

UNC13D monoclonal antibody (M05), clone 2C7

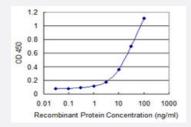
Catalog # H00201294-M05 Size 100 ug

Applications



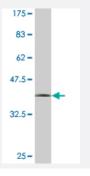
Western Blot (Cell lysate)

UNC13D monoclonal antibody (M05), clone 2C7. Western Blot analysis of UNC13D expression in Jurkat (Cat # L017V1).



Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged UNC13D is 0.3 ng/ml as a capture antibody.



Western Blot detection against Immunogen (36.41 KDa).

Specification

Product Description

Mouse monoclonal antibody raised against a partial recombinant UNC13D.



Product Information

Immunogen	UNC13D (NP_954712.1, 2 a.a. ~ 100 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
Sequence	ATLLSHPQQRPPFLRQAIKIRRRRVRDLQDPPPQMAPEIQPPSHHFSPEQRALLYEDALYTVLHRL GHPEPNHVTEASELLRYLQEAFHVEPEEHQQTL
Host	Mouse
Reactivity	Human
Isotype	lgG2a Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (36.41 KDa).
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 (Cell lysate)

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Protocol Download

Western Blot (Recombinant protein)

Protocol Download

Sandwich ELISA (Recombinant protein)

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Protocol Download

ELISA

Gene Info — UNC1		4		ч.	\sim $^{\prime}$	\sim	\neg
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Entrez GenelD	<u>201294</u>
GeneBank Accession#	NM_199242
Protein Accession#	NP_954712.1



Product Information

Gene Name	UNC13D
Gene Alias	FHL3, HLH3, HPLH3, Munc13-4
Gene Description	unc-13 homolog D (C. elegans)
Omim ID	608897 608898
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a protein that is a member of the UNC13 family, containing similar domain structure as other family members but lacking an N-terminal phorbol ester-binding C1 domain present in other Munc13 proteins. The protein appears to play a role in vesicle maturation during exocytosis and is involved in regulation of cytolytic granules secretion. Mutations in this gene are associated with familial hemophagocytic lymphohistiocytosis type 3, a genetically heterogeneous, rare a utosomal recessive disorder. [provided by RefSeq
Other Designations	unc-13 homolog D

Disease

- Arthritis
- Genetic Predisposition to Disease
- Lymphohistiocytosis
- Macrophage Activation Syndrome