

GLA Polyclonal Antibody

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|-------------------------|--------|----------------------|-----------------------|
| Catalog No. | A13987 | Category | Polyclonal Antibodies |
| Applications | WB | Observed MW | 49kDa |
| Cross-reactivity | Human | Calculated MW | 48kDa |

Immunogen Information

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|-------------------|---|
| Immunogen | Recombinant fusion protein containing a sequence corresponding to amino acids 150-429 of human GLA (NP_000160.1). |
| Gene ID | 2717 |
| Swiss prot | P06280 |
| Synonyms | GLA; GALA; galactosidase alpha |

Product information

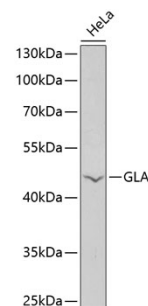
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|----------------------------|--|
| Source | Rabbit |
| Isotype | IgG |
| Purification method | Affinity purification |
| Storage | Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3. |

Background

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.

Recommended Dilutions

WB 1:500 -
1:2000



Western blot - GLA Polyclonal Antibody (A13987)