产品名称: AGTR1 Rabbit Polyclonal Antibody

产品货号: APRab06685



## 产品概述 (Summary)

产品名称 (Production Name) AGTR1 Rabbit Polyclonal Antibody

描述 (Description) Rabbit polyclonal Antibody

宿主 (Host) Rabbit 应用 (Application) WB,ELISA

种属反应性 (Reactivity) Human, Mouse, Rat

#### 产品性能 (Performance)

偶联物 (Conjugation)Unconjugated修饰 (Modification)Unmodified

同种型 (Isotype) IgG

克隆 (Clonality) Polyclonal 形式 (Form) Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% 储存溶液 (Buffer)

New type preservative N.

纯化方式 (Purification) Affinity purification

## 免疫原信息 (Immunogen)

基因名 (Gene Name) AGTR1 AGTR1A AGTR1B AT2R1 AT2R1B

Type-1 angiotensin II receptor (AT1AR) (AT1BR) (Angiotensin II type-1

别名 (Alternative Names) receptor) (AT1)

基因 ID (Gene ID) 185.0

蛋白 ID (SwissProt ID) P30556.Synthesized peptide derived from human AGTR1 Polyclonal

# 产品应用(Application)

**稀释比 (Dilution Ratio)** WB 1:500-1:2000,ELISA 1:10000-1:20000

蛋白分子量 (Molecular Weight) 41kDa

# 研究背景 (Background)

Web:https://www.enkilife.cn E-mail:order@enkilife.cn (销售) tech@enkilife.cn (技支持) Tel:027-87002838

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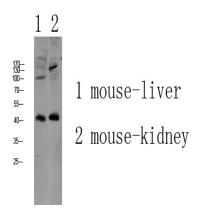


Angiotensin II is a potent vasopressor hormone and a primary regulator of aldosterone secretion. It is an important effector controlling blood pressure and volume in the cardiovascular system. It acts through at least two types of receptors. This gene encodes the type 1 receptor which is thought to mediate the major cardiovascular effects of angiotensin II. This gene may play a role in the generation of reperfusion arrhythmias following restoration of blood flow to ischemic or infarcted myocardium. It was previously thought that a related gene, denoted as AGTR1B, existed; however, it is now believed that there is only one type 1 receptor gene in humans. Multiple alternatively spliced transcript variants have been reported for this gene. [provided by RefSeq, Jul 2012], disease:Defects in AGTR1 are a cause of renal tubular dysgenesis (RTD) [MIM:267430]. RTD is an autosomal recessive severe disorder of renal tubular development characterized by persistent fetal anuria and perinatal death, probably due to pulmonary hypoplasia from early-onset oligohydramnios (the Potter phenotype)., function:Receptor for angiotensin II. Mediates its action by association with G proteins that activate a phosphatidylinositol-calcium second messenger system., online information:Angiotensin receptor entry, online information:The Singapore human mutation and polymorphism database, PTM:C-terminal Ser or Thr residues may be phosphorylated., similarity:Belongs to the G-protein coupled receptor 1 family., tissue specificity:Liver, lung, adrenal and adrenocortical adenomas.

#### 研究领域 (Research Area)

Calcium; Neuroactive ligand-receptor interaction; Vascular smooth muscle contraction; Renin-angiotensin system;

#### 图片 (Image Data)



Western blot analysis of various lysate, antibody was diluted at 1000. Secondary antibody was diluted at 1:20000

# 注意事项 (Note)

For research use only.

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