

Product Data Sheet

Anti-p18 INK4c Antibody

Catalog #	Source	Reactivity	Applications			
CPA1213	Rabbit	H, M, R, Mk	WB, IH			
Description		Rabbit polyclonal antibody to p18 INK4c				
Immunogen	I	KLH-conjugated synthetic per	tide encompassing a sequence within the C-term			
	I	region of human p18 INK4c. T	he exact sequence is proprietary.			
Purification	-	The antibody was purified by	immunogen affinity chromatography.			
Specificity	I	Recognizes endogenous level	s of p18 INK4c protein.			
Clonality	I	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	i	and 0.01% sodium azide.				
Dilution		WB (1/500 - 1/1000), IH (1/100	- 1/200)			
Gene Symbol		CDKN2C				
Alternative Na	ames	CDKN6; Cyclin-dependent kin	ase 4 inhibitor C; Cyclin-dependent kinase 6 inhibitor;			
	I	p18-INK4c; p18-INK6				
Entrez Gene		1031 (Human)				
SwissProt	I	P42773 (Human)				
Storage/Stabi	lity	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid			
	1	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

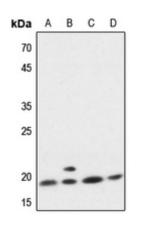
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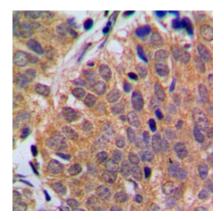


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Western blot analysis of p18 INK4c expression in mouse lung (A), mouse liver (B), rat lung (C), rat liver (D) whole cell lysates. (Predicted band size: 18 kD; Observed band size: 18 kD)



Immunohistochemical analysis of p18 INK4c staining in human breast 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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