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# Phospho-Na,K-ATPase α1 (Ser16) Antibody



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Applications: WB	Reactivity:	Sensitivity: Endogenous	MW (kDa): 100	Source: Rabbit	UniProt ID: #P05023	Entrez-Gene Id: 476	
Product Usage Information	Application			Dilution			
	Western Blotting			1:1000			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.					
Specificity / Sensiti	i <b>vity</b> Pho	Phospho-Na,K-ATPase α1 (Ser16) Antibody recognizes endogenous levels of Na,K-ATPase α1 only when					

the protein, corresponding to Ser11 of the mature cleaved form.

Species predicted to react based on 100% sequence homology:

Mouse, Bovine, Pia

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser16 of rat Na,K-ATPase  $\alpha$ 1. Antibodies are purified using protein A and peptide affinity chromatography.

phosphorylated at Ser16. The residue number, Ser16, is based on the sequence of the immature form of

### **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  $\alpha$  subunit and a  $\beta$  subunit (reviewed in 1). Several phosphorylation sites have been identified for the  $\alpha$ 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  $\alpha$  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**

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- 2. Féraille, E. et al. (1999) Mol Biol Cell 10, 2847-59.
- 3. Fisone, G. et al. (1994) J Biol Chem 269, 9368-73.
- 4. Feschenko, M.S. and Sweadner, K.J. (1995) J Biol Chem 270, 14072-7.
- 5. Beguin, P. et al. (1994) J Biol Chem 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 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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