# **Elabscience**®

# Human AGT Antibody Pair Set

Catalog No.	E-KAB-0543	Applications	ELISA
Synonyms	ANHU;SERPINA8		

#### Kit components & Storage

Title	Specifications	Storage
Human AGT Capture Antibody	1 vial, 100 µ g	Store at $-20^{\circ}$ C for one year.
		Avoid freeze/thaw cycles.
Human AGT Detection Antibody (Biotin)	1 vial, 50 μL	Store at $-20^{\circ}$ C for one year.
		Avoid freeze/thaw cycles.

Note: Centrifuge before opening to ensure complete recovery of vial contents.

### **Product Information**

Items		Characteristic (E-KAB-0543)		
		Human AGT Capture Antibody	Human AGT Detection Antibody (Biotin)	
Immunogen	Immunogen	Recombinant Human AGT protien	Recombinant Human AGT protien	
Information	Swissprot	P01019		
Product details	Reactivity	Human	Human	
	Host	Rabbit	Rabbit	
	Conjugation	Unconjugated	Biotin	
	Concentration	0.5 mg/mL	/	
	Buffer	PBS with 0.04% Proclin 300; 50%	PBS with 0.04% Proclin 300; 1%	
		glycerol; pH 7.5	protective protein; 50% glycerol; pH	
			7.5	
	Purify	Antigen Affinity	Protein A or G	
	Specificity	Detects Human AGT in ELISAs.		

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### Applications

Human AGT Sandwich ELISA Assay:

	Recommended	Reagent	Images
	Concentration/Dilution		
ELISA	0.5-4 μg/mL	Human AGT Capture	
Capture		Antibody	10
			- t ensity
ELISA	1:1000-1:10000	Human AGT Detection	Optical Dansity
Detection		Antibody (Biotin)	
			0.1
			1 10 100 1000 Human AGT Concentration(ng/mL)
			ruman AGT Concentration(ng/mL)

Note: This standard curve is only for demonstration purposes. A standard curve should be generated for each assay!

## Background

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 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.

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