For Research Use Only

## CoraLite® Plus 488-conjugated CLN3 Monoclonal antibody



**Purification Method:** 

Protein G purification

Excitation/Emission maxima

CloneNo.:

wavelengths: 488 nm / 515 nm

1E10A9

Catalog Number: CL488-67957

**Basic Information** 

Catalog Number: GenBank Accession Number:

CL488-67957 BC002394
Size: GeneID (NCBI):

100ul, Concentration: 1000 µg/ml by 1201

lanodrop; Full Name:

Source: ceroid-lipofuscinosis, neuronal 3

MouseCalculated MW:Isotype:438 aa, 48 kDaIgG1Observed MW:Immunogen Catalog Number:50 kDa

AG31402

**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 16p12.1and 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. Avoid exposure to light. Stable for one year after shipment.

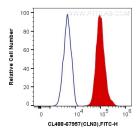
Storage Buffer

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

Aliquoting is unnecessary for -20°C storage

\*\*\* 20ul sizes contain 0.1% BSA

## 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).