For Research Use Only

CLN3 Monoclonal antibody

Catalog Number: 67957-1-Ig



Basic Information

Catalog Number: GenBank Accession Number: 67957-1-lg BC002394

 67957-1-lg
 BC002394

 Size:
 GeneID (NCBI):

 1000 μ g/ml
 1201

Source: UNIPROT ID: Recommended Dilutions: Mouse Q13286 WB 1:5000-1:50000

Isotype: Full Name:

IgG1 ceroid-lipofuscinosis, neuronal 3

Immunogen Catalog Number: Calculated MW: 438 aa, 48 kDa
Observed MW:

Observed N 50 kDa

Applications

Tested Applications:

WB, ELISA

Species Specificity:

Human

Positive Controls:

WB: HeLa cells, HepG2 cells, NCCIT cells, NCI-H1299

Purification Method:

Protein G purification

CloneNo.:

1E10A9

cells, A549 cells, Jurkat cells

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:

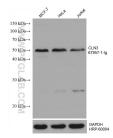
Store at -20°C. Stable for one year after shipment.

Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

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.