



# 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

### SLC25A19 (Human) Recombinant Protein (P01)

**Catalog Number:** H00060386-P01

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

**Product Description:** Human SLC25A19 full-length ORF ( AAH01075, 1 a.a. - 320 a.a.) recombinant protein with GST-tag at N-terminal.

**Sequence:**

MVGYDPKPDGRNNTKFQVAVAGSVSGLVTRALISPF  
VIKIRFQLQHERLSRSDPSAKYHGILQASRQILQEEGPT  
AFWKGHVPAQILSIGYGAQVFLSFEMLTENVHRGSVY  
DAREFSVHFVCGGLAACMATLTVHPVDVLRTRFAAQG  
EPKVYNTLRHAVGTMYSRSEGPQVFYKGLAPTLIAIFPY  
AGLQFSCYSSLKHLYKWAIPAEGKKNENLQNLCCGSG  
AGVISKTLTYPLDLFKRLQVGGFEHARAAGQVRRY  
KGLMDCAKQVLQKEGALGFFKGLSPSLLKAALSTGFM  
FFSYEFFCNVHFCMNRTASQR

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

**Theoretical MW (kDa):** 60.94

**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:** 60386

**Gene Symbol:** SLC25A19

**Gene Alias:** DNC, MCPHA, MUP1, TPC

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