

Product Data Sheet

Recombinant Anti-MOV10 Rabbit mAb

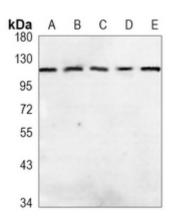
Catalog #	Course	Poortivity	Applications	
Catalog #	Source	Reactivity	Applications	
CMA4342	Rabbit	H, M, R	WB, IF/IC	
Description	Re	ecombinant rabbit monoc	lonal antibody to MOV10	
Immunogen	KL	H-conjugated synthetic p.	eptide encompassing a sequence within human MOV1	0.
	Th	ne exact sequence is prop	rietary.	
Purification	Th	ne antibody was purified b	by immunogen affinity chromatography.	
Specificity	Re	ecognizes endogenous lev	els of MOV10 protein	
Clonality	М	onoclonal		
Conjugation				
Form	Lic	quid in PBS containing 50	% glycerol, 0.2% BSA and 0.01% sodium azide.	
Dilution	W	B (1/500 - 1/1000), IF/IC (1	/50 - 1/200)	
Gene Symbol	М	OV10		
Alternative Na	ames Kl	AA1631; Putative helicase	e MOV-10; Moloney leukemia virus 10 protein	
Entrez Gene	43	843 (Human); 17454 (Μοι	use)	
SwissProt	Q	9HCE1 (Human); P23249	(Mouse)	
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ery 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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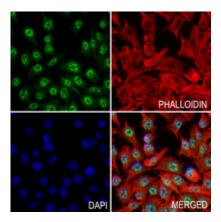




For research purposes only, not for human use

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Western blot analysis of MOV10 expression in HepG2 (A), MCF7 (B), mouse liver (C), mouse kidney (D), rat liver (E) whole cell lysates. (Predicted band size: 113 kD; Observed band size: 114 kD)



Immunofluorescent analysis of MOV10 staining in A375 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 an AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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