

## **Product Data Sheet**

### Anti-PALLD Antibody

Catalog #	Source	Reactivity	Applications	
CPA3394	Rabbit	H, M, R	WB, IH	
Description	Rat	bit polyclonal antibody	to PALLD	
Immunogen	KLF	I-conjugated synthetic p	eptide encompassing a sequence within the center	
	reg	ion of human PALLD. The	e exact sequence is proprietary.	
Purification	The	e antibody was purified b	y immunogen affinity chromatography.	
Specificity	Rec	cognizes endogenous lev	els of PALLD protein.	
Clonality	Pol	yclonal		
Conjugation				
Form	Liqu	uid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	and	d 0.01% sodium azide.		
Dilution	WB	8 (1/500 - 1/1000), IH (1/10	00 - 1/200)	
Gene Symbol	PAL	LD		
Alternative Names KIAA0992; Palladin; SIH002; Sarcoma antigen NY-SAR-77		Sarcoma antigen NY-SAR-77		
Entrez Gene	230	23022 (Human); 72333 (Mouse)		
SwissProt	Q81	WX93 (Human); Q9ET54	(Mouse); POC5E3 (Rat)	
Storage/Stabi	<b>lity</b> Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	free	eze/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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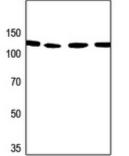
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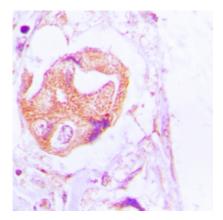
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# **Product Data Sheet**

kDa A B C D



Western blot analysis of PALLD expression in Hela (A), A375 (B), mouse muscle (C), mouse kidney (D) whole cell lysates. (Predicted band size: 150 kD; Observed band size: 122 kD)



Immunohistochemical analysis of PALLD staining in human lung 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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