

# Anti-GPR86 Antibody

Rabbit polyclonal antibody to GPR86 Catalog # AP60602

#### **Product Information**

Application WB, IP, IF/IC
Primary Accession Q9BPV8
Other Accession Q9D8I2

**Reactivity** Human, Mouse, Rat, Monkey

Host Rabbit
Clonality Polyclonal
Calculated MW 40789

#### **Additional Information**

**Gene ID** 53829

Other Names GPR86; GPR94; P2Y purinoceptor 13; P2Y13; G-protein coupled receptor 86;

G-protein coupled receptor 94

**Target/Specificity** Recognizes endogenous levels of GPR86 protein.

Dilution WB~~WB (1/500 - 1/1000), IF/IC (1/100 - 1/500), IP (1/10 - 1/100) IP~~N/A

IF/IC~~N/A

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

**Storage** Store at -20 °C.Stable for 12 months from date of receipt

#### **Protein Information**

Name P2RY13

**Synonyms** GPR86, GPR94

**Function** Receptor for ADP. Coupled to G(i)-proteins. May play a role in hematopoiesis

and the immune system.

**Cellular Location** Cell membrane; Multi-pass membrane protein.

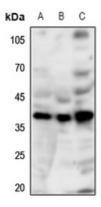
**Tissue Location** Strong expression in spleen and adult brain. Lower expression in placenta,

lung, liver, spinal cord, thymus, small intestine, uterus, stomach, testis, fetal brain, and adrenal gland. Not detected in pancreas, heart, kidney, skeletal muscle, ovary or fetal aorta. Clearly detected in lymph node and bone marrow, weakly detected in peripheral blood mononuclear cells (PBMC) and in peripheral blood leukocytes (PBL), but not detected in polymorphonuclear

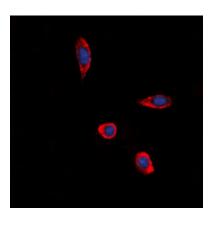
## **Background**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human GPR86. The exact sequence is proprietary.

### **Images**



Western blot analysis of GPR86 expression in HEK293T (A), Hela (B), HepG2 (C) whole cell lysates.



Immunofluorescent analysis of GPR86 staining in HeLa 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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