



# FXYD2 Polyclona Antibody

<b>Catalog No</b>	YP-Ab-10911
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Rat;Mouse;
<b>Applications</b>	IHC; ELISA
<b>Gene Name</b>	FXYD2 ATP1C ATP1G1
<b>Protein Name</b>	FXYD2
<b>Immunogen</b>	Synthesized peptide derived from human FXYD2 AA range: 10-90
<b>Specificity</b>	This antibody detects endogenous levels of human FXYD2
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from mouse ascites by affinity-chromatography using specific immunogen.
<b>Dilution</b>	IHC-p 1:50-200, ELISA(peptide)1:5000-20000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	Sodium/potassium-transporting ATPase subunit gamma (Na(+)/K(+)) ATPase subunit gamma;FXYD domain-containing ion transport regulator 2;Sodium pump gamma chain)
<b>Observed Band</b>	
<b>Cell Pathway</b>	Membrane ; Single-pass type III membrane protein .
<b>Tissue Specificity</b>	Expressed in the distal convoluted tubule in the kidney. Found on basolateral membranes of nephron epithelial cells.
<b>Function</b>	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**