

Product Data Sheet

Anti-hnRNP L Antibody

Catalog #	Source	Reactivity	Applications		
CPA7066	Rabbit	Н, М	WB, IH, IF/IC		
Description	Ra	Rabbit polyclonal antibody to hnRNP L			
Immunogen	KI	LH-conjugated synthetic p	eptide encompassing a sequence within the center		
	re	egion of human hnRNP L.	The exact sequence is proprietary.		
Purification	Tł	he antibody was purified k	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of hnRNP L protein.		
Clonality	Po	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	VB (1⁄500 - 1⁄1000), IH (1⁄1	00 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	н	NRNPL			
Alternative Na	ames H	NRPL; Heterogeneous nuc	lear ribonucleoprotein L; hnRNP L		
Entrez Gene	32	191 (Human); 15388 (Mou	ise)		
SwissProt	P:	14866 (Human); Q8R081 (Mouse)		
Storage/Stabi	lity Sł	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fr	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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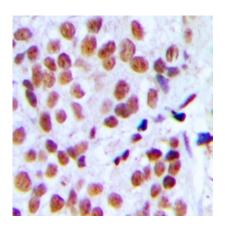
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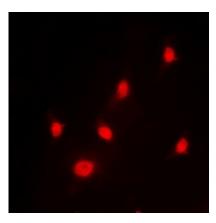
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Western blot analysis of hnRNP L expression in HuvEc (A), HepG2 (B), NIH3T3 (C) whole cell lysates. (Predicted band size: 64 kD; Observed band size: 68 kD)



Immunohistochemical analysis of hnRNP L staining in human breast 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 hnRNP L staining in HeLa 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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