## **CLN5 Antibody**



## **PACO22260**

Human

Source:

## **Product Information**

Size: Protein Background:

100ul This gene is one of eight which have been associated with neuronal ceroid

Reactivity: 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.

Rabbit Gene ID:

Isotype: CLN5

lgG Uniprot

**Applications:** O75503

ELISA, WB Synonyms:

**Recommended dilutions:** Ceroid-lipofuscinosis neuronal protein 5; Protein CLN5;

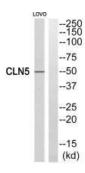
ELISA:1:2000-1:10000, WB:1:500-1:3000 **Immunogen:** 

Synthesized peptide derived from internal of human CLN5.

Storage:

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

## **Product Images**



Western blot analysis of extracts from LOVO cells, using CLN5 antibody.