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GBA (YD16799) Rabbit mAb

货号: AYD14795

产品信息

反应	Human
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB IHC-P
推荐浓度	
理论分子量	60kDa
实测分子量	
形式	Liquid
保存条件	Store at -20℃. Avoid freeze / thaw cycles. Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	MCF7
细胞定位	Lysosome membrane
纯化	亲和纯化

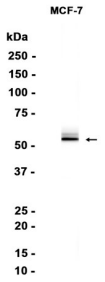
抗原信息

抗原信息	

靶点信息

研究背景	This gene encodes a lysosomal membrane protein that cleaves the beta-glucosidic linkage of glycosylceramide, an intermediate in glycolipid metabolism. Mutations in this gene cause Gaucher disease, a lysosomal storage disease characterized by an accumulation of glucocerebrosides. A related pseudogene is approximately 12 kb downstream of this gene on chromosome 1. Alternative splicing results in multiple transcript variants.
基因ID	2629
基因名	GBA1
Swiss	P04062 (https://www.uniprot.org/uniprotkb/P04062/entry)
别名	GBA (YD16799),GBA (YD16799) Rabbit mAb,GBA1,Acid beta-glucosidase,Alglucerase,Beta-glucocerebrosidase,Beta-glucosylceramidase 1,Cholesterol glucosyltransferase,Cholesteryl-beta-glucosidase,D-glucosyl-N-acylsphingosine glucohydrolase,Glucosylceramidase beta 1

产品验证



实验步骤

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