

## Galactosidase alpha Rabbit mAb

Catalog No: #56651

Package Size: #56651-1 50ul #56651-2 100ul

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## Description

Product Name	Galactosidase alpha Rabbit mAb
Host Species	Rabbit
Clonality	Monoclonal
Isotype	Rabbit IgG
Purification	Affinity-chromatography
Applications	WB IHC ICC/IF IP FC
Species Reactivity	Human
Specificity	Galactosidase alpha Antibody detects endogenous levels of total Galactosidase alpha
Immunogen Description	A synthesized peptide derived from human Galactosidase alpha
Other Names	Alpha gal A; GALA; Galactosidase, alpha; GLA; Melibiase;
Accession No.	Uniprot:P06280
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Calculated MW	46kDa
Formulation	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Storage	Store at +4 $\Lambda$ C short term. Store at -20 $\Lambda$ C long term. Avoid freeze / thaw cycle.

## Application Details

WB: 1:500~1:2000

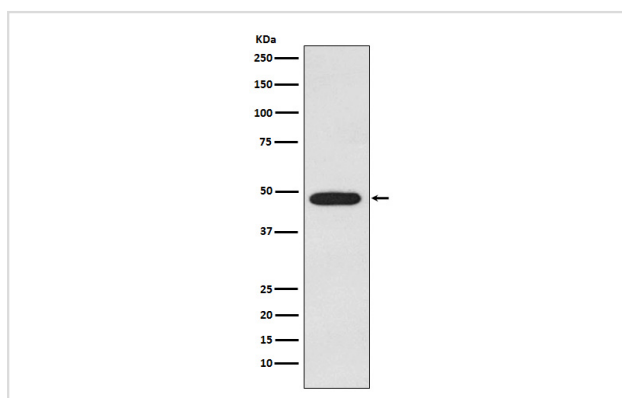
IHC: 1:50~1:200

ICC/IF: 1:50~1:200

IP: 1:50

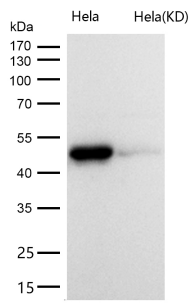
FC: 1:80

## Images



Western blot analysis of Galactosidase alpha expression in MCF-7 cell lysate.

All lanes use the Antibody at 1:1K dilution for 1 hour at room temperature.



## Product Description

Defects in GLA are the cause of Fabry disease (FD) [MIM:301500]. FD is a rare X-linked sphingolipidosis disease where glycolipid accumulates in many tissues. The disease consists of an inborn error of glycosphingolipid catabolism.

Note: This product is for in vitro research use only