

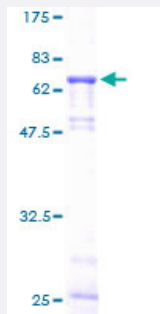
Full-Length

GLA (Human) Recombinant Protein (P01)

Catalog # H00002717-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human GLA full-length ORF (AAH02689, 1 a.a. - 429 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MQLRNPELHLGCALALRFLALVSWDIPGARALDNGLARTPTMGWLHWERFMCNLDCQEEDSCI
SEKLFMEMAELMVSEGWKDAGYEYLCIDDCWMAQRDSEGRQLQADPQRFPHGIRQLANYVHSK
GLKLGIVADVGNKTCAGFPGSFGYYDIDAQTFADWGVDLLKFDGCYCDSENLADGYKHMSLALN
RTGRSIVYSCEWPLYMWPFQKPNYTEIRQYCNHWRNFADIDDSWKSISILDWTSFNQERVDVAG
PGGWNDPDMVLVIGNFGLSWNQVQTQMALWAIMAAPLFMSNDLRHISPAKALLQDKDVIINQD
PLGKQGYQLRQGDNFEVWERPLSGLAWAVAMINRQEIGGPRSYTIAVASLGKGVACNPACFITQL
LPVKRKLGIFYEWTSLRSHINPTGTVLLQLENTMQMSLKDLL

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

72.93

Preparation Method

[in vitro wheat germ expression system](#)

Purification

Glutathione Sepharose 4 Fast Flow

Quality Control Testing

12.5% SDS-PAGE Stained with Coomassie Blue.

Storage Buffer

50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

Storage Instruction

Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Note

Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — GLA

Entrez GeneID [2717](#)

GeneBank Accession# [BC002689](#)

Protein Accession# [AAH02689](#)

Gene Name GLA

Gene Alias GALA

Gene Description galactosidase, alpha

Omim ID [300644](#) [301500](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes a homodimeric glycoprotein that hydrolyses the terminal alpha-galactosyl moieties from glycolipids and glycoproteins. This enzyme predominantly hydrolyzes ceramide trihexoside, and it can catalyze the hydrolysis of melibiose into galactose and glucose. A variety of mutations in this gene affect the synthesis, processing, and stability of this enzyme, which causes Fabry disease, a rare lysosomal storage disorder that results from a failure to catabolize alpha-D-galactosyl glycolipid moieties. [provided by RefSeq]

Other Designations OTTHUMP00000023679|agalsidase alfa|alpha-D-galactosidase A|alpha-D-galactoside galactohydrolase 1|alpha-gal A|alpha-galactosidase A|melibiase

Publication Reference

- [Integrated approach to produce a recombinant, his-tagged human alpha-galactosidase A in mammalian cells.](#)

Corchero JL, Mendoza R, Lorenzo J, Rodriguez-Sureda V, Dominguez C, Vazquez E, Ferrer-Miralles N, Villaverde A.

Biotechnology Progress 2011 Sep; 27(5):1206.

Application: WB, Rabbit, Serum

Pathway

- [Galactose metabolism](#)
- [Glycerolipid metabolism](#)
- [Glycosphingolipid biosynthesis - globo series](#)
- [Lysosome](#)
- [Sphingolipid metabolism](#)

Disease

- [Cardiomyopathy](#)
- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)
- [Fabry Disease](#)
- [Hypertrophy](#)
- [Lipid Metabolism](#)
- [Mitochondrial Myopathies](#)
- [Stroke](#)