

Product Data Sheet

Anti-p57 Kip2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA9706	Mouse	н	IH		
Description		Mouse monoclonal antibod	y to p57 Kip2		
Immunogen		KLH-conjugated synthetic po	ptide encompassing a sequence within human p57		
		Kip2. The exact sequence is	proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of p57 Kip2 protein.		
Clonality		Monoclonal			
Conjugation					
Form		Mouse IgG3. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium			
		azide.			
Dilution		IH (1/100 - 1/300)			
Gene Symbol		CDKN1C			
Alternative Na	ames	KIP2; Cyclin-dependent kina	se inhibitor 1C; Cyclin-dependent kinase inhibitor p57;		
		р57Кір2			
Entrez Gene		1028 (Human)			
SwissProt		P49918 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
		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

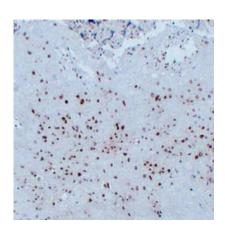
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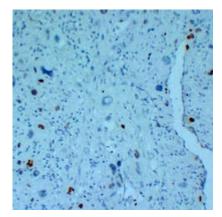
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For research purposes only, not for human use

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Immunohistochemical analysis of p57 Kip2 staining in human placenta 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



Immunohistochemical analysis of p57 Kip2 staining in human partial hydatidiform mole 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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