a1 protein only when phosphorylated at Tyr10. The antibody cross-reacts with an induced 75-80 kDa

Phoenba

sequence homology:

Phospho-Na,K-ATPase α1 (Tyr10) (E1Y9C) Rabbit mAb							
Stor					Orders:	877-616-CELL (2355) orders@cellsignal.com	
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For Research Use On	ly. Not for Use in	Diagnostic Proc	edures.				
Applications: WB	Reactivity: H M	Sensitivity: Endogenous	MW (kDa): 100	Source/Isotype: Rabbit IgG	UniProt ID: #P05023	Entrez-Gene Id: 476	
Product Usage Information	Application			Dilution			
	We	estern Blotting			1:1000		
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		Phospho-Na,K-ATPase α 1 (Tyr10) (E1Y9C) Rabbit mAb recognizes endogenous levels of Na,K-ATPase					

doublet of unknown origin.

Species predicted to Rat, Bovine, Dog, Pig react based on 100%

Source / Purification	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to
	residues surrounding Tyr10 of human Na,K-ATPase α 1 protein.

Background The Na,K-ATPase is an integral membrane heterodimer belonging to the P-type ATPase family. This ion channel uses the energy derived from ATP hydrolysis to maintain membrane potential by driving sodium export and potassium import across the plasma membrane against their electrochemical gradients. It is composed of a catalytic α subunit and a β subunit (reviewed in 1). Several phosphorylation sites have been identified for the α 1 subunit. Tyr10 is phosphorylated by an as yet undetermined kinase (2), Ser16 and Ser23 are phosphorylated by PKC, and Ser943 is phosphorylated by PKA (3-5). All of these sites have been implicated in the regulation of enzyme activity in response to hormones and neurotransmitters, altering trafficking and kinetic properties of Na,K-ATPase. Altered phosphorylation in response to angiotensin II stimulates activity in the rat proximal tubule (6). Na,K-ATPase is also involved in other signal transduction pathways. Insulin regulates its localization in differentiated primary human skeletal muscle cells, and this regulation is dependent on ERK1/2 phosphorylation of the α subunit (7). Na,K-ATPase and Src form a signaling receptor complex that affects regulation of Src kinase activity and, subsequently, its downstream effectors (8,9).

Background References	1. Therien, A.G. and Blostein, R. (2000) Am J Physiol Cell Physiol 279, C541-66.
-	2. Féraille, E. et al. (1999) <i>Mol Biol Cell</i> 10, 2847-59.
	3. Fisone, G. et al. (1994) <i>J Biol Chem</i> 269, 9368-73.
	 Feschenko, M.S. and Sweadner, K.J. (1995) J Biol Chem 270, 14072-7.
	5. Beguin, P. et al. (1994) <i>J Biol Chem</i> 269, 24437-45.
	6. Yingst, D.R. et al. (2004) Am J Physiol Renal Physiol 287, F713-21.
	7. Al-Khalili, L. et al. (2004) J Biol Chem 279, 25211-8.
	8. Tian, J. et al. (2006) Mol Biol Cell 17, 317-26.
	9. Liang, M. et al. (2006) J Biol Chem 281, 19709-19.

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 Cross-Reactivity Key	WB: Western Blotting

1/1/24, 2:29 PM	Phospho-Na,K-ATPase α 1 (Tyr10) (E1Y9C) Rabbit mAb (#13566) Datasheet Without Images Cell Signaling Te
	 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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