

# **Product Data Sheet**

### Anti-Caspase 6 p18 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1147	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	Rabbit polyclonal antibody to Caspase 6 p18			
Immunogen	KL	H-conjugated synthetic pe	otide encompassing a sequence within the center		
	re	gion of human Caspase 6 p	18. The exact sequence is proprietary.		
Purification	Th	ne antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	s of Caspase 6 p18 protein.		
Clonality	Pc	olyclonal			
Conjugation					
Form	Lic	quid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/100	) - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	CA	ASP6			
Alternative Na	ames M	CH2; Caspase-6; CASP-6; A	poptotic protease Mch-2		
Entrez Gene	83	39 (Human); 12368 (Mouse	)		
SwissProt	P5	55212 (Human); O08738 (N	louse); O35397 (Rat)		
Storage/Stabi	lity Sh	nipped at 4°C. Upon deliver	y 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

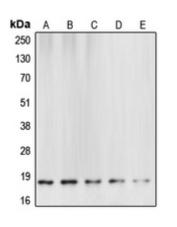
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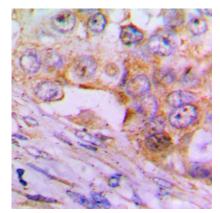




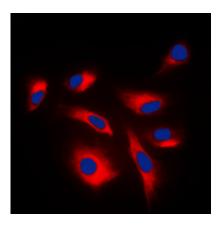
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Western blot analysis of Caspase 6 p18 expression in HeLa (A), mouse brain (B), rat kidney (C), NIH3T3 staurosporine-treated (D), Jurkat etoposide-treated (E) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 33 kD)



Immunohistochemical analysis of Caspase 6 p18 staining in human lung 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.



Immunofluorescent analysis of Caspase 6 p18 staining in NIH3T3 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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