

GBE1 Polyclonal Antibody

| Catalog No. | A6599 | Category | Polyclonal Antibodies |
|----------------------------|--|---------------|--|
| Applications | WB | Observed MW | 75kDa |
| Cross-reactivity | Human, Mouse, Rat | Calculated MW | 80kDa |
| Immunogen Information | | | Recommended Dilutions |
| Immunogen | Recombinant fusion protein containing a sequence corresponding to amino acids 1-300 of human GBE1 (NP_000149.3). | | WB 1:500 - 1:2000 |
| Gene ID | Q04446 | | Hero ^{C2} HARD Mouse heart inter kildred |
| Swiss prot | | | 130KD- 70KD- 70KD- |
| Synonyms | GBE1; APBD; GBE; GSD4; 1,4-alpha-glucan-branch ing enzyme | | |
| Product information | | | 55KD- |
| Source | Rabbit | | |
| Isotype | IgG Western blot - GBET Polyce Antibody (A6599) Affinity purification | | Western blot - GBE1 Polyclonal Antibody (A6599) |
| Purification method | | | |
| Storage | Store at -20°C. Avoid freeze / thaw cyc Buffer: PBS with 0.02% sodium azide, 5 | | |

Background

The protein encoded by this gene is a glycogen branching enzyme that catalyzes the transfer of alpha-1,4-linked glucosyl units from the outer end of a glycogen chain to an alpha-1,6 position on the same or a neighboring glycogen chain. Branching of the chains is essential to increase the solubility of the glycogen molecule and, consequently, in reducing the osmotic pressure within cells. Highest level of this enzyme are found in liver and muscle. Mutations in this gene are associated with glycogen storage disease IV (also known as Andersen's disease).