## ម្ចុ ស្តុ MMP-13 (E4W3T) Rabbit mAb





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Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 60	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P45452	Entrez-Gene Id: 4322		
Product Usage Information	2	<b>Application</b> Western Blotting			Dilution 1:1000			
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. <i>Do not aliquot the antibody.</i>						
Specificity/Sen	sitivity	MMP-13 (E4W3T) Rabbit mAb recognizes endogenous levels of total human MMP13 protein.						
<b>Source / Purification</b> Monoclonal antibody is produced by immunizing animals with a recombinant protein carboxy terminus of human MMP13 protein.				n specific to the				
Background		MMP-13 (collagenase 3) belongs to the matrix metalloproteinase (MMP) superfamily of enzymes that targets many extracellular proteins, including other proteases, growth factors, cell surface receptors, and adhesion molecules (1,2). MMP-13 is a member of a subgroup of collagenases (including MMP-1, MMP-8, and MMP-18) that play an even more important function targeting fibrillar collagen. MMP-13 is synthesized as a latent proenzyme, and proteolytic removal of the inhibitory propeptide domain is required for enzyme activation. MMP-13 protein levels are regulated at the transcriptional level via specific transcription factors and promoter DNA methylation (3,4). MMP-13 preferentially cleaves Type II collagen, and research studies have shown that aberrant upregulation of MMP-13 activity can lead to cartilage loss and osteoarthritis (5,6). In addition, MMP-13 has been shown to promote cancer development, in part through enhancing tumor angiogenesis and metastases (7-9), suggesting that collagenase activity may serve as a useful marker of tumor progression (10).						
Background R	eferences	<ol> <li>McCawley, L.J. and Matrisian, L.M. (2001) <i>Curr Opin Cell Biol</i> 13, 534-40.</li> <li>Cathcart, J. et al. (2015) <i>Genes Dis</i> 2, 26-34.</li> <li>Chan, C.M. et al. (2017) <i>J Biol Chem</i> 292, 1625-36.</li> <li>Hashimoto, K. et al. (2013) <i>J Biol Chem</i> 288, 10061-72.</li> <li>Elayyan, J. et al. (2017) <i>FASEB J</i> 31, 3116-25.</li> <li>Wang, M. et al. (2011) <i>Ann N Y Acad Sci</i> 1240, 61-9.</li> <li>Mendonsa, A.M. et al. (2015) <i>Mol Cancer</i> 14, 49.</li> <li>Wang, C. et al. (2015) <i>Oncotarget</i> 6, 2903-16.</li> <li>Kudo, Y. et al. (2012) <i>J Biol Chem</i> 287, 38716-28.</li> <li>Brinckerhoff, C.E. et al. (2000) <i>Clin Cancer Res</i> 6, 4823-30.</li> </ol>						
Species Reacti	vity	Species reactivity is determined by testing in at least one approved 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						
Cross-Reactivit	ty Key	H: Human						
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