## **PLA2G4B Antibody**



## PACO62959

Reactivity:

Human

Source:

## **Product Information**

Size: Protein Background:

50ul Calcium-dependent phospholipase A2 that selectively hydrolyzes glycerophospholipids

in the sn-2 position with a preference for arachidonoyl phospholipids. Has a much weaker activity than PLA2G4A. Isoform 3 has calcium-dependent activity against

palmitoyl-arachidonyl-phosphatidylethanolamine and low level lysophospholipase

activity but no activity against phosphatidylcholine. Isoform 5 does have activity against

phosphatidylcholine.

Rabbit Gene ID:

**Isotype:** PLA2G4B

lgG **Uniprot** 

**Applications:** POC869

ELISA, WB, IHC, IF Synonyms:

Recommended dilutions: Cytosolic phospholipase A2 beta (cPLA2-beta) (EC 3.1.1.4) (Phospholipase A2 group

IVB), PLA2G4B

ELISA:1:2000-1:10000, WB:1:1000-1:5000, IHC:1:200-1:500, IF:1:50-1:200

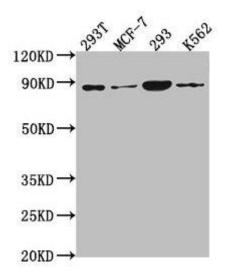
Recombinant Human Cytosolic phospholipase A2 β protein (514-781AA).

Storage:

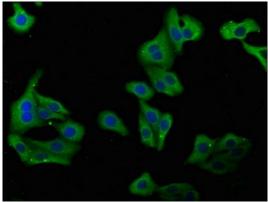
Immunogen:

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

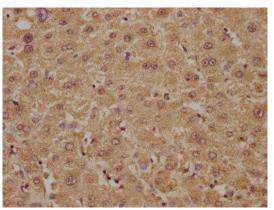
## **Product Images**



Western Blot. Positive WB detected in: 293T whole cell lysate, MCF-7 whole cell lysate, 293 whole cell lysate, K562 whole cell lysate. All lanes: P0C869 antibody at 1:2000. Secondary. Goat polyclonal to rabbit IgG at 1/50000 dilution. Predicted band size: 88, 101, 55, 115 kDa. Observed band size: 88 kDa.



Immunofluorescence staining of HepG2 cells with PACO62959 at 1:100, 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 PACO62959 diluted at 1:300 and staining in paraffinembedded human liver 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.