For Research Use Only

CLN3 Monoclonal antibody

Catalog Number:67957-1-Ig



| Basic Information | Catalog Number: 67957-1-lg | GenBank Accession Number: BC002394 | Purification Method: Protein G purification | | | | |
|---------------------------------|---|---------------------------------------|--|------------------------|--|---|--|
| | Size: 150ul, Concentration: 1000 µg/ml by Nanodrop; Source: Mouse Isotype: | GenelD (NCBI): | CloneNo.: 1E10A9 Recommended Dilutions: WB 1:5000-1:50000 | | | | |
| | | | | Applications | Tested Applications: | Positive Controls: WB : HeLa cells, HepG2 cells, NCCIT cells, NCI-H1299 cells, A549 cells, Jurkat cells | |
| | | | | | WB, 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 -20°C. Storage Buffer: PBS with 0.02% sodium azide and 50' | % glycerol pH 7.3. | |
| *** 20ul sizes contain 0.1% BSA | Aliquoting is unnecessary for -20 $^{\circ}$ C s | ••• | | | | | |
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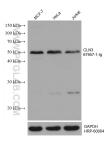
 For technical support and original validation data for this product please contact:

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Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 67957-1-1g (CLN3 antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours.