



FXYD2 Polyclonal Antibody

Catalog No	YP-Ab-10911
Isotype	IgG
Reactivity	Human;Rat;Mouse;
Applications	IHC; ELISA
Gene Name	FXYD2 ATP1C ATP1G1
Immunogen	Synthesized peptide derived from human FXYD2 AA range: 10-90
Specificity	This antibody detects endogenous levels of human FXYD2
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from mouse ascites by affinity-chromatography using specific immunogen.
Dilution	IHC-p 1:50-200, ELISA(peptide)1:5000-20000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	ATP1C;ATP1G1;FXYD 2;FXYD domain-containing ion transport regulator 2; Sodium/potassium-;FXYD2;FXYD2 ATP1C ATP1G1;Sodium/potassium-transporting ATPase subunit gamma (Na+)/K(+) ATPase subunit gamma; Sodium pump gamma chain)
Observed Band	7-10 kDa
Calculated Molecular Weight	7 kDa
Cell Pathway	Membrane ; Single-pass type III membrane protein .
Tissue Specificity	Expressed in the distal convoluted tubule in the kidney. Found on basolateral membranes of nephron epithelial cells.
Function	disease:Defects in FXYD2 are the cause of hypomagnesemia type 2 (HOMG2) [MIM:154020]; also known as dominant renal hypomagnesemia or hypomagnesemia with hypocalciuria. HOMG2 is a disorder due to primary renal wasting of magnesium. Plasma levels of other electrolytes are normal. The only abnormality found, in addition to low magnesium levels, is lowered renal excretion of calcium resulting in hypocalciuria.,function:May be involved in forming the receptor site for cardiac glycoside binding or may modulate the transport function of the sodium ATPase.,sequence caution:Wrong choice of frame.,similarity:Belongs to the FXYD family.,subunit:Composed of three



subunits: alpha (catalytic), beta and gamma.,tissue specificity:Expressed in the distal convoluted tubule in the kidney. Found on basolateral membranes of nephron epithelial cells.,

Background

FXYP domain containing ion transport regulator 2(FXYP2) Homo sapiens This gene encodes a member of the FXYP family of transmembrane proteins. This particular protein encodes the sodium/potassium-transporting ATPase subunit gamma. Mutations in this gene have been associated with Renal Hypomagnesemia-2. Alternatively spliced transcript variants have been described. Read-through transcripts have been observed between this locus and the upstream FXYP domain-containing ion transport regulator 6 (FXYP6, GeneID 53826) locus.[provided by RefSeq, Feb 2011],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images