

hnRNP C1/C2 Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H Mk	Sensitivity: Endogenous	MW (kDa): 40	Source/Isotype: Rabbit	UniProt ID: #P07910	Entrez-Gene Id: 3183	
Product Usage Information		Application Western Blotting			Dilution 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity/Sensitivity		hnRNP C1/C2 Antibody recognizes endogenous levels of total hnRNP C1/C2 protein. This antibody does not cross-react with other hnRNPCL1 or RALY proteins.					
Species predicted to react based on 100% sequence homology		Hamster, Xenopus, Bovine, Dog, Pig, Horse, Guinea Pig					
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human hnRNP C1/C2 protein. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		Heterogeneous nuclear ribonucleoprotein C1/C2 (hnRNP C1/C2) has multiple biological functions including transcriptional regulation, DNA repair, and RNA processing. hnRNP C1/C2 acts as a 'molecular ruler' in the mRNA processing pathway, committing nascent transcripts from the chromatin template to the mRNA export pathway once the nascent transcript becomes longer than 200-300 nucleotides (1). hnRNP C1/C2 associates with SWI/SNF and NurD family members to form the locus control region (LCR)-associated remodeling complex (LARC), which binds to β-globin gene promoter to prevent transcriptional silencing. Studies indicate that without hnRNP C1/C2, LARC does not associate with its target DNA sequence (2,3). hnRNP C1/C2 and other hnRNP family members interact with DNA damage response (DDR) proteins (4). hnRNP proteins regulate double stranded break (DSB) repair by promoting either homologous recombination (HR) or non-homologous end joining (NHEJ) (4). hnRNP C1/C2 downregulates the expression of miR-21, which leads to the increased expression of programmed cell death 4 (PDCD4) protein in glioblastoma multiforme (GBM) (5). Research studies have shown that silencing of hnRNP C1/C2 renders GBM cells more susceptible to apoptosis (5).					
Background Re	ferences	2. Huang, L. et al. (201 3. Mahajan, M.C. et al. 4. Haley, B. et al. (2009	1. McCloskey, A. et al. (2012) <i>Science</i> 335, 1643-6. 2. Huang, L. et al. (2011) <i>Mol Cell Biol</i> 31, 3472-84. 3. Mahajan, M.C. et al. (2005) <i>Proc Natl Acad Sci U S A</i> 102, 15012-7. 4. Haley, B. et al. (2009) <i>Int J Radiat Biol</i> 85, 643-55. 5. Park, Y.M. et al. (2012) <i>Mol Cell Biol</i> 32, 4237-44.				
Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot)						western blot).	

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting

Cross-Reactivity Key H: Human Mk: Monkey

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