

## **Product Data Sheet**

## Anti-PIR121 Antibody

Catalog #	Source	Reactivity	Applications		
CPA7491	Rabbit	Н, М	WB, IH		
Description	I	Rabbit polyclonal antibody t	o PIR121		
Immunogen	I	KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	ı	region of human PIR121. Th	e exact sequence is proprietary.		
Purification	-	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	els of PIR121 protein.		
Clonality	I	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/2000), IH (1/50	- 1/200)		
Gene Symbol	(	CYFIP2			
Alternative Na	ames l	KIAA1168; PIR121; Cytoplas	nic FMR1-interacting protein 2; p53-inducible protein		
	-	121			
Entrez Gene		26999 (Human); 76884 (Mouse)			
SwissProt	(	Q96F07 (Human); Q5SQX6 (	Mouse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	f	freeze/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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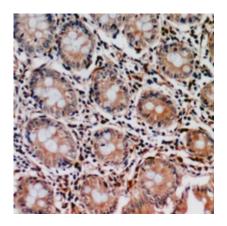
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Western blot analysis of PIR121 expression in HepG2 (A), Hela (B) whole cell lysates. (Predicted band size: 148 kD; Observed band size: 148 kD)



Immunohistochemical analysis of PIR121 staining in human colon 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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