

## GPD1L Antibody (N-term)

Catalog\_no: AB0821

Applications: IHC-P, WB

Reactivity: H, M

Category: 抗原抗体

Size:  $100\mu L/50\mu L$ 

Immunogen: HUMAN:44-73

Specificity: This GPD1L antibody is generated from rabbits immunized with a KLH conjugated

synthetic peptide between 44-73 amino acids from the N-terminal region of human

GPD1L.

Dilution: WB,1:1000;IHC-P,1:50~100;FC,1:10~50;

Purification: Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This

antibody is purified through a protein A column, followed by peptide affinity

purification.

Other\_name: Glycerol-3-phosphate dehydrogenase 1-like protein, GPD1-L, GPD1L, KIAA0089

Isotype: Rabbit Ig

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).

reference: Rose, J.E., et al. Mol. Med. 16 (7-8), 247-253 (2010): Liu, M., et al. Circ. Res.

105(8):737-745(2009) Valdivia, C.R., et al. Am. J. Physiol. Heart Circ. Physiol. 297 (4),

H1446-H1452 (2009) : Makiyama, T., et al. Circ. J. 72(10):1705-1706(2008) Lon