

## Product Information

### MemDX™ Membrane Protein Human SLC25A19 (Solute carrier family 25 member 19) for Antibody Discovery

Cat. No.: **MP1183X**

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

This product is a 60.94 kDa Human SLC25A19 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

SLC25A19

##### Protein Length

Full-length

##### Molecular Weight

60.94 kDa

##### TMD

6

##### Sequence

MVGYDPKPDGRNNTKFQVAVAGSVSGLVTRALISPFDVIRFQLQHERLSRSDPSAKYHGILQASRQILQEEGPTAFWKGHVPAQ

#### 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

##### Protein Format

Liposome

##### Form

Liquid

### **Purification**

Glutathione Sepharose 4 Fast Flow

### **Buffer**

50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0

### **Storage**

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

## **Target**

### **Target Protein**

SLC25A19

### **Full Name**

Solute carrier family 25 member 19

### **Introduction**

This gene encodes a mitochondrial protein that is a member of the solute carrier family. Although this protein was initially thought to be the mitochondrial deoxynucleotide carrier involved in the uptake of deoxynucleotides into the matrix of the mitochondria, further studies have demonstrated that this protein instead functions as the mitochondrial thiamine pyrophosphate carrier, which transports thiamine pyrophosphates into mitochondria. Mutations in this gene cause microcephaly, Amish type, a metabolic disease that results in severe congenital microcephaly, severe 2-ketoglutaric aciduria, and death within the first year. Multiple alternatively spliced variants, encoding the same protein, have been identified for this gene.

### **Alternative Names**

DNC; TPC; MUP1; MCPHA; THMD3; THMD4; mitochondrial thiamine pyrophosphate carrier; mitochondrial uncoupling protein 1; solute carrier family 25 (mitochondrial deoxynucleotide carrier), member 19; solute carrier family 25 (mitochondrial thiamine pyrophosphate carrier), member 19

### **Gene ID**

[60386](#)

### **UniProt ID**

[Q9HC21](#)