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anti-PLA2G4E antibody (AA 604-868)

2 Images



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Quantity:	100 μL	
Target:	PLA2G4E	
Binding Specificity:	AA 604-868	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This PLA2G4E antibody is un-conjugated	
Application:	ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)	

Product Details

Immunogen:	Recombinant Human Cytosolic phospholipase A2 epsilon protein (604-868AA)	
Isotype:	IgG	
Cross-Reactivity:	Human	
Purification:	Antigen Affinity Purified	

Target Details

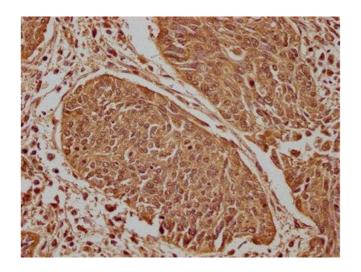
Target:	PLA2G4E	
Alternative Name:	PLA2G4E (PLA2G4E Products)	
Background:	Background: Calcium-dependent phospholipase A2 that selectively hydrolyzes	
	glycerophospholipids in the sn-2 position.	

Target Details

	Aliases: PLA2G4E antibody, Cytosolic phospholipase A2 epsilon antibody, cPLA2-epsilon antibody, EC 3.1.1.4 antibody, Phospholipase A2 group IVE antibody	
UniProt:	Q3MJ16	
Pathways:	Inositol Metabolic Process, VEGF Signaling	

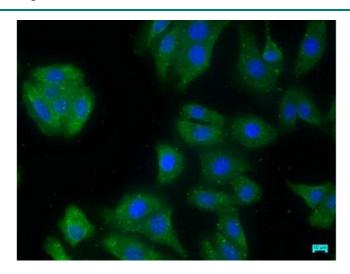
Application Details		
Application Notes:	Recommended dilution: IHC:1:20-1:200, IF:1:50-1:200,	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Buffer:	Preservative: 0.03 % Proclin 300	
	Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4	
Preservative:	ProClin	
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be	
	handled by trained staff only.	
Storage:	-20 °C,-80 °C	
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.	

Images



Immunohistochemistry

Image 1. IHC image of ABIN7149408 diluted at 1:100 and staining in paraffin-embedded human cervical cancer 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 30 min 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.



Immunofluorescence

Image 2. Immunofluorescence staining of HepG2 cells with ABIN7149408 at 1:50, 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).