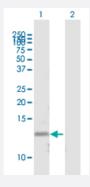


MaxPab@

# CD59 purified MaxPab mouse polyclonal antibody (B05P)

Catalog # H00000966-B05P Size 50 ug

## **Applications**



#### Western Blot (Transfected lysate)

Western Blot analysis of CD59 expression in transfected 293T cell line (<u>H00000966-T07</u>) by CD59 MaxPab polyclonal antibody.

Lane 1: CD59 transfected lysate(14.08 KDa).

Lane 2: Non-transfected lysate.

Specification	
Product Description	Mouse polyclonal antibody raised against a full-length human CD59 protein.
Immunogen	CD59 (AAH01506, 1 a.a. ~ 128 a.a) full-length human protein.
Sequence	MGIQGGSVLFGLLLVLAVFCHSGHSLQCYNCPNPTADCKTAVNCSSDFDACLITKAGLQVYNKC WKFEHCNFNDVTTRLRENELTYYCCKKDLCNFNEQLENGGTSLSEKTVLLLVTPFLAAAWSLHP
Host	Mouse
Reactivity	Human
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

### **Applications**





Western Blot (Transfected lysate)

Western Blot analysis of CD59 expression in transfected 293T cell line (<u>H00000966-T07</u>) by CD59 MaxPab polyclonal antibody.

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**Protocol Download** 

Gene Info — CD59	
Entrez GenelD	966
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 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 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
- Macular Degeneration