

Product Data Sheet

Recombinant Anti-hnRNP E2 Rabbit mAb

Catalog #	Source	Reactivity	Applications
CMA4572	Rabbit	н	WB, IH, IF/IC
Description	F	Recombinant rabbit mono	clonal antibody to hnRNP E2
Immunogen	k	KLH-conjugated synthetic p	eptide encompassing a sequence within human hnRNP
	E	E2 protein. The exact sequ	ence is proprietary.
Purification	T	The antibody was purified	by immunogen affinity chromatography.
Specificity	F	Recognizes endogenous lev	els of hnRNP E2 protein
Clonality	Ν	Monoclonal	
Conjugation			
Form	L	iquid in PBS containing 50-	% glycerol, 0.2% BSA and 0.01% sodium azide.
Dilution	١	WB (1/500 - 1/1000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	F	PCBP2	
Alternative N	ames F	Poly(rC)-binding protein 2;	Alpha-CP2; Heterogeneous nuclear ribonucleoprotein E2;
	ł	nnRNP E2	
Entrez Gene 5094 (Human)		5094 (Human)	
SwissProt	C	Q15366 (Human)	
Storage/Stabi	i lity S	Shipped at 4 $^\circ$ C. Upon del	very aliquot and store at -20 $^\circ$ 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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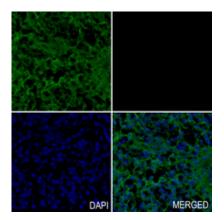
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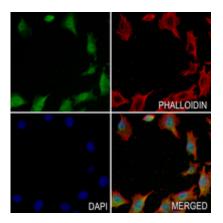
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Western blot analysis of hnRNP E2 expression in MCF7 (A), K562 (B), Hela (C) whole cell lysates. (Predicted band size: 38, 35 kD; Observed band size: 38, 35 kD)



Immunohistochemical analysis of hnRNP E2 staining in human tonsil 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 hnRNP E2 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 hidified chamber. Cells were washed with PBST and incubated with a 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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