## <sup>502- Top 207</sup> β-Actin (8H10D10) Mouse mAb



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Applications: Reactivity: W, W-S, IHC-P, IF-IC, H M R Hm Mk Dg FC-FP	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 45	<b>Source/Isotype:</b> Mouse IgG2b	<b>UniProt ID:</b> #P60709	<b>Entrez-Gene Id:</b> 60
Product Usage Information	<b>Application</b> Western Blotting Simple Western™ Immunohistochemistry (Paraffin) Immunofluorescence (Immunocytochemistry) Flow Cytometry (Fixed/Permeabilized)			<b>Dilution</b> 1:1000 1:10 - 1:50 1:3000 - 1:12000 1:400 - 1:1600 1:800 - 1:3200	
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/Sensitivity	β-Actin (8H10D10) Mouse mAb detects endogenous levels of total β-actin protein. Due to the high sequence identity between the cytoplasmic actin isoforms, β-actin and cytoplasmic γ-actin, this antibody may cross-react with cytoplasmic γ-actin. It does not cross-react with α-skeletal, α-cardiac, α-vascular smooth, or γ-enteric smooth muscle isoforms.				
Source / Purification	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to amino-terminal residues of human β-actin.			orresponding to	
Background	Actin, a ubiquitous eukaryotic protein, is the major component of the cytoskeleton. At least six isoforms are known in mammals. Nonmuscle $\beta$ - and $\gamma$ -actin, also known as cytoplasmic actin, are ubiquitously expressed, controlling cell structure and motility (1). While all actin isoforms are highly homologous, cytoplasmic $\beta$ - and $\gamma$ -actin protein sequences differ by only four biochemically similar amino acids (2). For this reason, antibodies raised to $\beta$ -actin may cross-react with $\gamma$ -actin, and vice versa. $\alpha$ -cardiac and $\alpha$ -skeletal actin are expressed in striated cardiac and skeletal muscles, respectively; two smooth muscle, respectively. These actin isoforms regulate the contractile potential of muscle cells (1). Actin exists mainly as a fibrous polymer, F-actin. In response to cytoskeletal reorganizing signals during processes such as cytokinesis, endocytosis, or stress, cofilin promotes fragmentation and depolymerization of F-actin, resulting in an increase in the monomeric globular form, G-actin (3). The ARP2/3 complex stabilizes F-actin fragments and promotes formation of new actin filaments (3). Research studies have shown that actin is hyperphosphorylated in primary breast tumors (4). Cleavage of actin under apoptotic conditions has been observed <i>in vitro</i> and in cardiac and skeletal muscle, as shown in research studies (5-7). Actin cleavage by caspase-3 may accelerate ubiquitin/proteasome-dependent muscle proteolysis (7).				
Background References	<ol> <li>Herman, I.M. (1993) <i>Curr. Opin. Cell Biol.</i> 5, 48-55.</li> <li>Perrin, B.J. and Ervasti, J.M. (2010) <i>Cytoskeleton (Hoboken)</i> 67, 630-4.</li> <li>Condeelis, J. (2001) <i>Trends Cell Biol</i> 11, 288-93.</li> <li>Lim, Y.P. et al. (2004) <i>Clin Cancer Res</i> 10, 3980-7.</li> <li>Kayalar, C. et al. (1996) <i>Proc Natl Acad Sci U S A</i> 93, 2234-8.</li> <li>Communal, C. et al. (2002) <i>Proc Natl Acad Sci U S A</i> 99, 6252-6.</li> <li>Du, J. et al. (2004) <i>J Clin Invest</i> 113, 115-23.</li> </ol>				
Species Reactivity	Species reactivity is c	letermined by testin	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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key	W: Western Blotting W-S: Simple Western™ IHC-P: Immunohistochemistry (Paraffin) IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)				

Cross-Reactivity Key	H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey Dg: Dog		
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