Revision 1	
va,K-ATPase β1 (D6U8Q) Rabbit mAb	Cell Signaling TECHNOLOGY
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Applications: W	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 45-55	Source/Isotype: Rabbit IgG	UniProt ID: #P05026	Entrez-Gene Id: 481	
Product Usage Information	2	Application Western Blotting			Dilution 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/Ser	nsitivity	Na,K-ATPase β 1 (D6U8Q) Rabbit mAb recognizes endogenous levels of total Na,K-ATPase β 1 protein.					
Source / Purifi	cation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro200 of human Na,K-ATPase β1 protein.					
Background		channel uses the ene sodium export and po gradients. It is compo phosphorylation sites undetermined kinase PKA (3-5). All of these hormones and neuro phosphorylation in re ATPase is also involve differentiated priman phosphorylation of th affects regulation of S Na,K-ATPase β1 is the and translocation of t mediates the trans-di integrity of tight junct of Na,K-ATPase (18). F Hedgehog signaling p	rgy derived from A otassium import ac sed of a catalytic of a catalytic of shave been identifi (2), Ser16 and Ser2 sites have been im transmitters, alteri sponse to angioter d in other signal tr y human skeletal m e α subunit (7). Na fric kinase activity a non-catalytic subu he catalytic α subu merization of Na,K tions (13-17). Gluta Research studies ha bathway and may b e encoding Na,K-AT	The heterodimer belonging TP hydrolysis to maintain ross the plasma membra is ubunit and a β subunit ed for the a1 subunit. Ty 23 are phosphorylated by plicated in the regulation ing trafficking and kinetic rafficking and kinetic regulation rafficking and kinetic rafficking and k	i membrane potent ane against their ele (reviewed in 1). Sev r10 is phosphorylat γ PKC, and Ser943 is n of enzyme activity properties of Na,K- in the rat proximal sulin regulates its le ilation is dependen a signaling receptor vnstream effectors required for stabiliz rane(10-12). Na,K-A oring cells where it ase β 1 regulates th ase β 1 regulates th ase β 1 regulates th ase β 1 urgulates the ase β 1 sa target of g tumor developme	ial by driving ectrochemical veral eed by an as yet sphosphorylated by in response to -ATPase. Altered tubule (6). Na,K- ocalization in t on ERK1/2 complex that (8,9). ation, maturation, TPase β 1 also regulates the e ion pump activity the Sonic ent and progression	
Background R	eferences	 2. Féraille, E. et al. (19 3. Fisone, G. et al. (19) 4. Feschenko, M.S. an 5. Beguin, P. et al. (19) 6. Yingst, D.R. et al. (21) 7. Al-Khalili, L. et al. (22) 8. Tian, J. et al. (2006) 9. Liang, M. et al. (2006) 9. Liang, M. et al. (2001) 10. Beggah, A.T. et al. 11. Hasler, U. et al. (19) 12. Rajasekaran, S.A. 13. Rajasekaran, A.K. 15. Bab-Dinitz, E. et al 16. Tokhtaeva, E. et al 	99) Mol Biol Cell 10 94) J Biol Chem 269 d Sweadner, K.J. (19 94) J Biol Chem 269 004) Am J Physiol F 1004) J Biol Chem 281, (1997) J Biol Chem 281, (1997) J Biol Chem 27 et al. (2004) Mol Bii et al. (2004) Mol Bii and Rajasekaran, S J. (2009) Biochemis J. (2001) J Biol Chem 12) Am J Physiol Ce (2012) Free Radice 1 5) Mol Cancer 14, 1	 a) 9368-73. b) J Biol Chem 270, 140 b) J Biol Chem 270, 140 c) 24437-45. Renal Physiol 287, F713-2. 7-26. 7-26. 19709-19. 272, 10318-26. 3, 30826-35. ol Cell 15, 3224-32. ol Cell 12, 3717-32. A. (2003) Am J Physiol Retry 48, 8684-91. 286, 25801-12. ell Physiol 302, C1271-81. siol Med 53, 2263-8. 59. 	72-7. 1. enal Physiol 285, F3	88-96.	

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	W: Western Blotting
Cross-Reactivity Key	H: Human M: Mouse R: Rat
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