



**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

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## CD59 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # : H00000966-T02

規格 : [ 100 uL ]

[List All](#)

### Specification

Transfected 293T

Cell Line:

Plasmid: pCMV-CD59 full-length

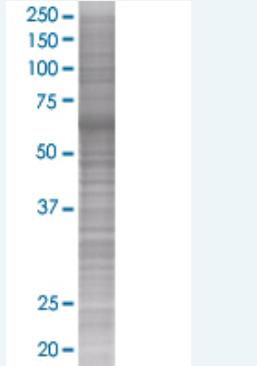
Host: Human

Theoretical MW 14.19

(kDa):

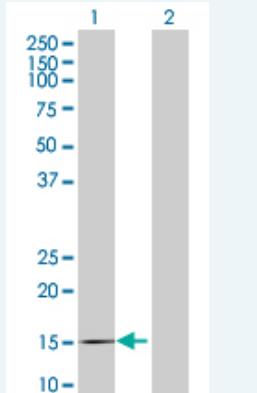
**Quality Control Testing:** Transient overexpression cell lysate was tested with Anti-CD59 antibody (H00000966-B02) by Western Blots.

#### SDS-PAGE Gel



CD59 transfected lysate.

#### Western Blot



Lane 1: CD59 transfected lysate ( 14.19 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

### Application Image

Western Blot

## Western Blot

### Gene Information

Entrez GeneID: [966](#)

GeneBank [NM\\_000611](#)

Accession#:

Protein [NP\\_000602](#)

Accession#:

Gene Name: CD59

Gene Alias: 16.3A5,1F5,EJ16,EJ30,EL32,FLJ38134,FLJ92039,G344,HRF-20,HRF20,MAC-IP,MACIF,MEM43,MGC2354,MIC11,MIN1,MIN2,MIN3,MIRL,MSK21,p18-20

Gene CD59 molecule, complement regulatory protein

Description:

Omim ID: [107271](#)

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

**Gene Summary:** This gene encodes a cell surface glycoprotein that regulates complement-mediated cell lysis, and it is involved in lymphocyte signal transduction. This protein is a potent inhibitor of the complement membrane attack complex, whereby it binds complement C8 and/or C9 during the assembly of this complex, thereby inhibiting the incorporation of multiple copies of C9 into the complex, which is necessary for osmolytic pore formation. This protein also plays a role in signal transduction pathways in the activation of T cells. Mutations in this gene cause CD59 deficiency, a disease resulting in hemolytic anemia and thrombosis, and which causes cerebral infarction. Multiple alternatively spliced transcript variants, which encode the same protein, have been identified for this gene. [provided by RefSeq]

Other 20 kDa homologous restriction factor,CD59 antigen,CD59 antigen p18-20 (antigen identified by monoclonal antibodies 16.3A5, EJ16, EJ30, EL32 and G344),CD59 glycoprotein,Ly-6-like protein,T cell-activating protein,human leukocyte antigen MIC11,lymphocytic a

### Gene Pathway

Complement and coagulation cascades [Hematopoietic cell lineage](#)

### Related Disease

[Genetic Predisposition to Disease](#) [Lymphoma](#), [Non-Hodgkin Macular Degeneration](#)