# Product Name: GBA (1P9) Rabbit Monoclonal Antibody Catalog #: AMRe11321

# **Summary**

**Production Name** GBA (1P9) Rabbit Monoclonal Antibody

**Description** Rabbit Monoclonal Antibody

HostRabbitApplicationWB,IHC-PReactivityHuman

# **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

Clonality Monoclonal Form Liquid

**Storage** Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.

Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% New type

**Buffer** preservative N and 50% glycerol. Store at +4°C short term. Store at -20°C long term.

Avoid freeze / thaw cycle.

**Purification** Affinity purification

## **Immunogen**

Gene Name GBA

Alglucerase; betaGC; GBA1; GCase; GCB; GLUC; Glucosylceramidase; Imiglucerase;

 Gene ID
 2629.0

 SwissProt ID
 P04062.

# **Application**

**Dilution Ratio** WB 1:1000-1:5000, IHC-P/IF-P 1:50-1:100

Molecular Weight 60kDa

# **Background**

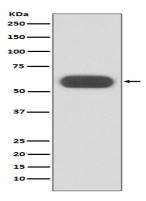
# Product Name: GBA (1P9) Rabbit Monoclonal Antibody Catalog #: AMRe11321



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 reticuloendothelial system. Glucosylceramidase that catalyzes, within the lysosomal compartment, the hydrolysis of glucosylceramide/GlcCer into free ceramide and glucose (PubMed: <a href="http://www.uniprot.org/citations/9201993" target=" blank">9201993</a>, PubMed:<a href="http://www.uniprot.org/citations/24211208" target=" blank">24211208</a>, PubMed:<a href="http://www.uniprot.org/citations/15916907" target="\_blank">15916907</a>). Thereby, plays a central role in the degradation of complex lipids and the turnover of cellular membranes (PubMed: <a href="http://www.uniprot.org/citations/27378698" target=" blank" > 27378698 < /a>). Through the production of ceramides, participates in the PKC-activated salvage pathway of ceramide formation (PubMed: <a href="http://www.uniprot.org/citations/19279011" target=" blank">19279011</a>). Also plays a role in cholesterol metabolism (PubMed: <a href="http://www.uniprot.org/citations/24211208" target=" blank">24211208</a>, PubMed:<a href="http://www.uniprot.org/citations/26724485" target=" blank">26724485</a>). May either catalyze the glucosylation of cholesterol, through a transglucosylation reaction that transfers glucose from glucosylceramide to cholesterol (PubMed: <a href="http://www.uniprot.org/citations/24211208" target=" blank">24211208</a>, PubMed: <a href="http://www.uniprot.org/citations/26724485" target=" blank">26724485</a>). The short chain saturated C8:0-GlcCer and the mono-unsaturated C18:0-GlcCer being the most effective glucose donors for that transglucosylation reaction (PubMed: <a href="http://www.uniprot.org/citations/24211208" target=" blank">24211208</a>). Under specific conditions, may alternatively catalyze the reverse reaction, transferring glucose from cholesteryl-beta-D-glucoside to ceramide (PubMed: <a href="http://www.uniprot.org/citations/26724485" target=" blank">26724485</a>). Finally, may also hydrolyze cholesteryl- beta-D-glucoside to produce D-glucose and cholesterol (PubMed: <a href="http://www.uniprot.org/citations/24211208" target=" blank">24211208</a>, PubMed:<a href="http://www.uniprot.org/citations/26724485" target=" blank">26724485</a>).

#### **Research Area**

### **Image Data**



Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838

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

### Note

For research use only.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838