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CBX4 Antibody -20C l268



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Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 78	Source/Isotype: Rabbit	UniProt ID: #000257	Entrez-Gene Id: 8535		
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:100			
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/Ser	sitivity	CBX4 Antibody recognizes endogenous levels of total CBX4 protein. In Western blot analysis, the antibody detects 32kDa and 55kDa proteins of unknown identity.						
Source / Purifi	ource / Purification Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresp residues surrounding Pro166 of human CBX4 protein. Antibodies are purified by protein A a affinity chromatography.							
Background Background R	eferences	The polycomb group (PcG) proteins contribute to the maintenance of cell identity, stem cell self- renewal, cell cycle regulation, and oncogenesis by maintaining the silenced state of genes that promote cell lineage specification, cell death, and cell-cycle arrest (1-4). PcG proteins exist in two complexes that cooperate to maintain long-term gene silencing through epigenetic chromatin modifications. The first complex, EED-EZH2, is recruited to genes by DNA-binding transcription factors and methylates histone H3 on Lys27. This histone methyl-transferase activity requires the Ezh2, Eed, and Suz12 subunits of the complex (5). Histone H3 methylation at Lys27 facilitates the recruitment of the second complex, PRC1, which ubiquitinylates histone H2A on Lys119 (6). CBX4 is a component of the PRC1 complex, which together with Ring1 strongly enhances the E3 ubiquitin ligase activity of the Ring2 catalytic subunit (7,8). CBX4 itself is a SUMO E3 ligase, and its function influences EMT, DNA damage response, tumor angiogenesis, and self-renewal (9-13). 1. Boyer, L.A. et al. (2006) <i>Nature</i> 441, 349-53. 2. Lee, T.I. et al. (2006) <i>Nature</i> 441, 349-53. 3. Cao, R. et al. (2002) <i>Science</i> 298, 1039-43. 4. Müller, J. et al. (2002) <i>Science</i> 298, 1039-43. 5. Cao, R. and Zhang, Y. (2004) <i>Mol Cell</i> 15, 57-67. 6. Wang, H. et al. (2004) <i>Nature</i> 431, 873-8. 7. Satijn, D.P. et al. (1997) <i>Mol Cell</i> Biol 17, 6076-86. 8. Cao, R. et al. (2003) <i>Mol Cell</i> 20, 845-54. 9. Kagey, M.H. et al. (2007) <i>Biochem J</i> 405, 369-78. 11. Ismail, I.H. et al. (2007) <i>Biochem J</i> 405, 369-78.						
		12. Li, J. et al. (2014) <i>Ca</i> 13. Klauke, K. et al. (20						
Species Reacti	vity	Species reactivity is de	termined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	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 K	ey	W: Western Blotting IP: Immunoprecipitation						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat Mk: Monkey						
Trademarks ar	nd Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.						

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