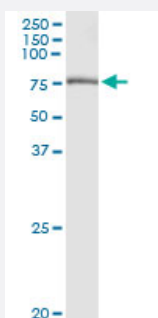


# GBE1 (Human) IP-WB Antibody Pair

Catalog # H00002632-PW1

Size 1 Set

## Applications



Immunoprecipitation of GBE1 transfected lysate using rabbit polyclonal anti-GBE1 and Protein A Magnetic Bead ([U0007](#)), and immunoblotted with mouse purified polyclonal anti-GBE1.

## Specification

|                                      |   |
|--------------------------------------|---|
| <b>Product Description</b>           | This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.  |
| <b>Reactivity</b>                    | Human   |
| <b>Interspecies Antigen Sequence</b> | Mouse (91%); Rat (91%)  |
| <b>Quality Control Testing</b>       | Immunoprecipitation-Western Blot (IP-WB)<br>Immunoprecipitation of GBE1 transfected lysate using rabbit polyclonal anti-GBE1 and Protein A Magnetic Bead ( <a href="#">U0007</a> ), and immunoblotted with mouse purified polyclonal anti-GBE1. |
| <b>Supplied Product</b>              | Antibody pair set content:<br>1. Antibody pair for IP: rabbit polyclonal anti-GBE1 (300 ul)<br>2. Antibody pair for WB: mouse purified polyclonal anti-GBE1 (50 ug)   |
| <b>Storage Instruction</b>           | Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.  |

## Applications

- Immunoprecipitation-Western Blot

[Protocol Download](#)

## Gene Info — GBE1

Entrez GeneID [2632](#)

Gene Name GBE1

Gene Alias GBE

Gene Description glucan (1,4-alpha-), branching enzyme 1

Omim ID [232500 607839](#)

Gene Ontology [Hyperlink](#)

**Gene Summary** 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). [provided by RefSeq]

**Other Designations** amylo-(1,4 to 1,6) transglucosidase|amylo-(1,4 to 1,6) transglycosylase|glycogen branching enzyme|glycogen storage disease type IV

## Pathway

- [Metabolic pathways](#)
- [Starch and sucrose metabolism](#)

## Disease

- [Asthma](#)
- [Cardiovascular Diseases](#)
- [Coronary Disease](#)
- [Diabetes Mellitus](#)

- [Edema](#)
- [Genetic Predisposition to Disease](#)
- [Metabolic Diseases](#)
- [Metabolic Syndrome X](#)
- [Obesity](#)
- [Tobacco Use Disorder](#)