

CLN5 polyclonal antibody

Catalog: BS72228

Host: Rabbit

Reactivity: Human, Mouse, Rat

BackGround:

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.

Product:

1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

~ 47 kDa

Swiss-Prot:

O75503

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

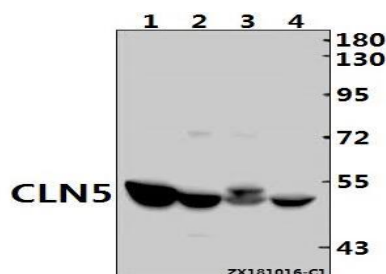
WB, 1:500 - 1:2000 | IHC, 1:50 - 1:200

Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

Polyclonal Antibodies

DATA:

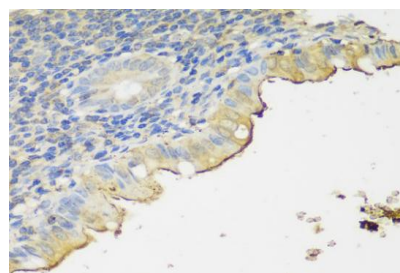
Western blot (WB) analysis of CLN5 pAb at 1:500 dilution

Lane1:PMVEC whole cell lysate(40ug)

Lane2:AML-12 whole cell lysate(40ug)

Lane3:L02 whole cell lysate(40ug)

Lane4:A549 whole cell lysate(40ug)



Immunohistochemistry of paraffin-embedded human appendix using CLN5 antibody at dilution of 1:150. Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.

Note:

For research use only, not for use in diagnostic procedure.

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