

Anti-hNRNP C (Phospho-S260) Antibody

Catalog #	Source	Reactivity	Applications
CPA7008	Rabbit	H, M, R, Rb	WB, IH
Description	Rabbit polyclonal antibody to hNRNP C (Phospho-S260)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S260 of human hNRNP C protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of hNRNP C protein only when phosphorylated at S260.		
Clonality	Polyclonal		
Conjugation			
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
Dilution	WB (1/500 - 1/1000), IH (1/100 - 1/200)		
Gene Symbol	HNRNPC		
Alternative Names	HNRPC; Heterogeneous nuclear ribonucleoproteins C1/C2; hnRNP C1/C2		
Entrez Gene	3183 (Human); 15381 (Mouse)		
SwissProt	P07910 (Human); Q9Z204 (Mouse); G3V9R8 (Rat)		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/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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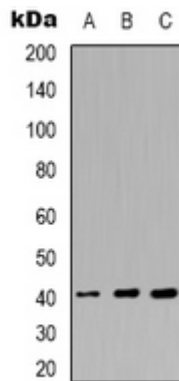
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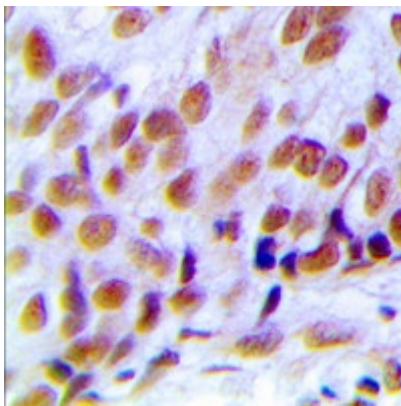
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Product Data Sheet



Western blot analysis of hNRNP C (Phospho-S260) expression in HeLa (A), K562 (B), HEK293T (C) whole cell lysates.
(Predicted band size: 33 kD; Observed band size: 40 kD)



Immunohistochemical analysis of hNRNP C (Phospho-S260) 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.

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