2010 AMPKβ1/2 (57C12) Rabbit mAb 010



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Applications: W, IHC-P, IF-IC, FC- FP	Reactivity: H M R Hm Mk	Sensitivity: Endogenous	MW (kDa): 30, 38	Source/Isotype: Rabbit IgG	UniProt ID: #Q9Y478, #O43741	Entrez-Gene Id: 5564, 5565
Product Usage Information		Application Western Blotting Immunohistochemis Immunofluorescence Flow Cytometry (Fixe	(Immunocytochem	istry)		Dilution 1:1000 1:100 1:50 1:100
Storage), 150 mM NaCl, 100 μ ot aliquot the antibody	g/ml BSA, 50% glycerc y.	l and less than
Specificity/Sensitivity		AMPKbeta1/2 (57C12) Rabbit mAb detects endogenous levels of both total AMPKβ1 and β2 proteins. The antibody does not cross-react with other related proteins.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the sequence surrounding His233 residues of human $AMPK\beta1$.				
Background		key role in the regula catalytic α subunit an genes (α 1, 2; β 1, 2; γ' environmental stress association with acce activation loop, and t phosphorylated at Tr significance of these translationally modifi Ser101, Ser108, and S AMPK activation, whi mutations in AMPK γ binding sites (CBS or and cause glycogen a that AMPK not only r synthesis and cell gro (1). AMPKbeta1 and A contribute equally to shows minimal expres	tion of energy home of regulatory β and γ 1, 2, 3) (2). The kinase , such as heat shock ssory proteins STRA his phosphorylation ar258 and Ser485 (fo phosphorylation events and Ser485 (fo phosphorylation events and Ser485 (fo phosphorylation ar subunits have been Bateman domains). accumulation in hear egulates the metabor with through EF2 an MPK beta2 share ap AMPK activity. AMPH ssion in kidney and	ostasis (1). AMPK is a subunits, each of white is activated by an elet , hypoxia, and ischemi D and MO25, phospho is required for AMPK at r α1; Ser491 for α2). The the start of the second orylation at Ser108 of orylation at Ser108 of the Ser24/25 and Ser182 identified, most of white Mutations at these site t or skeletal muscle (1 lism of fatty acids and d TSC2/mTOR pathwa proximately 70% sequits (β1 is predominantly effective second seco	yeast to plants and ani heterotrimeric comple- ich is encoded by two of vated AMP/ATP ratio d ia (1). The tumor suppri- orylates AMPK α at Thr1 activation (3-5). AMPK α he upstream kinase an cidated (6). The β 1 sub- horylation including S6 the β 1 subunit seems 2 affects AMPK localiza- ich are located in the p es lead to reduction of ,2). Accumulating evid I glycogen, but also mo ys, as well as blood flo ence homology. Both i expressed in the liver a nparison, AMPK β 2 is h nd lung.	x composed of a or three distinct ue to cellular and ressor LKB1, in 72 in the a is also d the biological ount is post- er24/25, Ser96, to be required for tion (7). Several outative AMP/ATP AMPK activity ence indicates odulates protein w via eNOS/nNOS soforms nd brain, and
Background Re	eferences	1. Hardie, D.G. (2004) 2. Carling, D. (2004) 3. Hawley, S.A. et al. (4. Lizcano, J.M. et al. (5. Shaw, R.J. et al. (20 6. Woods, A. et al. (20 7. Warden, S.M. et al.	<i>rends Biochem Sci 2</i> 1996) <i>J Biol Chem 27</i> (2004) <i>EMBO J</i> 23, 83 04) <i>Proc Natl Acad S</i> 103) <i>J Biol Chem</i> 278,	9, 18-24. 1, 27879-87. 3-43. <i>ci USA</i> 101, 3329-35. 28434-42.		
Species Reactiv	vity	Species reactivity is d	etermined by testing	g in at least one appro	ved application (e.g., v	vestern 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) IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)				
Cross-Reactivity Key	H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey				
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