## **INPP5E Antibody**



## PACO60356

Reactivity:

## **Product Information**

Size: Protein Background:

50ug Converts phosphatidylinositol 3,4,5-trisphosphate (PtdIns 3,4,5-P3) to PtdIns-P2, and

phosphatidylinositol 4,5-bisphosphate to phosphatidylinositol 4-phosphate. Specific for

lipid substrates, inactive towards water soluble inositol phosphates.

Human Gene ID:

Source: INPP5E

Rabbit **Uniprot** 

**Isotype:** Q9NRR6

lgG Synonyms:

**Applications:** 72 kDa inositol polyphosphate 5-phosphatase (EC 3.1.3.36) (Phosphatidylinositol 4,5-

ELISA, IHC, IF bisphosphate 5-phosphatase) (Phosphatidylinositol polyphosphate 5-phosphatase type

IV), INPP5E

Recommended dilutions: Immunogen:

ELISA:1:2000-1:10000, IHC:1:200-1:500,

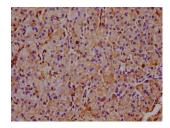
IF:1:50-1:200

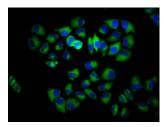
Recombinant Human 72 kDa inositol polyphosphate 5-phosphatase protein (1-133AA).

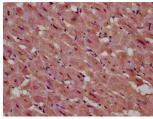
Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

## **Product Images**







IHC image of PACO60356 diluted at 1:400 and staining in paraffinembedded human pancreatic tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunofluorescence staining of Hela cells with PACO60356 at 1:133, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

IHC image of PACO60356 diluted at 1:400 and staining in paraffinembedded human heart tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.