#5256 Store at -20C

MW (kDa)

Phospho-AMPKα (Thr172) (40H9) Rabbit mAb (Biotinylated)



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62

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Applications: WB	Reactivity: H M R Hm Mk Dm Sc	Sensitivity: Endogenous	MW (kDa): 62	Source/Isotype: Rabbit IgG	UniProt ID: #Q13131, #P54646	Entrez-Gene Id: 5562, 5563
Product Usage Application Information Western Blotting				Dilution 1:1000		
Storage	Supplied in 136 mM NaCl, 2.6 mM KCl, 150% glycerol. Store at -20°C. Do not aliq			12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and quot the antibodies.		
Specificity / Sens	phos		72. The antibody	detects both α1 and α	etects endogenous AMPk 12 isoforms of the catalytic	
Species predicte react based on 1 sequence homol	00%	ken, Zebrafish, Bov	vine, Pig			
Source / Purifica		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Thr172 of human AMPK α protein.				
Product Descript	antib	This Cell Signaling Technology antibody is conjugated to biotin under optimal conditions. The biotinylated antibody is expected to exhibit the same species cross-reactivity as the unconjugated Phospho-AMPKα (Thr172) (40H9) Rabbit mAb #2535.				

Background

AMP-activated protein kinase (AMPK) is highly conserved from yeast to plants and animals and plays a key role in the regulation of energy homeostasis (1). AMPK is a heterotrimeric complex composed of a catalytic α subunit and regulatory β and γ subunits, each of which is encoded by two or three distinct genes $(\alpha 1, 2; \beta 1, 2; \gamma 1, 2, 3)$ (2). The kinase is activated by an elevated AMP/ATP ratio due to cellular and environmental stress, such as heat shock, hypoxia, and ischemia (1). The tumor suppressor LKB1, in association with accessory proteins STRAD and MO25, phosphorylates AMPKα at Thr172 in the activation loop, and this phosphorylation is required for AMPK activation (3-5). AMPKα is also phosphorylated at Thr258 and Ser485 (for α 1; Ser491 for α 2). The upstream kinase and the biological significance of these phosphorylation events have yet to be elucidated (6). The β1 subunit is post-translationally modified by myristoylation and multi-site phosphorylation including Ser24/25, Ser96, Ser101, Ser108, and Ser182 (6,7). Phosphorylation at Ser108 of the β1 subunit seems to be required for AMPK activation, while phosphorylation at Ser24/25 and Ser182 affects AMPK localization (7). Several mutations in AMPKy subunits have been identified, most of which are located in the putative AMP/ATP binding sites (CBS or Bateman domains). Mutations at these sites lead to reduction of AMPK activity and cause glycogen accumulation in heart or skeletal muscle (1,2). Accumulating evidence indicates that AMPK not only regulates the metabolism of fatty acids and glycogen, but also modulates protein synthesis and cell growth through EF2 and TSC2/mTOR pathways, as well as blood flow via eNOS/nNOS (1).

Background References

- 1. Hardie, D.G. (2004) J Cell Sci 117, 5479-87.
- 2. Carling, D. (2004) Trends Biochem Sci 29, 18-24.
- 3. Hawley, S.A. et al. (1996) J Biol Chem 271, 27879-87.
- 4. Lizcano, J.M. et al. (2004) EMBO J 23, 833-43.
- 5. Shaw, R.J. et al. (2004) Proc Natl Acad Sci USA 101, 3329-35.
- 6. Woods, A. et al. (2003) J Biol Chem 278, 28434-42.
- 7. Warden, S.M. et al. (2001) Biochem J 354, 275-83.

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

WB: Western Blotting

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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