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CLN5 (YD15600) Rabbit mAb

货号: **AYD15032**

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB IHC-P ICC/IF
推荐浓度	
理论分子量	41kDa
实测分子量	
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	Mouse heart,Rat heart
细胞定位	Lysosome, Membrane
纯化	亲和纯化

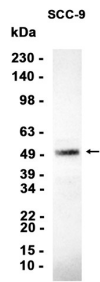
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靶点信息

研究背景	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.
基因ID	1203
基因名	CLN5
Swiss	O75503 (https://www.uniprot.org/uniprotkb/O75503/entry)
别名	CLN5 (YD15600), CLN5 (YD15600) Rabbit mAb, CLN5, Ceroid-lipofuscinosis neuronal protein 5, Palmitoyl protein thioesterase CLN5, S-depalmitoylase CLN5, BMPS

产品验证



实验步骤

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