

## UMOD polyclonal antibody

Catalog # PAB15392      Size 2 mL

### Specification

<b>Product Description</b>	Goat polyclonal antibody raised against UMOD.
<b>Immunogen</b>	Human UMOD.
<b>Host</b>	Goat
<b>Reactivity</b>	Human
<b>Specificity</b>	This product is suitable for use as a precipitating antibody. Cross-reactivity to other species may exist. Antibody activity to other serum proteins is not present.
<b>Form</b>	Lyophilized
<b>Recommend Usage</b>	ELISA (1:400-1:2000) Immunohistochemistry (1:50-1:200) Western Blot (1:100-1:500) The optimal working dilution should be determined by the end user.
<b>Storage Buffer</b>	Lyophilized from 20 mM sodium phosphate buffer, 140 mM NaCl, pH 7.3 (0.05% sodium azide).
<b>Storage Instruction</b>	Store at 4°C on dry atmosphere. After reconstitution with 2 mL of deionized water, store at -20°C. Aliquot to avoid repeated freezing and thawing.
<b>Note</b>	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

### Applications

- Western Blot
- Enzyme-linked Immunoabsorbent Assay
- Immunodiffusion

## Gene Info — UMOD

Entrez GeneID	<a href="#">7369</a>
Gene Name	UMOD
Gene Alias	ADMCKD2, FJHN, HNFJ, MCKD2, THGP, THP
Gene Description	uromodulin
Omim ID	<a href="#">162000</a> <a href="#">191845</a> <a href="#">603860</a> <a href="#">609886</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	<p>This gene encodes uromodulin, the most abundant protein in normal urine. Its excretion in urine follows proteolytic cleavage of the ectodomain of its glycosyl phosphatidylinositol-anchored counterpart that is situated on the luminal cell surface of the loop of Henle. Uromodulin may act as a constitutive inhibitor of calcium crystallization in renal fluids. Excretion of uromodulin in urine may provide defense against urinary tract infections caused by uropathogenic bacteria. Defects in this gene are associated with the autosomal dominant renal disorders medullary cystic kidney disease-2 (MCKD2) and familial juvenile hyperuricemic nephropathy (FJHN). These disorders are characterized by juvenile onset of hyperuricemia, gout, and progressive renal failure. While several transcript variants may exist for this gene, the full-length nature of only two have been described to date. These two represent the major variants of this gene and encode the same isoform. [provided by RefSeq]</p>
Other Designations	OTTHUMP00000162212[Tamm-Horsfall glycoprotein]uromodulin (uromucoid, Tamm-Horsfall glycoprotein)uromucoid

## Disease

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