CLN2 antibody

Catalog No: #22320



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| CLN2 antibody | |
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| Rabbit | |
| Polyclonal | |
| Purified by antigen-affinity chromatography. | |
| WB IHC IF | |
| Hu | |
| Recombinant protein | |
| Recombinant protein fragment contain a sequence corresponding to a region within amino acids 66 and 294 of | |
| human TPP1 | |
| CLN2 | |
| Swiss-Prot:O14773Gene ID:1200 | |
| O14773 | |
| 1200; | |
| 0.6mg/ml | |
| Supplied in 0.1M Tris-buffered saline with 10% Glycerol (pH7.0). 0.01% Thimerosal was added as a | |
| preservative. | |
| Store at -20°C for long term preservation (recommended). Store at 4°C for short term use. | |
| | |

Application Details

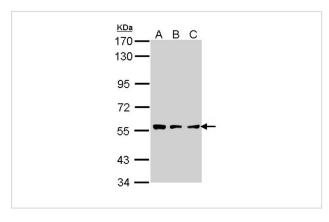
Predicted MW: 61kd

Western blotting: 1:500-1:3000

Immunohistochemistry: 1:100-1:250

Immunofluorescence: 1:100-1:200

Images



Sample (30 ug of whole cell lysate)

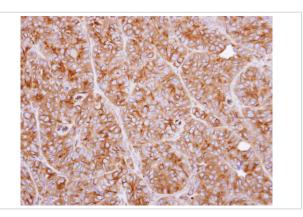
A: A431

B: H1299

C: Hela

7.5% SDS PAGE

Primary antibody diluted at 1: 1000



Immunohistochemical analysis of paraffin-embedded SW480 xenograft, using TPP1 antibody at 1: 100 dilution.



Immunofluorescence analysis of methanol-fixed A431, using TPP1 antibody at 1: 200 dilution.

Background

This gene encodes a member of the sedolisin family of serine proteases. The protease functions in the lysosome to cleave N-terminal tripeptides from substrates, and has weaker endopeptidase activity. It is synthesized as a catalytically-inactive enzyme which is activated and auto-proteolyzed upon acidification. Mutations in this gene result in late-infantile neuronal ceroid lipofuscinosis, which is associated with the failure to degrade specific neuropeptides and a subunit of ATP synthase in the lysosome. [provided by RefSeq]

Note: This product is for in vitro research use only