c-Myc (D84C12) Rabbit mAb



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Applications: W, W-S, IF-IC	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 57-65	Source/Isotype: Rabbit IgG	UniProt ID: #P01106	Entrez-Gene Id: 4609	
Product Usage Information		Application Western Blotting Simple Western™ Immunofluorescence (Immunocytochemistry)		Dilution 1:1000 1:50 - 1:250 1:400 - 1:1600			
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 aliquot the antibody.					
Specificity/Ser	nsitivity	c-Myc (D84C12) Rabbit mAb detects endogenous levels of total c-Myc protein. This antibody is not recommended for detection of Myc-tagged fusion proteins (use Cell Signaling Technology cat. #2276 or #2278).					
Species predic based on 100% homology	ted to react 6 sequence	Dog, Pig					
Source / Purifi	cation	Monoclonal antibody is produced with a synthetic peptide corresponding to residues surrounding Asp30 of human c-Myc1 protein.					
Background		Members of the Myc/Max/Mad network function as transcriptional regulators with roles in various aspects of cell behavior, including proliferation, differentiation, and apoptosis (1). These proteins share a common basic-helix-loop-helix leucine zipper (bHLH-ZIP) motif required for dimerization and DNA- binding. Max was originally discovered based on its ability to associate with c-Myc and found to be required for the ability of Myc to bind DNA and activate transcription (2). Subsequently, Max has been viewed as a central component of the transcriptional network, forming homodimers as well as heterodimers with other members of the Myc and Mad families (1). The association between Max and either Myc or Mad can have opposing effects on transcriptional regulation and cell behavior (1). The Mad family consists of four related proteins; Mad1, Mad2 (Mxi1), Mad3, and Mad4, and the more distantly related members of the bHLH-ZIP family, Mnt and Mga. Like Myc, the Mad proteins are tightly regulated with short half-lives. In general, Mad family members interfere with Myc-mediated processes, such as proliferation, transformation, and prevention of apoptosis by inhibiting transcription (3,4).					
Background R	eferences	1. Baudino, T.A. and Cleveland, J.L. (2001) <i>Mol Cell Biol</i> 21, 691-702. 2. Blackwood, E.M. and Eisenman, R.N. (1991) <i>Science</i> 251, 1211-7. 3. Henriksson, M. and Lüscher, B. (1996) <i>Adv Cancer Res</i> 68, 109-82. 4. Grandori, C. et al. (2000) <i>Annu Rev Cell Dev Biol</i> 16, 653-99.					
Species Reacti	vity	Species reactivity is do	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).	
Western Blot B	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 K	ey	W: Western Blotting W-S: Simple Western™ IF-IC: Immunofluorescence (Immunocytochemistry)					
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat					
Trademarks a	nd Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.					
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