For Research Use Only

CLN3 Monoclonal antibody, PBS Only

Catalog Number: 67957-1-PBS



Basic Information

Catalog Number:

GenBank Accession Number: BC002394 Purification Method: Protein G purification

67957-1-PBS

GeneID (NCBI):

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Size: 1mg/ml

1201

CloneNo.: 1E10A9

Source: Mouse

UNIPROT ID: Q13286 Full Name:

Isotype: IgG1

ceroid-lipofuscinosis, neuronal 3

Immunogen Catalog Number: AG31402

Calculated MW: 438 aa, 48 kDa

Observed MW:

50 kDa

Applications

Tested Applications:

WB, Indirect ELISA

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 that is involved in microtubule-dependent, anterograde transport of late endosomes and lysosomes. The CLN3 gene is located on chromosome 16p12.1and 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 homodimeric complex (PMID: 10356317; 17286803).

Storage

Storage:

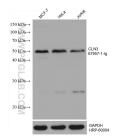
Store at -80°C.

The product is shipped with ice packs. Upon receipt, store it immediately at -80°C $\,$

Storage Buffer:

PBS Only

Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 67957-1-lg (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. This data was developed using the same antibody clone with 67957-1-PBS in a different storage buffer formulation.