# **G6PC Polyclonal Antibody**

Catalog Number: E-AB-92332



Note: Centrifuge before opening to ensure complete recovery of vial contents.

## Description

**Reactivity** Mouse,Rat

**Immunogen** A synthetic peptide of human G6PC

Host Rabbit
Isotype IgG

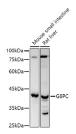
Purification Affinity purification
Conjugation Unconjugated

**Formulation** PBS with 0.05% proclin300,50% glycerol,pH7.3.

**Applications Recommended Dilution** 

**WB** 1:500-1:2000

### Data



Western blot analysis of extracts of various cell lines using G6PC Polyclonal Antibody at 1:1000 dilution.

Observed MW:40KDa Calculated Mw:40KDa

## **Preparation & Storage**

Storage Store at -20°C. Avoid freeze/thaw cycles.

### **Background**

Glucose-6-phosphatase (G6Pase) is a multi-subunit integral membrane protein of the endoplasmic reticulum that is composed of a catalytic subunit and transporters for G6P, inorganic phosphate, and glucose. This gene (G6PC) is one of the three glucose-6-phosphatase catalytic-subunit-encoding genes in human: G6PC, G6PC2 and G6PC3.

Glucose-6-phosphatase catalyzes the hydrolysis of D-glucose 6-phosphate to D-glucose and orthophosphate and is a key enzyme in glucose homeostasis, functioning in gluconeogenesis and glycogenolysis. Mutations in this gene cause glycogen storage disease type I (GSD1). This disease, also known as von Gierke disease, is a metabolic disorder characterized by severe hypoglycemia associated with the accumulation of glycogen and fat in the liver and kidneys.

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Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017

Web: www.elabscience.com Email: techsupport@elabscience.com