

## Basic Information

<b>Product Name</b>	Anti-Beta Galactosidase/GLB1 Antibody	
<b>Gene Name</b>	GLB1	
<b>Source</b>	Rabbit	
<b>Clonality</b>	Polyclonal	
<b>Isotype</b>	IgG	
<b>Species Reactivity</b>	human	
<b>Tested Application</b>	WB, IHC, ICC/IF, FCM, ELISA	
<b>Contents</b>	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
<b>Immunogen</b>	E.coli-derived human GLB1/Beta-galactosidase recombinant protein (Position: Q46-K655).	
<b>Concentration</b>	500 ug/ml	
<b>Purification</b>	Immunogen affinity purified.	
<b>Observed MW</b>	65-85 kDa	
<b>Dilution Ratios</b>	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-400
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400
	Flow Cytometry (Fixed):	1:50-200
	Enzyme linked immunosorbent assay (ELISA):	1:100-1000
	(Boiling the paraffin sections in 10mM citrate buffer,pH6.0,or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

## Storage

12 months from date of receipt, -20°C as supplied.

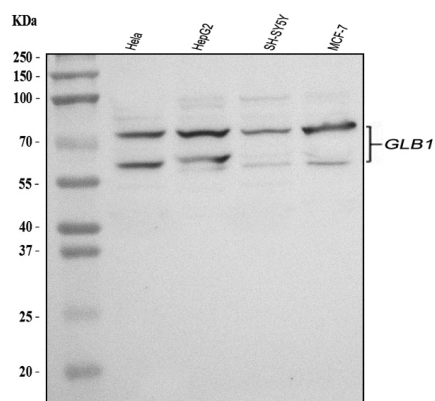
## Background Information

Galactosidase, beta 1, also known as GLB1, is a protein which in humans is encoded by the GLB1 gene. It is mapped to 3p22.3. This gene encodes a member of the glycosyl hydrolase 35 family of proteins. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed to generate the mature lysosomal enzyme. This enzyme catalyzes the hydrolysis of a terminal beta-linked galactose residue from ganglioside substrates and other glycoconjugates. Mutations in this gene may result in GM1-gangliosidosis and Morquio B syndrome.

## Reference

Anti-Beta Galactosidase/GLB1 Antibody被引用在2文献中。

## Selected Validation Data



Western blot analysis of anti- GLB1 antibody (A01829-2). The sample well of each lane was loaded with 30ug of sample under reducing conditions.

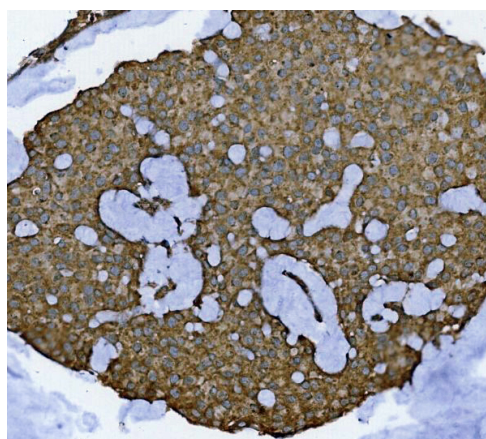
Lane 1: human HeLa whole cell lysates,

Lane 2: human HepG2 whole cell lysates,

Lane 3: human SH-SY5Y whole cell lysates,

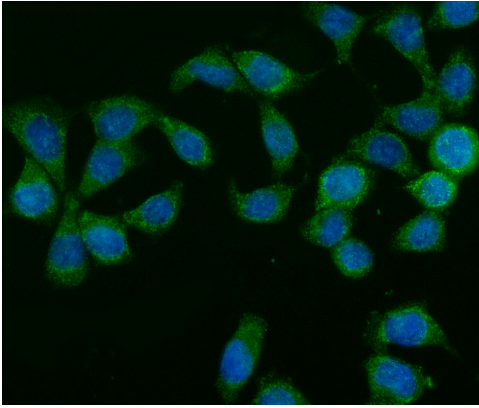
Lane 4: human MCF-7 whole cell lysates.

Use rabbit anti- GLB1 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog#EK1002). A specific band was detected for GLB1 at approximately 65-85KD. The expected band size for GLB1 is at 76KD.



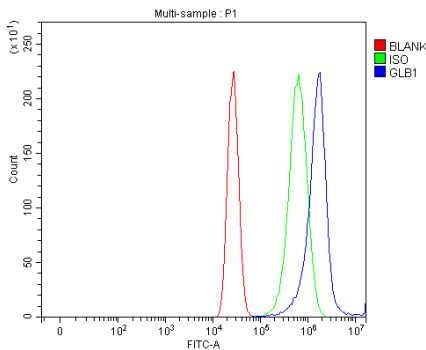
IHC analysis of Beta Galactosidase/GLB1 using anti-Beta Galactosidase/GLB1 antibody (A01829-2).

Beta Galactosidase/GLB1 was detected in a paraffin-embedded section of human breast cancer tissue. The tissue section was incubated with rabbit anti-Beta Galactosidase/GLB1 Antibody (A01829-2) at a dilution of 1:200 and developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB (Catalog # AR1027) as the chromogen.



ICC/IF analysis of Beta Galactosidase/GLB1 using anti-Beta Galactosidase/GLB1 antibody (A01829-2).

Beta Galactosidase/GLB1 was detected in an immunocytochemical section of Caco-2 cells. The section was incubated with rabbit anti-Beta Galactosidase/GLB1 Antibody (A01829-2) at a dilution of 1:100. Fluoro488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of MCF-7 cells using anti-Beta Galactosidase/GLB1 antibody (A01829-2).

Overlay histogram showing MCF-7 cells stained with A01829-2 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Beta Galactosidase/GLB1 Antibody (A01829-2) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.