production, Protein Array

#### Expression Systems

*in vitro* wheat germ expression system

#### Tag

GST-tag at N-terminal

#### Form

Liquid

#### Purification

**Buffer**

50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer

**Storage**

Store at +4°C for up to one week or several months at -80°C

**Target****Target Protein**

ABCD3

**Full Name**

ATP binding cassette subfamily D member 3

**Introduction**

The protein encoded by this gene is a member of the superfamily of ATP-binding cassette (ABC) transporters. ABC proteins transport various molecules across extra- and intra-cellular membranes. ABC genes are divided into seven distinct subfamilies (ABC1, MDR/TAP, MRP, ALD, OABP, GCN20, White). This protein is a member of the ALD subfamily, which is involved in peroxisomal import of fatty acids and/or fatty acyl-CoAs in the organelle. All known peroxisomal ABC transporters are half transporters which require a partner half transporter molecule to form a functional homodimeric or heterodimeric transporter. This peroxisomal membrane protein likely plays an important role in peroxisome biogenesis. Mutations have been associated with some forms of Zellweger syndrome, a heterogeneous group of peroxisome assembly disorders. Alternative splicing results in multiple transcript variants encoding distinct isoforms.

**Alternative Names**

ABC43; PMP70; PXMP1; ATP-binding cassette, sub-family D, member 3; OTTHUMP00000012428; Peroxisomal membrane protein-1 (70kD), dJ824O18.1 (ATP-binding cassette, sub-family D (ALD), member 3 (PMP70, PXMP1)); peroxisomal membrane protein 1 (70kD, Zellweger syndrome)

**Gene ID**

[5825](#)

**UniProt ID**

[P28288](#)