





Human angiotensinogen (aGT) ELISA Kit

Product Code	CSB-E08564h	
Abbreviation	AGT	
Target Name	angiotensinogen (serpin peptidase inhibitor, clade A, member 8)	
Uniprot No.	P01019	
Alias	ANHU, FLJ92595, FLJ97926, SERPINA8, alpha-1 antiproteinase, antitrypsin angiotensin I angiotensin II angiotensinogen pre-angiotensinogen serine (or cysteine) proteinase inhibitor	
Product Type	ELISA Kit	
Immunogen Species	Homo sapiens (Human) serum, plasma, urine,cell culture supernates 78 pg/mL-5000 pg/mL	
Sample Types		
Detection Range		
Sensitivity	19.5 pg/mL	
Assay Time	1-5h	
Sample Volume	50-100ul	
Detection Wavelength	450 nm	
Lead Time	3-5 working days after you place the order, and it takes another 3-5 days for delivery via DHL or FedEx.	
Research Area	Cardiovascular	
Gene Names	AGT	
Tag Info	quantitative	
Protein Description	Sandwich	
Description	This Human AGT ELISA Kit was designed for the quantitative measurement of Human AGT protein in serum, plasma, urine,cell culture supernates. It is a Sandwich ELISA kit, its detection range is 78 pg/mL-5000 pg/mL and the sensitivity is 19.5 pg/mL.	
Target Details	This protein, pre-angiotensinogen or angiotensinogen precursor, is expressed in	

Target Details

This protein, 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 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.







Product Precision

Intra-assay Precision (Precision within an assay): CV%<8%

Three samples of known concentration were tested twenty times on one plate to assess.

Inter-assay Precision (Precision between assays): CV%<10%

Three samples of known concentration were tested in twenty assays to assess.

Linearity

To assess the linearity of the assay, samples were spiked with high concentrations of human aGT in various matrices and diluted with the Sample Diluent to produce samples with values within the dynamic range of the assay.

?	Sample	Serum(n=4)
1:200	Average %	90
1.200	Range %	85-100
1:400	Average %	97
1.400	Range %	91-105
1:800	Average %	98
1.000	Range %	93-110
1:1600	Average %	93
1.1000	Range %	85-105

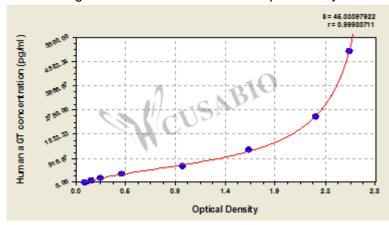
Recovery

The recovery of human aGT spiked to levels throughout the range of the assay in various matrices was evaluated. Samples were diluted prior to assay as directed in the Sample Preparation section.

Sample Type	Average % Recovery	Range
Serum (n=5)	90	85-100
EDTA plasma (n=4)	97	90-105

Typical

These standard curves are provided for demonstration only. A standard curve should be generated for each set of samples assayed.



pg/ml OD1 OD2 Average Corrected

5000	2.533 2.589 2.561	2.467
2500	2.234 2.268 2.251	2.157
1250	1.618 1.628 1.623	1.529
625	1.005 1.014 1.010	0.916
312	0.423 0.453 0.438	0.344
156	0.236 0.243 0.240	0.146
78	0.149 0.156 0.153	0.059
0	0.091 0.096 0.094	?



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Msds

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