Revision 4	
DUSP4/MKP2 (D9A5) Rabbit mAb	
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Applications: W, W-S	Reactivity: H Mk	Sensitivity: Endogenous	MW (kDa): 42	Source/Isotype: Rabbit	UniProt ID: #Q13115	Entrez-Gene Id: 1846	
Product Usage Information	2	Application Western Blotting Simple Western™		1:1	l ution 000 50 - 1:250		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliguot the antibody.					
Specificity/Sensitivity		DUSP4/MKP2 (D9A5) Rabbit mAb recognizes endogenous levels of total DUSP4 protein.					
Source / Purifi	cation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro168 of human DUSP4 protein.					
Background		MAP kinases are inactivated by dual-specificity protein phosphatases (DUSPs) that differ in their substrate specificity, tissue distribution, inducibility by extracellular stimuli, and cellular localizati DUSPs, also known as MAPK phosphatases (MKPs), specifically dephosphorylate both threonine a tyrosine residues in MAPK P-loops and have been shown to play important roles in regulating the function of the MAPK family (1,2). At least 13 members of the family (DUSP1-10, DUSP14, DUSP12) display unique substrate specificities for various MAP kinases (3). MAPK phosphatases typically contain an amino-terminal rhodanese-fold responsible for DUSP docking to MAPK family members and a carboxy-terminal catalytic domain (4). These phosphatases can play important role specificities for surious specification of the response of cancer to chemotherapy (6).				ular localization. h threonine and egulating the SP14, DUSP16, and osphatases MAPK family important roles in sis (5). In addition,	
		DUSP4 (MKP2, hVH2) is a nuclear dual-specificity phosphatase that is a negative regulator of Erk1/2 signaling by dephosphorylating and inactivating Erk1/2 in response to growth factors (7,8). Treatment with mitogen or expression of activating mutations of Ras (G12V) or Raf (V600E) promote increased expression of DUSP4 and a coincident decrease in phospho-Erk in the nucleus (9). In contrast, numerous studies have detected decreased expression of DUSP4 in a variety of tumor types, resulting in increased signaling via the Ras/Erk pathway, enhanced tumor growth, and decreased drug sensitivity (10-12). DUSP4/MKP2 also plays an important role in regulating the immune system where it has been implicated in regulating T and B cell proliferation and apoptosis, and adaptive and inflammatory responses (13-16).					
Background R	eferences	1. Camps, M. et al. (20 2. Theodosiou, A. and 3. Salojin, K. and Orav 4. Tanoue, T. et al. (20 5. Dickinson, R.J. and I 6. Wu, G.S. (2007) <i>Can</i> 7. Peng, D.J. et al. (201 8. Lawan, A. et al. (201 9. Cagnol, S. and Riva 10. Chitale, D. et al. (20 12. Balko, J.M. et al. (20 12. Balko, J.M. et al. (20 13. Ramesh, S. et al. (2 14. Cornell, T.T. et al. (15. Huang, C.Y. et al. (20 16. Yu, M. et al. (2012)	Ashworth, A. (2002 ecz, T. (2007) J Leul 02) J Biol Chem 277 Keyse, S.M. (2006) J cer Metastasis Rev 0) Cell Cycle 9, 465 (1) J Biol Chem 286, rd, N. (2012) Oncog 009) Oncogene 28, 10) Cancer Res 70, 012) Nat Med 18, 1 2008) EMBO Rep 9, 2010) Infect Immun 2012) Eur J Immun	 Cenome Biol 3, REVIEV (coc Biol 81, 860-9. , 22942-9. Cell Sci 119, 4607-15. 26, 579-85. 0-5. 12933-43. tene,. 2773-83. 1689-99. 052-9. 990-7. n 78, 2868-76. b) 42, 476-88. 	v53009.		
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).	
Western Blot E	Buffer	IMPORTANT: For west TBS, 0.1% Tween® 20		membrane with diluted shaking, overnight.	primary antibody i	n 5% w/v BSA, 1X	

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Applications Key	W: Western Blotting W-S: Simple Western™				
Cross-Reactivity Key	H: Human Mk: Monkey				
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