

**Na,K-ATPase  $\alpha$ 1 (D4Y7E) Rabbit mAb**

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**For Research Use Only. Not for Use in Diagnostic Procedures.**

<b>Applications:</b> W, IHC-P, IF-IC	<b>Reactivity:</b> H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 100	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P05023	<b>Entrez-Gene Id:</b> 476
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**Product Usage Information****Application**

Western Blotting  
Immunohistochemistry (Paraffin)  
Immunofluorescence (Immunocytochemistry)

**Dilution**

1:1000  
1:250  
1:100

**Storage**

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100  $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at  $-20^{\circ}\text{C}$ . Do not aliquot the antibody.

For a carrier free (BSA and azide free) version of this product see product #99935.

**Specificity/Sensitivity**

Na,K-ATPase  $\alpha$ 1 (D4Y7E) Rabbit mAb recognizes endogenous levels of total Na,K-ATPase  $\alpha$ 1 protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro12 of human Na,K-ATPase  $\alpha$ 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  $\alpha$  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**

1. Therien, A.G. and Blostein, R. (2000) *Am J Physiol Cell Physiol* 279, C541-66.
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  $4^{\circ}\text{C}$  with gentle shaking, overnight.

**Applications Key**

**W:** Western Blotting **IHC-P:** Immunohistochemistry (Paraffin) **IF-IC:** Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key**

**H:** Human **Mk:** Monkey

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