CD59 (1-129) 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # H00000966-T07 Size 100 uL

Applications



Specification	
Transfected Cell Line	293T
Plasmid	pCMV-CD59 full-length
Host	Human
Theoretical MW (kDa)	14.19



Product Information

Quality Control Testing	Transient overexpression cell lysate was tested with Anti-CD59 antibody (H00000966-B05P) by We		
	stern 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% Bro mophenol blue)		
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.		

Applications

• Western Blot

Gene Info — CD59	
Entrez GenelD	<u>966</u>
GeneBank Accession#	BC001506
Protein Accession#	<u>AAH01506</u>
Gene Name	CD59
Gene Alias	16.3A5, 1F5, EJ16, EJ30, EL32, FLJ38134, FLJ92039, G344, HRF-20, HRF20, MAC-IP, MACI F, MEM43, MGC2354, MIC11, MIN1, MIN2, MIN3, MIRL, MSK21, p18-20
Gene Description	CD59 molecule, complement regulatory protein
Omim ID	<u>107271</u>
Gene Ontology	<u>Hyperlink</u>
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 thi s 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 pathw ays 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 s pliced transcript variants, which encode the same protein, have been identified for this gene. [prov ided by RefSeq



Product Information

Other Designations

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

Pathway

- Complement and coagulation cascades
- Hematopoietic cell lineage

Disease

- Genetic Predisposition to Disease
- Lymphoma
- <u>Macular Degeneration</u>