

Applications: W, IP	<b>Reactivity:</b> H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 60	Source/Isotype: Rabbit	<b>UniProt ID:</b> #O94992	<b>Entrez-Gene Id:</b> 10614
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 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. Do not aliquot the antibody.				
Specificity/Sensitivity		HEXIM1 Antibody recognizes endogenous levels of total HEXIM1 protein.				
Species predicted to react based on 100% sequence homology		Bovine, Dog, Horse				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human HEXIM1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Hexamethylene bis-acetamide-inducible protein 1 (HEXIM1) was originally identified in vascular smooth muscle cells as a protein that is upregulated upon treatment with the differentiating agent hexamethylene bisacetamide (1). HEXIM1 binds 7SK RNA, a highly abundant non-coding RNA, and together they act as a potent inhibitor of positive transcription elongation factor b (P-TEFb) (2,3). P-TEFb phosphorylates the C-terminal domain of the largest subunit of RNA polymerase II and is an important regulator of transcription elongation (4-8). 7SK RNA-bound HEXIM1 interacts with the cyclin T1 subunit of P-TEFb, sequestering P-TEFb in an inactive form leading to transcription inhibition (2,3). The regulation of the relative ratio of inactive to active P-TEFb in the cell by HEXIM1/7SK RNA is thought to play a critical role in regulation of a wide range of cellular gene expression programs such as estrogen and glucocorticoid receptor regulated genes (9-12).				
Background References		<ol> <li>Ouchida, R. et al. (2003) <i>Genes Cells</i> 8, 95-107.</li> <li>Michels, A.A. et al. (2004) <i>EMBO J</i> 23, 2608-19.</li> <li>Yik, J.H. et al. (2003) <i>Mol Cell</i> 12, 971-82.</li> <li>Buratowski, S. (2009) <i>Mol Cell</i> 36, 541-6.</li> <li>Lenasi, T. and Barboric, M. <i>RNA Biol</i> 7, 145-50.</li> <li>Pirngruber, J. et al. (2009) <i>Cell Cycle</i> 8, 3636-42.</li> <li>Wada, T. et al. (1998) <i>EMBO J</i> 17, 7395-403.</li> <li>Yamada, T. et al. (2006) <i>Mol Cell</i> 21, 227-37.</li> <li>Peterlin, B.M. et al. (2012) <i>Wiley Interdiscip Rev RNA</i> 3, 92-103.</li> <li>Ketchart, W. et al. (2011) <i>Oncogene</i> 30, 3563-9.</li> <li>Ogba, N. et al. (2008) <i>Cancer Res</i> 68, 7015-24.</li> <li>Shimizu, N. et al. (2005) <i>Proc Natl Acad Sci U S A</i> 102, 8555-60.</li> </ol>				
Species Reactiv	/ity	Species reactivity is det	ermined by testing	g 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 BSA, 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 Mk: Monkey				
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