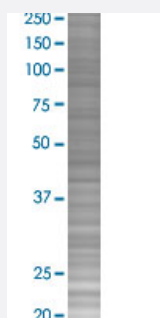


SGCE 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # H00008910-T02

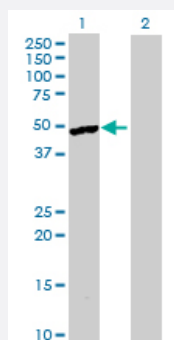
Size 100 uL

Applications



SDS-PAGE Gel

SGCE transfected lysate.



Western Blot

Lane 1: SGCE transfected lysate (49.90 KDa)

Lane 2: Non-transfected lysate.

Specification

Transfected Cell Line	293T
Plasmid	pCMV-SGCE full-length
Host	Human
Theoretical MW (kDa)	49.9
Interspecies Antigen Sequence	Mouse (88); Rat (96)

Quality Control Testing

Transient overexpression cell lysate was tested with Anti-SGCE antibody ([H00008910-B01P](#)) by Western Blots.
SDS-PAGE Gel
SGCE transfected lysate.
Western Blot
Lane 1: SGCE transfected lysate (49.90 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.

Applications

- Western Blot

Gene Info — SGCE

Entrez GeneID[8910](#)**GeneBank Accession#**[NM_003919.1](#)**Protein Accession#**[NP_003910.1](#)**Gene Name**

SGCE

Gene Alias

DYT11, ESG

Gene Description

sarcoglycan, epsilon

Omim ID[159900 604149](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

This gene encodes a member of the sarcoglycan family. Sarcoglycans are transmembrane components in the dystrophin-glycoprotein complex which help stabilize the muscle fiber membranes and link the muscle cytoskeleton to the extracellular matrix. Mutations in this gene have been associated with myoclonus-dystonia syndrome. Alternative splicing results in multiple transcript variants

Other Designations

dystonia 11, myoclonic

Disease

- [Chromosome Aberrations](#)
- [Chromosome Deletion](#)
- [Cognition](#)
- [Dystonia](#)
- [Dystonic Disorders](#)
- [Genetic Predisposition to Disease](#)
- [Myoclonus](#)
- [Obsessive-Compulsive Disorder](#)
- [Schizophrenia](#)
- [Tourette Syndrome](#)