产品名称: LSHR Rabbit Polyclonal Antibody

产品货号: APRab13459



## 产品概述 (Summary)

产品名称 (Production Name) LSHR 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) LHCGR LCGR LGR2 LHRHR

Lutropin-choriogonadotropic hormone receptor (LH/CG-R) (Luteinizing 別名 (Alternative Names)

hormone receptor) (LHR) (LSH-R)

基因 ID (Gene ID) 3973.0

蛋白 ID (SwissProt ID) P22888.Synthesized peptide derived from human LSHR 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)** 80kDa

## 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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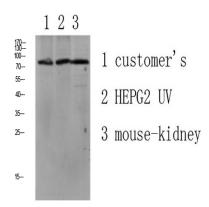


This gene encodes the receptor for both luteinizing hormone and choriogonadotropin. This receptor belongs to the G-protein coupled receptor 1 family, and its activity is mediated by G proteins which activate adenylate cyclase. Mutations in this gene result in disorders of male secondary sexual character development, including familial male precocious puberty, also known as testotoxicosis, hypogonadotropic hypogonadism, Leydig cell adenoma with precocious puberty, and male pseudohermaphtoditism with Leydig cell hypoplasia. [provided by RefSeq, Jul 2008], alternative products: Additional isoforms seem to exist, disease: Defects in LHCGR are a cause of familial male precocious puberty (FMPP) [MIM:176410]; also known as testotoxicosis. In FMPP the receptor is constitutively activated., disease: Defects in LHCGR are a cause of Leydig cell hypoplasia (LCH) [MIM:152790]. LCH is an autosomal recessive disease characterized by male pseudohermaphroditism. In LCH the testes are small with marked immaturity of the Leydig cells which correlates with undetectable plasma testosterone levels and elevated gonadotropins., function: Receptor for lutropin-choriogonadotropic hormone. The activity of this receptor is mediated by G proteins which activate adenylate cyclase., online information: Glycoprotein-hormone Receptors Information System, similarity: Belongs to the G-protein coupled receptor 1 family., similarity: Belongs to the G-protein coupled receptor 1 family. FSH/LSH/TSH subfamily., similarity: Contains 7 LRR (leucine-rich) repeats., tissue specificity: Gonadal and thyroid cells.,

#### 研究领域 (Research Area)

Calcium; Neuroactive ligand-receptor interaction;

#### 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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