

For Research Use Only

CLN3 Monoclonal antibody

Catalog Number: 67957-1-Ig



Basic Information

Catalog Number: 67957-1-Ig	GenBank Accession Number: BC002394	Purification Method: Protein G purification
Size: 150ul , Concentration: 1000 ug/ml by Nanodrop;	GeneID (NCBI): 1201	CloneNo.: 1E10A9
Source: Mouse	UNIPROT ID: Q13286	Recommended Dilutions: WB 1:5000-1:50000
Isotype: IgG1	Full Name: ceroid-lipofuscinosis, neuronal 3	
Immunogen Catalog Number: AG31402	Calculated MW: 438 aa, 48 kDa	
	Observed MW: 50 kDa	

Applications

Tested Applications: WB, ELISA	Positive Controls: WB : HeLa cells, HepG2 cells, NCCIT cells, NCI-H1299 cells, A549 cells, Jurkat cells
Species Specificity: Human	

Background Information

Neuronal ceroid lipofuscinosis (NCL, or Batten disease) refers to a group of lethal pediatric neurodegenerative diseases originating from mutations in one of the thus far identified 13 CLN genes (Ceroid Lipofuscinosis, Neuronal type; CLN1 to CLN14) (PMID: 25051496). CLN3 is a multi-membrane-spanning protein involved in the microtubule-dependent, anterograde transport of late endosomes and lysosomes. The CLN3 gene is located on chromosome 16p12.1 and produces three mRNA splicing variants. The 438-amino-acid CLN3 protein has a calculated molecular weight of 48 kDa. It has been reported that CLN3 can be glycosylated and form a homodimeric complex (PMID: 10356317; 17286803).

Storage

Storage:
Store at -20°C. Stable for one year after shipment.
Storage Buffer:
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

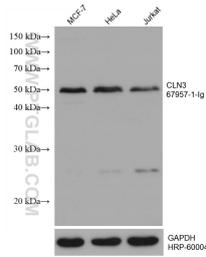
For technical support and original validation data for this product please contact:

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Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 67957-1-Ig (CLN3 antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours. The membrane was stripped and reblotted with HRP-conjugated GAPDH Monoclonal antibody (HRP-60004) as loading control.