



# 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

### CD59 (Human) Recombinant Protein

**Catalog Number:** P9097

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

**Product Description:** Human CD59 (P13987, 26 a.a. - 102 a.a.) partial-length recombinant protein with His tag at N-Terminus expressed in *Escherichia coli*.

**Sequence:**

MGSSHHHHHHSSGLVPRGSHMGSLQCYNCPNPTAD  
CKTAVNCSSDFDAKLITKAGLQVYNKCWKFEHCNFND  
VTTRLRENELTYCYCKKDLNCFNEQLEN.

**Host:** *Escherichia coli*

**Theoretical MW (kDa):** 11.3

**Protocols:** See our web site at  
<http://www.abnova.com/support/protocols.asp> or product  
page for detailed protocols

**Form:** Liquid

**Preparation Method:** *Escherichia coli* expression  
system

**Purity:** > 85% by SDS-PAGE.

**Storage Buffer:** 20mM Tris-HCl buffer (pH 8.0), 0.15M  
NaCl, 1mM DTT and 10% glycerol.

**Storage Instruction:** Store at -20°C. Aliquot the product  
after reconstitution to avoid repeated freezing/thawing  
cycles.

**Entrez GeneID:** 966

**Gene Symbol:** 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 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]