GBA Rabbit mAb

Catalog No: #59345

Package Size: #59345-1 50ul #59345-2 100ul



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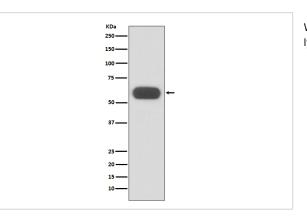
Description

Product Name	GBA Rabbit mAb
Host Species	Rabbit
Clonality	Monoclonal
sotype	Rabbit IgG
Purification	Affinity-chromatography
Applications	WB IHC Ms
Species Reactivity	Human Rat
Specificity	GBA Antibody detects endogenous levels of total GBA
mmunogen Description	A synthesized peptide derived from human GBA
Other Names	Alglucerase; betaGC; GBA1; GCase; GCB; GLUC; Glucosylceramidase; Imiglucerase;
Accession No.	Uniprot:P04062
Jniprot	P04062
Formulation	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Storage	Store at +4°C short term. Store at -20°C long term. Avoid freeze / thaw cycle.

Application Details

WB 1:500~1:2000 IHC 1:50~1:200

Images



Western blot analysis of GBA expression in U87-MG cell lysate.

Product Description

Defects in GBA are the cause of Gaucher disease (GD) [MIM:230800]; also known as glucocerebrosidase deficiency. GD is the most prevalent lysosomal storage disease, characterized by accumulation of glucosylceramide in the reticulo-endothelial system.

Background

Defects in GBA are the cause of Gaucher disease (GD) [MIM:230800]; also known as glucocerebrosidase deficiency. GD is the most prevalent lysosomal storage disease, characterized by accumulation of glucosylceramide in the reticulo-endothelial system.

Note: This product is for in vitro research use only