For Research Use Only

CoraLite® Plus 488-conjugated CLN3 Monoclonal antibody

www.ptglab.com

Catalog Number: CL488-67957

Basic Information

Catalog Number: GenBank Accession Number:

CL488-67957 BC002394 GeneID (NCBI):

100ul, Concentration: 1000 ug/ml by 1201 Nanodrop:

UNIPROT ID: Q13286 Mouse

Isotype: ceroid-lipofuscinosis, neuronal 3

lgG1 Calculated MW: Immunogen Catalog Number: 438 aa, 48 kDa AG31402 Observed MW:

50 kDa

Full Name:

Purification Method:

Protein G purification CloneNo.:

1E10A9 Excitation/Emission maxima

wavelengths: 493 nm / 522 nm

Applications

Tested Applications:

FC (Intra)

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~16p12.1 and~produces~three~mRNA~splicing~variants.~The~438-amino-acid~CLN3~protein~has~a~calculated~likely and the contraction of themolecular weight of 48 kDa. It has been reported that CLN3 can be glycosylated and form homodimeric complex (PMID: 10356317; 17286803).

Storage

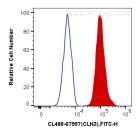
Store at -20°C. Avoid exposure to light. Stable for one year after shipment.

Storage Buffer:

PBS with 50% Glycerol, 0.05% Proclin300, 0.5% BSA, pH 7.3.

Aliquoting is unnecessary for -20°C storage

Selected Validation Data



1X10^6 HeLa cells were intracellularly stained with 0.4 ug CoraLite® Plus 488 Anti-Human CLN3 (CL488-67957, Clone:1E10A9) (red), or 0.4 ug Control Antibody. Cells were fixed with 4% PFA and permeabilized with Flow Cytometry Perm Buffer (PF00011-C).