AGXT2 Antibody

PACO57460

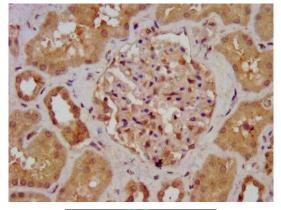


Product Information	
Size:	Protein Background:
50ug	Can metabolize asymmetric dimethylarginine (ADMA) via transamination to alpha- keto-delta-(NN-dimethylguanidino) valeric acid, (DMGV). ADMA is a potent inhibitor of nitric-oxide (NO) synthase, and this activity provides mechanism through which the kidney regulates blood pressure.
Reactivity:	
Human	
Source:	Gene ID:
Rabbit	AGXT2
lsotype:	Uniprot
lgG	Q9BYV1
Applications:	Synonyms:
ELISA, WB, IHC, IF	Alanineglyoxylate aminotransferase 2, mitochondrial (AGT 2) (EC 2.6.1.44) ((R)-3- amino-2-methylpropionatepyruvate transaminase) (EC 2.6.1.40) (Beta-ALAAT II) (Beta-
Recommended dilutions:	alanine-pyruvate aminotransferase) (D-AIBAT), AGXT2, AGT2
ELISA:1:2000-1:10000, WB:1:500-1:5000, IHC:1:500-1:1000, IF:1:200-1:500	Immunogen:
	Recombinant Human Alanineglyoxylate aminotransferase 2, mitochondrial protein (196-328AA).
	Storage:

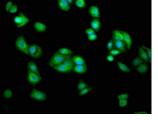
Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

	102
$_{\rm 120KD}\!\rightarrow$	Ŷ
90KD→	
50KD \rightarrow	-
$_{35 \text{KD}} \rightarrow$	
$25 \text{KD} \rightarrow$	
$_{20\text{KD}} \rightarrow$	

Western Blot. Positive WB detected in: LO2 whole cell lysate. All lanes: AGXT2 antibody at 4.9µg/ml. Secondary. Goat polyclonal to rabbit IgG at 1/50000 dilution. Predicted band size: 58, 50 kDa. Observed band size: 50 kDa.



IHC image of PACO57460 diluted at 1:700 and staining in paraffinembedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of HepG2 cells with PACO57460 at 1:233, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).