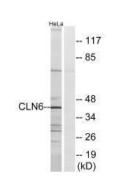
## **CLN6** Antibody

PACO23411



Product Information	
Size:	Protein Background:
100ul	This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses (NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive, neurodegenerative disorders affecting children. The genes responsible likely encode proteins involved in the degradation of post-translationally modified proteins in lysosomes. The primary defect in NCL disorders is thought to be associated with lysosomal storage function.
Reactivity:	
Human	
Source:	
Rabbit	Gene ID:
lsotype:	CLN6
IgG	Uniprot
Applications:	Q9NWW5
ELISA, WB, IHC	Synonyms:
Recommended dilutions:	ceroid-lipofuscinosis neuronal protein 6;
ELISA:1:2000-1:10000, WB:1:500-1:3000, IHC:1:50-1:100	Immunogen:
	Synthesized peptide derived from internal of human CLN6.
	Storage:
	Debbine Charles to the first dealer (1) the state of Color) and 7.4.450 and

Rabbit IgG in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.



Western blot analysis of extracts from HeLa cells, using CLN6 antibody.

Immunohistochemistry analysis of paraffin-embedded human cervix tissue using CLN6 antibody.

