

Anti-PLA2G4A (pS505) Antibody

Rabbit polyclonal antibody to PLA2G4A (pS505) Catalog # AP60038

Product Information

Application WB, IF/IC, IHC

Primary Accession P47712
Other Accession P47713

Reactivity Human, Mouse, Rat, Monkey

Host Rabbit
Clonality Polyclonal
Calculated MW 85239

Additional Information

Gene ID 5321

Other Names CPLA2; PLA2G4; Cytosolic phospholipase A2; cPLA2; Phospholipase A2 group

IVA

Target/Specificity Recognizes endogenous levels of PLA2G4A (pS505) protein.

Dilution WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 - 1/500)

IF/IC~~N/A IHC~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 -

1/500)

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 PLA2G4A

Synonyms CPLA2, PLA2G4

Function Has primarily calcium-dependent phospholipase and lysophospholipase

activities, with a major role in membrane lipid remodeling and biosynthesis of

lipid mediators of the inflammatory response (PubMed: 10358058,

PubMed: 14709560, PubMed: 16617059, PubMed: 17472963,

PubMed:<u>18451993</u>, PubMed:<u>27642067</u>, PubMed:<u>7794891</u>, PubMed:<u>8619991</u>, PubMed:<u>8702602</u>, PubMed:<u>9425121</u>). Plays an important role in embryo implantation and parturition through its ability to trigger prostanoid

production (By similarity). Preferentially hydrolyzes the ester bond of the fatty acyl group attached at sn-2 position of phospholipids (phospholipase A2 activity) (PubMed:10358058, PubMed:17472963, PubMed:18451993,

PubMed:<u>7794891</u>, PubMed:<u>8619991</u>, PubMed:<u>9425121</u>). Selectively hydrolyzes sn-2 arachidonoyl group from membrane phospholipids, providing the precursor for eicosanoid biosynthesis via the cyclooxygenase pathway (PubMed:10358058, PubMed:17472963, PubMed:18451993, PubMed: 7794891, PubMed: 9425121). In an alternative pathway of eicosanoid biosynthesis, hydrolyzes sn-2 fatty acyl chain of eicosanoid lysophopholipids to release free bioactive eicosanoids (PubMed: 27642067). Hydrolyzes the ester bond of the fatty acyl group attached at sn-1 position of phospholipids (phospholipase A1 activity) only if an ether linkage rather than an ester linkage is present at the sn-2 position. This hydrolysis is not stereospecific (PubMed:7794891). Has calcium-independent phospholipase A2 and lysophospholipase activities in the presence of phosphoinositides (PubMed: 12672805). Has O-acyltransferase activity. Catalyzes the transfer of fatty acyl chains from phospholipids to a primary hydroxyl group of glycerol (sn-1 or sn-3), potentially contributing to monoacylglycerol synthesis (PubMed: 7794891).

Cellular Location

Cytoplasm. Golgi apparatus membrane. Nucleus envelope Note=Translocates to intracellular membranes in a calcium-dependent way.

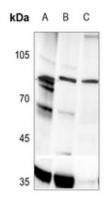
Tissue Location

Expressed in various cells and tissues such as macrophages, neutrophils, fibroblasts and lung endothelium. Expressed in platelets (at protein level) (PubMed:25102815)

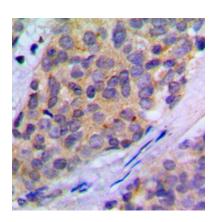
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human PLA2G4A (pS505). The exact sequence is proprietary.

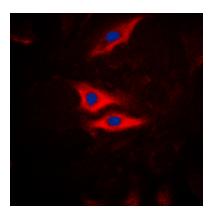
Images



Western blot analysis of PLA2G4A (pS505) expression in Hela (A), mouse brain (B), rat kidney (C) whole cell lysates.



Immunohistochemical analysis of PLA2G4A (pS505) staining in human breast cancer 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 PLA2G4A (pS505) 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 humidified 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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