Angiotensin antibody

Catalog No: #22961

Description



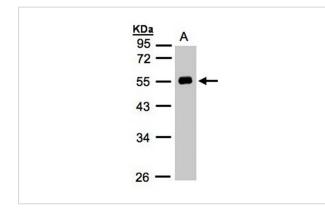
Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

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Product Name	Angiotensin antibody		
Host Species	Rabbit		
Clonality	Polyclonal		
Purification	Purified by antigen-affinity chromatography.		
Applications	WB IHC IF		
Species Reactivity	Ни		
Immunogen Type	Recombinant protein		
Immunogen Description	Recombinant protein fragment contain a sequence corresponding to a region within amino acids 1 and 446 of		
	Human AGT		
Target Name	Angiotensin		
Other Names	ANHU; FLJ92595; FLJ97926; SERPINA8		
Accession No.	Swiss-Prot:P01019Gene ID:183		
Uniprot	P01019		
GenelD	183;		
Concentration	1mg/ml		
Formulation	Supplied in 0.1M Tris-buffered saline with 10% Glycerol (pH7.0). 0.01% Thimerosal was added as a		
	preservative.		
Storage	Store at -20°C for long term preservation (recommended). Store at 4°C for short term use.		

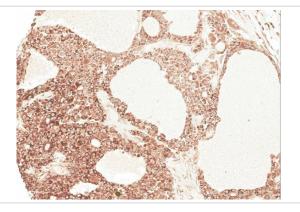
Application Details

Predicted MW: 53kd		
Western blotting: 1:500-1:3000		
Immunohistochemistry: 1:100-1:250		
Immunofluorescence: 1:100-1:200		

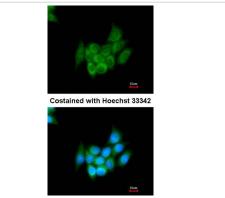
Images



Sample(30 ug whole cell lysate) A: Hep G2 10% SDS PAGE Primary antibody diluted at 1: 1000



Immunohistochemical analysis of paraffin-embedded Hepatocellular carcinoma Huh7 xenograft, using AGT antibody at 1: 100 dilution.



Immunofluorescence analysis of methanol-fixed HepG2, using Angiotensinogen antibody at 1: 200 dilution.

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. [provided by RefSeq]

Note: This product is for in vitro research use only