Revision 1

Store at -20C

#4968



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Applications: W, IHC-P	Reactivity: H M R Mk Z	Sensitivity: Endogenous	MW (kDa): 45	Source/Isotype: Rabbit	UniProt ID: #P60709, #P68133, #P63261, #P68032, #P62736, #P63267	Entrez-Gene Id: 60, 58, 71, 70, 59, 72
Product Usage Information	1	Application Western Blotting Immunohistochemistr	y (Paraffin)		Dilution 1:1000 1:50 - 1:2	
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/Sensitivity		Pan-Actin Antibody detects endogenous levels of total actin (all isoforms). The antibody also detects the 30 kDa actin fragment cleaved at glutamate 107.				
Species predicted to react based on 100% sequence homology		D. melanogaster, Xenopus, Bovine, Pig				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp244 of human beta-actin. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Actin, a ubiquitous eukaryotic protein, is the major component of the cytoskeleton. At least six isoforms are known in mammals. Nonmuscle β - and γ -actin, also known as cytoplasmic actin, are ubiquitously expressed, controlling cell structure and motility (1). While all actin isoforms are highly homologous, cytoplasmic β - and γ -actin protein sequences differ by only four biochemically similar amino acids (2). For this reason, antibodies raised to β -actin may cross-react with γ -actin, and vice versa. α -cardiac and α -skeletal actin are expressed in striated cardiac and skeletal muscles, respectively; two smooth muscle actins, α - and γ -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 Re	eferences	1. Herman, I.M. (1993) 2. Perrin, B.J. and Erva: 3. Condeelis, J. (2001) 4. Lim, Y.P. et al. (2004) 5. Kayalar, C. et al. (199 6. Communal, C. et al. 7. Du, J. et al. (2004) <i>J</i> C	sti, J.M. (2010) <i>Cyto</i> . Trends Cell Biol 11, <i>Clin Cancer Res</i> 10 96) <i>Proc Natl Acad S</i> (2002) <i>Proc Natl Ac</i>	skeleton (Hoboken) 67 288-93. , 3980-7. cci U S A 93, 2234-8. ad Sci U S A 99, 6252-6		
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot 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 Key		W: Western Blotting IHC-P: Immunohistochemistry (Paraffin)				

Cross-Reactivity Key	H: Human M: Mouse R: Rat Mk: Monkey Z: Zebrafish				
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