产品名称: LKB1 (phospho Thr189) Rabbit Polyclonal

**Antibody** 

产品货号: APRab04960



### 产品概述 (Summary)

产品名称 (Production Name) LKB1 (phospho Thr189) 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

## 产品性能 (Performance)

偶联物 (Conjugation) Unconjugated 修饰 (Modification) Phosphorylated

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) STK11

STK11; LKB1; PJS; Serine/threonine-protein kinase STK11; Liver kinase B1; 别名 (Alternative Names)

LKB1; hLKB1; Renal carcinoma antigen NY-REN-19

基因 ID (Gene ID) 6794.0

Q15831.The antiserum was produced against synthesized peptide derived

蛋白ID (SwissProt ID) from human LKB1 around the phosphorylation site of Thr189. AA range:155-

204

### 产品应用 (Application)

稀释比 (Dilution Ratio) WB 1:500-1:2000,ELISA 1:5000-1:10000

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**蛋白分子量 (Molecular Weight)** 65kDa

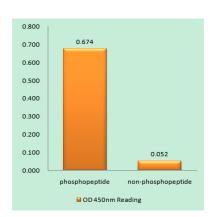
### 研究背景 (Background)

This gene, which encodes a member of the serine/threonine kinase family, regulates cell polarity and functions as a tumor suppressor. Mutations in this gene have been associated with Peutz-Jeghers syndrome, an autosomal dominant disorder characterized by the growth of polyps in the gastrointestinal tract, pigmented macules on the skin and mouth, and other neoplasms. Alternate transcriptional splice variants of this gene have been observed but have not been thoroughly characterized. [provided by RefSeq, Jul 2008],catalytic activity:ATP + a protein = ADP + a phosphoprotein., cofactor: Magnesium or manganese., disease: Defects in STK11 are a cause of Peutz-Jeghers syndrome (PJS) [MIM:175200]. PJS is a rare hereditary disease in which there is predisposition to benign and malignant tumors of many organ systems. PJS is an autosomal dominant disorder characterized by melanocytic macules of the lips, multiple gastrointestinal hamartomatous polyps and an increased risk for various neoplasms, including gastrointestinal cancer., disease: Defects in STK11 have been associated with testicular tumors [MIM:273300]. It includes germ cell tumor (GCT) or testicular germ cell tumor (TGCT), enzyme regulation: Activated by binding of a complex consisting of CAB39 and STRAD or CAB39 and ALS2CR2, function: Essential role in G1 cell cycle arrest. Phosphorylates and activates members of the AMPK-related subfamily of protein kinases. Tumor suppressor., online information: PJS entry, PTM: Phosphorylated by a cAMP-dependent protein kinase., similarity: Belongs to the protein kinase superfamily,, similarity: Belongs to the protein kinase superfamily, CAMK Ser/Thr protein kinase family, LKB1 subfamily, similarity; Contains 1 protein kinase domain.,subcellular location:Relocates to the cytoplasm when bound to CAB39 and STRAD or CAB39 and ALS2CR2., subunit: Found in a ternary complex composed of SMAD4, STK11 and STK11IP. Interacts with SMAD4 and STK11IP., tissue specificity: Ubiquitously expressed. Strongest expression in testis and fetal liver.,

#### 研究领域(Research Area)

Insulin Receptor; mTOR; AMPK

# 图片 (Image Data)



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-

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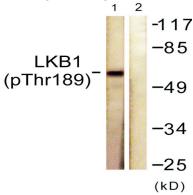
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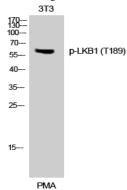






Western blot analysis of lysates from NIH/3T3 cells treated with PMA 125ng/ml 30 ', using LKB1 (Phospho-Thr189)

Antibody. The lane on the right is blocked with the phospho peptide.



Western Blot analysis of 3T3 cells using Phospho-LKB1 (T189) Polyclonal Antibody

#### 注意事项 (Note)

For research use only.

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