

## Product datasheet for AP51899PU-N

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OriGene Technologies, Inc.

## **GPAM (Center) Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** FC, WB

Recommended Dilution: ELISA: 1/1000.

**Western Blot:** 1/100 - 1/500. **Flow Cytometry:** 1/10 - 1/50.

Reactivity: Human Rabbit

**Isotype:** lg

**Clonality:** Polyclonal

Immunogen: KLH conjugated synthetic peptide between 433-463 amino acids from the Central region of

human GPAM

**Specificity:** This antibody reacts to GPAM.

Formulation: PBS

State: Aff - Purified

State: Liquid purified Ig fraction

Preservative: 0.09% (W/V) sodium azide

**Concentration:** lot specific

**Purification:** Affinity chromatography on Protein A

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

Predicted Protein Size: 93795 Da

**Gene Name:** glycerol-3-phosphate acyltransferase, mitochondrial

Database Link: Entrez Gene 57678 Human

Q9HCL2





**Background:** Glycerol-3-phosphate acyltransferase (GPAT; EC 2.3.1.15), which catalyzes the initial and

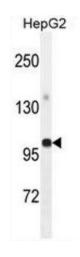
committing step in glycerolipid biosynthesis, is predicted to play a pivotal role in the regulation of cellular triacylglycerol and phospholipid levels. Two mammalian forms of GPAT have been identified on the basis of localization to either the endoplasmic reticulum or

mitochondria. [supplied by OMIM].

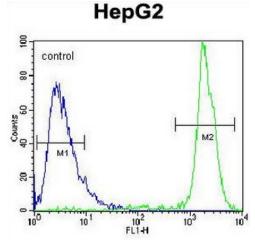
**Synonyms:** Glycerol-3-phosphate acyltransferase 1, KIAA1560

**Protein Pathways:** Glycerolipid metabolism, Glycerophospholipid metabolism, Metabolic pathways

## **Product images:**



GPAM Antibody (Center) western blot analysis in HepG2 cell line lysates (35ug/lane). This demonstrates the GPAM antibody detected the GPAM protein (arrow).



GPAM Antibody (Center) flow cytometric analysis of HepG2 cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.