



CLN3 rabbit pAb

Cat#: orb774308 (Manual)

For research use only. Not intended for diagnostic use.

Product Name CLN3 rabbit pAb

Host species Rabbit

Applications WB;ELISA

Species Cross-Reactivity Human; Rat; Mouse;

Recommended dilutions WB 1:500-2000 ELISA 1:5000-20000

Immunogen Synthesized peptide derived from human protein . at AA range: 221-270

Specificity CLN3 Polyclonal Antibody detects endogenous levels of protein.

Formulation Liquid in PBS containing 50% glycerol, and 0.02% sodium azide...

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name Battenin (Batten disease protein) (Protein CLN3)

Gene Name CLN3 BTS

Cellular localization Lysosome membrane; Multi-pass membrane protein. Late endosome.

Lysosome . Golgi apparatus . Golgi apparatus membrane . Golgi apparatus, Golgi stack . Golgi apparatus, trans-Golgi network . Cell membrane .

Recycling endosome. Membrane raft. Membrane, caveola. Early endosome

membrane . Cell junction, synapse, synaptosome . Late endosome membrane . Cytoplasmic vesicle, autophagosome . CLN3 is not present in late endosomes/lysosomes in fibroblasts and neurons (PubMed:15240864).

Trafficks from cell membrane to Golgi via endosomes (PubMed:15240864). Osmotic stress changes the subcellular localization of CLN3

(PubMed:23840424). Trafficks to intracellular compartments via the plasma membranet through AP3M1-dependent mechanisms (PubMed:14644441).



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Excluded from the synaptic vesicles (By simila

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

epitope-specific immunogen. chromatography using

Polyclonal **Clonality**

Concentration 1 mg/ml

Observed band 48kD

Human Gene ID 1201

Human Swiss-Prot Number Q13286

Alternative Names

Background

This gene encodes a protein that is involved in lysosomal function. Mutations in this, as well as other neuronal ceroid-lipofuscinosis (CLN) genes, cause neurodegenerative diseases commonly known as Batten disease or collectively known as neuronal ceroid lipofuscinoses (NCLs). Many alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Jul 2008],