

# **GPR92** polyclonal antibody

| Catalog: BS | S6' | 768 |
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Host:

Rabbit

Reactivity: Hu

: Human, Mouse, Rat

## **BackGround:**

Receptor for lysophosphatidic acid (LPA), a mediator of diverse cellular activities.

## **Product:**

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

**Molecular Weight:** 

## ~ 180 kDa

**Swiss-Prot:** 

#### Q9H1C0

## **Purification&Purity:**

The antibody was purified by immunogen affinity chromatography.

## **Applications:**

WB (1/500 - 1/1000), IHC (1/50 - 1/200), IF/ICC (1/50 - 1/200)

## **Storage&Stability:**

Store at  $4 \ \mathbb{C}$  short term. Aliquot and store at  $-20 \ \mathbb{C}$  long term. Avoid freeze-thaw cycles.

#### **Specificity:**

Recognizes endogenous levels of GPR92 protein.

#### **DATA:**



Western blot analysis of GPR92 expression in HuT78 (A), H1792 (B),

A375 (C), mouse spleen (D), mouse heart (E), rat spleen (F), rat heart (G) whole cell lysates.



Immunohistochemical analysis of GPR92 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of GPR92 staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. Note:

For research use only, not for use in diagnostic procedure.

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