# **GPD1L Rabbit pAb**

Catalog No.: A14392



### **Basic Information**

#### **Observed MW**

38kDa

### **Calculated MW**

38kDa

### Category

Primary antibody

### **Applications**

ELISA,WB

### **Cross-Reactivity**

Human, Mouse

## **Background**

The protein encoded by this gene catalyzes the conversion of sn-glycerol 3-phosphate to glycerone phosphate. The encoded protein is found in the cytoplasm, associated with the plasma membrane, where it binds the sodium channel, voltage-gated, type V, alpha subunit (SCN5A). Defects in this gene are a cause of Brugada syndrome type 2 (BRS2) as well as sudden infant death syndrome (SIDS).

## **Recommended Dilutions**

**WB** 

1:500 - 1:2000

## **Immunogen Information**

**Gene ID** 23171

Swiss Prot

Q8N335

### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-351 of human GPD1L (NP\_055956.1).

## **Synonyms**

GPD1-L; GPD1L

### **Contact**

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## **Product Information**

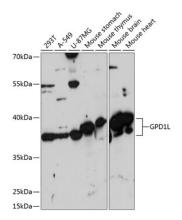
SourceIsotypePurificationRabbitIgGAffinity purification

### **Storage**

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

Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.

## **Validation Data**



Western blot analysis of various lysates using GPD1L Rabbit pAb (A14392) at 1:3000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 90s.