## For Research Use Only

## CoraLite® Plus 488-conjugated Alpha Galactosidase A Monoclonal antibody

Catalog Number:CL488-66121 Featured Product

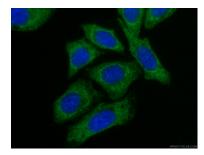


488-66121 e: Dul, Concentration: 1000 ug/ml by nodrop; urce: use type: i2a munogen Catalog Number: 7505	BC002689 GeneID (NCBI): 2717 UNIPROT ID: P06280 Full Name: galactosidase, alpha Calculated MW: 49 kDa Observed MW:	Protein A purification CloneNo.: 2B2C5 Recommended Dilutions: IF/ICC 1:50-1:500 Excitation/Emission maxima wavelengths: 493 nm / 522 nm
	49 kDa	
sted Applications: ICC ecies Specificity: nan	Positive ( IF/ICC : F	Controls: lepG2 cells,
minal, non-reducing alpha-D-gala actomannans and galactolipids. F icient activity of GLA. Enzyme rep	ctose residues in alpha-D-galactos abry disease is an X-linked lysoso lacement therapy (ERT) with GLA i	sides, including galactose oligosaccharides, mal storage disorder resulting from the
re at -20°C. Avoid exposure to ligh rage Buffer: 5 with 50% Glycerol, 0.05% Proclin	n300, 0.5% BSA, pH 7.3.	ent.
GLA, also named as Melibiase and Alpha-galactosidase A, belongs to the glycosyl hydrolase 27 family. It hydrolyst terminal, non-reducing alpha-D-galactose residues in alpha-D-galactosides, including galactose oligosaccharides galactomannans and galactolipids. Fabry disease is an X-linked lysosomal storage disorder resulting from the deficient activity of GLA. Enzyme replacement therapy (ERT) with GLA is currently the most effective therapeutic strategy for patients with Fabry disease, a lysosomal storage disease. Storage: Store at -20°C. Avoid exposure to light. Stable for one year after shipment. Storage Buffer: PBS with 50% Glycerol, 0.05% Proclin300, 0.5% BSA, pH 7.3. Aliquoting is unnecessary for -20°C storage		a, also named as Melibiase and Alpha-galactosidase A, belongs to the ninal, non-reducing alpha-D-galactose residues in alpha-D-galactos actomannans and galactolipids. Fabry disease is an X-linked lysoso icient activity of GLA. Enzyme replacement therapy (ERT) with GLA is tegy for patients with Fabry disease, a lysosomal storage disease. rage: re at -20°C. Avoid exposure to light. Stable for one year after shipme rage Buffer: with 50% Glycerol, 0.05% Proclin300, 0.5% BSA, pH 7.3.

For technical support and original validation data for this product please contact: T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free E: proteintech@ptglab.com in USA), or 1(312) 455-8498 (outside USA) W: ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

## Selected Validation Data



Immunofluorescent analysis of (-20°C Ethanol) fixed HepG2 cells using CL488-66121 (Alpha galactosidase A antibody) at dilution of 1:100.