Store at -20C

056



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

Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 100	Source/Isotype: Rabbit	UniProt ID: #Q08J23	Entrez-Gene Id: 54888
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		NSUN2 Antibody recognizes endogenous levels of total NSUN2 protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val690 of human NSUN2 protein. Antibodies are purified by peptide affinity chromatography.				
Background		Chemical modifications of RNA regulate many cellular processes. One particular RNA modification, 5- methylcytosine (5-mC), regulates ribosome assembly, translation, and RNA stability (1). In eukaryotes, this modification is added to RNA by the DNA methyltransferase homologue 2 protein (DNMT2; also known as TRDMT1), and also by members of the NOL1/NOP2/SUN domain (NSUN) family of proteins. NSUN proteins are putative S-adenosylmethionine (SAM)-dependent methyltransferases that carry out their enzymatic activity by utilizing two cysteine residues in their active sites (2). There are currently seven known members of this family, consisting of NOP2 (NSUN1) and NSUN2-7.				
		NSUN2 is an 86 kDa me methylates a variety of methylation of pre-tRN/ methylate mRNAs and s reported in multiple typ among others (9). The n phosphorylation at Serf diagnostic marker for A	tRNAs and other F A ^{Leu} , tRNA ^{Gly} , tRNA several ncRNAs, su ses of human canc nethyltransferase I 39 (10), and there	NA substrates (3). NSUN V ^{al} , tRNA ^{Leu} , and tRNA ^{A:} Ich as vtRNAs (6-8). Eleva ers, including breast, pr activity of NSUN2 is red is potential for NSUN2	N2 has been found ^{sp} (1,4,5). It has also ated NSUN2 proteir ostate, kidney, blad uced upon Aurora B	to play a role in the been found to hlevels have been lder, and liver, 3 kinase-mediated
Background Re	eferences	1. Trixl, L. and Lusser, A. 2. Bohnsack, K.E. et al. (3. Bourgeois, G. et al. (2 4. Brzezicha, B. et al. (20 5. Tuorto, F. et al. (2012) 6. Tang, H. et al. (2015) 7. Xing, J. et al. (2015) M 8. Hussain, S. et al. (201 9. Okamoto, M. et al. (20 10. Sakita-Suto, S. et al.	2019) <i>Genes (Base 015) PLoS One 10 006) Nucleic Acids) Nat Struct Mol Bi Aging (Albany NY) 10 Cell Biol 35, 404 3) Cell Rep 4, 255- 012) DNA Cell Biol</i>	e/) 10, pii: E102. doi: 10.3 e0133321. <i>Res</i> 34, 6034-43. o/ 19, 900-5. 7, 1143-58. I3-52. 61. 31, 660-71.		2.
Species Reactiv	/ity	Species reactivity is dete	ermined by testing	j in at least one approve	ed application (e.g.,	western blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human M: Mouse R: Rat Mk: Monkey				
Trademarks and Patents		Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.				

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