

Rabbit Anti-Uromucoid antibody

SL2189R

Product Name:	Uromucoid
Chinese Name:	
Alias:	UMOD; ADMCKD2; FJHN; HNFJ; HNFJ1; MCKD2; medullary cystic kidney disease 2 (autosomal dominant); Tamm Horsfall glycoprotein; Tamm Horsfall urinary glycoprotein; Tamm-Horsfall urinary glycoprotein; THGP; THP; Umod; UROM_MOUSE; uromodulin (uromucoid, Tamm-Horsfall glycoprotein); Uromodulin; Uromodulin, secreted form.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Cow,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow- Cyt=1µg/TestIF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	61/65kDa
Cellular localization:	The cell membraneSecretory protein
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from mouse MCKD2:351-450/642
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
Product Detail:	The protein encoded by this gene is the most abundant protein in mammalian urine under physiological conditions. Its excretion in urine follows proteolytic cleavage of the ectodomain of its glycosyl phosphatidylinosital-anchored counterpart that is situated on

the luminal cell surface of the loop of Henle. This protein may act as a constitutive inhibitor of calcium crystallization in renal fluids. Excretion of this protein in urine may provide defense against urinary tract infections caused by uropathogenic bacteria. Defects in this gene are associated with the renal disorders medullary cystic kidney disease-2 (MCKD2), glomerulocystic kidney disease with hyperuricemia and isosthenuria (GCKDHI), and familial juvenile hyperuricemic nephropathy (FJHN). Alternative splicing of this gene results in multiple transcript variants. [provided by RefSeq, Jul 2013].

Function:

Uromodulin: Functions in biogenesis and organization of the apical membrane of epithelial cells of the thick ascending limb of Henle's loop (TALH), where it promotes formation of complex filamentous gel-like structure providing the water barrier permeability. May serve as a receptor for binding and endocytosis for cytokines (IL-1, IL-2) and TNF. Facilitates neutrophil migration across renal epithelial (By similarity). Uromodulin, secreted form: Secreted into urine after proteolytically cleaveage. Into the urine, may contribute to colloid osmotic pressure, retards passage of positively charged electrolytes, prevents urinary tract infection and modulates formation of supersaturated salts and their crystals.

Subcellular Location:

Apical cell membrane; Lipid-anchor, GPI-anchor (By similarity). Basolateral cell membrane; Lipid-anchor, GPI-anchor (By similarity). Cell projection, cilium membrane (By similarity). Note=Only a small fraction is sorts to the basolateral pole of tubular epithelial cells compared to apical localization (By similarity). Uromodulin, secreted form: Secreted (By similarity).

Post-translational modifications: N-glycosylated.

Similarity:

Contains 3 EGF-like domains. Contains 1 ZP domain.

SWISS: Q91X17

Gene ID: 22242

Database links:

Entrez Gene: 7369Human

Omim: 191845Human

<u>SwissProt: P07911</u>Human











performed.

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