



# Mouse Anti-CLN6 monoclonal antibody, clone O500/35 (CABT-RM159)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Specificity</b>	Specifically detects human Ceroid-lipofuscinosis neuronal protein 6. It does not cross-react with other CLN proteins.
<b>Target</b>	CLN6
<b>Immunogen</b>	Recombinant fragments corresponding to 54, 47, and 30 amino acids from non-helical regions from the N-terminal, internal, and C-terminal regions of human Ceroid-lipofuscinosis neuronal protein 6 (CLN6).
<b>Isotype</b>	IgG2b, κ
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Clone</b>	O500/35
<b>Purification</b>	Protein G purified
<b>Conjugate</b>	unconjugated
<b>Applications</b>	IHC
<b>Molecular Weight</b>	35.92 kDa calculated.
<b>Format</b>	Liquid
<b>Size</b>	100 µg
<b>Buffer</b>	0.1 M Tris-Glycine (pH 7.4), 150 mM NaCl

<b>Preservative</b>	0.05% sodium azide
<b>Storage</b>	Stable for 1 year at 2-8°C from date of receipt.

## BACKGROUND

<b>Introduction</b>	Ceroid-lipofuscinosis neuronal protein 6 is encoded by the CLN6 gene in human. CLN6 is a multi-pass membrane protein that is localized to endoplasmic reticulum and contributes to lysosomal function and viability of neurons. It has no homology with known proteins or functional domains, however it is highly conserved across mammalian species. Mutations in CLN6 gene have been linked to ceroid lipofuscinosis, neuronal 6 and 4A that are characterized by intracellular accumulation of autofluorescent liposomal material, seizures, dementia, visual loss, and/or cerebral atrophy. Two isoforms of CLN6 have been described that are produced by alternative splicing.
<b>Keywords</b>	CLN6; ceroid-lipofuscinosis, neuronal 6, late infantile, variant; ceroid-lipofuscinosis neuronal protein 6; FLJ20561; HsT18960; nclf; CLN4A

## GENE INFORMATION

<b>Entrez Gene ID</b>	<a href="#">54982</a>
<b>UniProt ID</b>	<a href="#">Q9NWW5</a>