

Product Data Sheet

Anti-p27 Kip1 Antibody

Catalog #	Source	Reac	tivity	Applications		
CPA1209	Rabbit	H, R	,	WB, IH		
Description		Rabbit polyclonal antibody to p27 Kip1				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term				
		region of human p27 Kip1. The exact sequence is proprietary.				
Purification		The antibody	was purified by immunogen a	affinity chromatography.		
Specificity		Recognizes endogenous levels of p27 Kip1 protein.				
Clonality		Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glyc				
		and 0.01% so	dium azide.			
Dilution		WB (1/500 - 1/	1000), IH (1/100 - 1/200)			
Gene Symbol		CDKN1B				
Alternative Names		KIP1; Cyclin-dependent kinase inhibitor 1B; Cyclin-dependent kinase inhibitor p27;				
		p27Kip1				
Entrez Gene 1027		1027 (Human))27 (Human)			
SwissProt		P46527 (Huma	an)			
Storage/Stabi	lity	Shipped at 4 $^\circ$	C. Upon delivery aliquot and	d store at -20 $^\circ$ C for one year. Avoid		
		freeze/thaw c	ycles.			

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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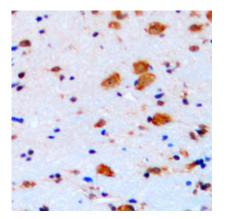
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130

For research purposes only, not for human use

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Western blot analysis of p27 Kip1 expression in MCF7 (A), HEK293T (B), PC12 (C) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 27 kD)



Immunohistochemical analysis of p27 Kip1 staining in human brain 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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