



# 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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## DTNA 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # : H00001837-T03

規格 : [ 100 uL ]

List All

### Specification

**Transfected Cell Line:** 293T

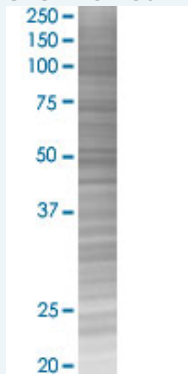
**Plasmid:** pCMV-DTNA full-length

**Host:** Human

**Theoretical MW (kDa):** 40.92

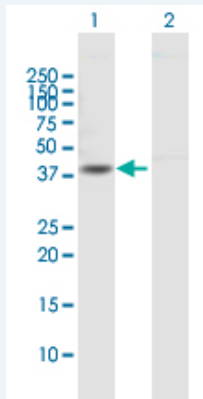
**Quality Control Testing:** Transient overexpression cell lysate was tested with Anti-DTNA antibody (H00001837-B01P) by Western Blots.

#### SDS-PAGE Gel



DTNA transfected lysate.

#### Western Blot



Lane 1: DTNA transfected lysate ( 40.92 KDa)

Lane 2: Non-transfected lysate.

**Storage Buffer:** 1X Sample Buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 300 mM 2-mercaptoethanol, 0.01% Bromophenol blue)

**Storage Instruction:** Store at -80°C. Aliquot to avoid repeated freezing and thawing.

**MSDS:**  [Download](#)

### Applications

## Western Blot

### Gene Information

**Entrez GeneID:** [1837](#)

**GeneBank  
Accession#:** [BC005300.1](#)

**Protein  
Accession#:** [AAH05300.1](#)

**Gene Name:** DTNA

**Gene Alias:** D18S892E,DRP3,DTN,FLJ96209,LVNC1

**Gene  
Description:** dystrobrevin, alpha

**Omim ID:** [601239](#), [604169](#), [606617](#)

**Gene Ontology:** [Hyperlink](#)

**Gene Summary:** The protein encoded by this gene belongs to the dystrobrevin subfamily of the dystrophin family. This protein is a component of the dystrophin-associated protein complex (DPC), which consists of dystrophin and several integral and peripheral membrane proteins, including dystroglycans, sarcoglycans, syntrophins and alpha- and beta-dystrobrevin. The DPC localizes to the sarcolemma and its disruption is associated with various forms of muscular dystrophy. Mutations in this gene are associated with left ventricular noncompaction with congenital heart defects. Multiple alternatively spliced transcript variants encoding different isoforms have been identified for this gene. [provided by RefSeq]

**Other  
Designations:** OTTHUMP00000163154,OTTHUMP00000163155,dystrobrevin alpha,dystrophin-related protein 3

### Related Disease

[Tobacco Use Disorder](#)

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