

## **Product Data Sheet**

## **Anti-NIP7** Antibody

Catalog #	Source	Reactivity	Applications			
CPA3060	Rabbit	H, M, R, B, C, D, Mk, F	P WB, IF/IC			
Description	Ra	bbit polyclonal antibody to	NIP7			
Immunogen	KL	H-conjugated synthetic pep	tide encompassing a sequence within the C-term			
	re	region of human NIP7. The exact sequence is proprietary.				
Purification	Th	e antibody was purified by i	immunogen affinity chromatography.			
Specificity	Re	cognizes endogenous levels	s of NIP7 protein.			
Clonality	Ро	lyclonal				
Conjugation						
Form	Lic	uid in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycer	ol,		
	an	d 0.01% sodium azide.				
Dilution	WI	B (1/500 - 1/1000), IF/IC (1/50	0 - 1/200)			
Gene Symbol	NI	P7				
Alternative Names 60S		S ribosome subunit biogene	ibosome subunit biogenesis protein NIP7 homolog; KD93			
Entrez Gene	51	51388 (Human); 66164 (Mouse); 192180 (Rat)				
SwissProt	QS	Q9Y221 (Human); Q9CXK8 (Mouse); Q9WV50 (Rat)				
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery	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

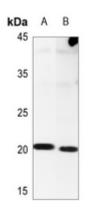
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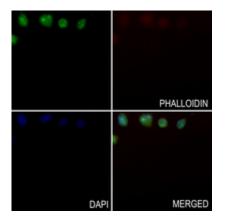
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Western blot analysis of NIP7 expression in A549 (A), DLD (B) whole cell lysates. (Predicted band size: 20 kD; Observed band size: 20 kD)



Immunofluorescent analysis of NIP7 staining in A549 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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