

## Product datasheet for TP721058XL

## OriGene Technologies, Inc.

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## Angiotensinogen (AGT) (NM\_000029) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human angiotensinogen (serpin peptidase inhibitor, clade A,

member 8) (AGT)

Asp34-Ala485

Species: Human Expression Host: HEK293

Expression cDNA Clone

or AA Sequence:

Tag: C-His

Predicted MW: 51.7 kDa

**Purity:** >95% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

Endotoxin: Endotoxin level is < 0.1 ng/μg of protein (< 1 EU/μg)

Storage: Store at -80°C.

Stability: Stable for at least 3 months from date of receipt under proper storage and handling

conditions.

RefSeq: <u>NP 000020</u>

Locus ID: 183

**UniProt ID:** <u>P01019</u>, <u>B0ZBE2</u>, <u>B2R5S1</u>

RefSeq Size: 2587 Cytogenetics: 1q42.2 RefSeq ORF: 1455

Synonyms: ANHU; hFLT1; SERPINA8





## Angiotensinogen (AGT) (NM\_000029) Human Recombinant Protein - TP721058XL

**Summary:** 

**Protein Families:** 

The protein encoded by this gene, pre-angiotensinogen or angiotensinogen precursor, is expressed in the liver and is cleaved by the enzyme renin in response to lowered blood pressure. The resulting product, angiotensin I, is then cleaved by angiotensin converting enzyme (ACE) to generate the physiologically active enzyme angiotensin II. The protein is involved in maintaining blood pressure, body fluid and electrolyte homeostasis, and in the pathogenesis of essential hypertension and preeclampsia. Mutations in this gene are associated with susceptibility to essential hypertension, and can cause renal tubular dysgenesis, a severe disorder of renal tubular development. Defects in this gene have also been associated with non-familial structural atrial fibrillation, and inflammatory bowel disease. [provided by RefSeq, Nov 2019]

Druggable Genome, Secreted Protein

**Protein Pathways:** Renin-angiotensin system