

Product Data Sheet

Anti-MDM2 Antibody

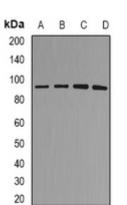
Catalog #	Source	Reactivity	Applications	
CPA7059	Rabbit	H, R, Mk	WB, IH, IF/IC, IP	
Description	F	Rabbit polyclonal antibody to	D MDM2	
Immunogen	ķ	KLH-conjugated synthetic pe	ptide encompassing a sequence within the center	
	r	region of human MDM2. The	e exact sequence is proprietary.	
Purification	Т	The antibody was purified by	r immunogen affinity chromatography.	
Specificity	F	Recognizes endogenous leve	ls of MDM2 protein.	
Clonality	F	Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	a	and 0.01% sodium azide.		
Dilution	١	WB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500), IP (1/10 - 1/100)	
Gene Symbol	Ν	MDM2		
Alternative Na	ames E	E3 ubiquitin-protein ligase N	dm2; Double minute 2 protein; Hdm2; Oncoprotein	
	Ν	Vdm2; p53-binding protein	Mdm2	
Entrez Gene	Z	4193 (Human)		
SwissProt	C	Q00987 (Human)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
	f	reeze/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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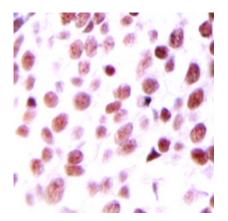




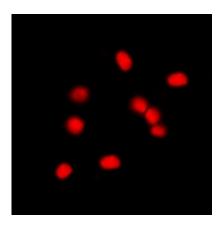
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Western blot analysis of MDM2 expression in SHSY5Y (A), A549 (B), rat brain (C), rat lung (D) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 90 kD)



Immunohistochemical analysis of MDM2 staining in human prostate 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 MDM2 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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