

# **Product Data Sheet**

## Anti-PLA2G4C Antibody

Catalog #	Source	Reactivity	Applications		
CPA3358	Rabbit	H, M, R	WB, IH		
Description	Ra	abbit polyclonal antibody t	D PLA2G4C		
Immunogen	KL	LH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	re	gion of human PLA2G4C.	he exact sequence is proprietary.		
Purification	Tł	ne antibody was purified b	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	ls of PLA2G4C protein.		
Clonality	Pc	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	PL	LA2G4C			
Alternative Na	ames Cy	ytosolic phospholipase A2	amma; cPLA2-gamma; Phospholipase A2 group IVC		
Entrez Gene	86	605 (Human)			
SwissProt	Q	9UP65 (Human)			
Storage/Stabi	<b>lity</b> Sh	nipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fre	eeze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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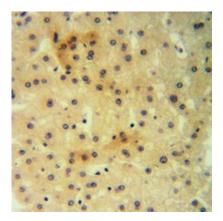
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**kDa** A B C D E 105 70 45 35 For research purposes only, not for human use

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Western blot analysis of PLA2G4C expression in Jurkat (A), H446 (B), mouse lung (C), rat lung (D), rat spleen (E) whole cell lysates. (Predicted band size: 60 kD; Observed band size: 80 kD)



Immunohistochemical analysis of PLA2G4C staining in human liver 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.

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