产品名称: Fibulin-3 Rabbit Polyclonal Antibody

产品货号: APRab10979



### 产品概述 (Summary)

产品名称 (Production Name) Fibulin-3 Rabbit Polyclonal Antibody

描述 (Description) Rabbit polyclonal Antibody

宿主 (Host) Rabbit

应用 (Application)WB,IHC,ICC/IF,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) EFEMP1

EFEMP1; FBLN3; FBNL; EGF-containing fibulin-like extracellular matrix protein **别名 (Alternative Names)** 

1; Extracellular protein S1-5; Fibrillin-like protein; Fibulin-3; FIBL-3

基因 ID (Gene ID) 2202.0

Q12805.The antiserum was produced against synthesized peptide derived 蛋白ID (SwissProt ID)

from human EFEMP1. AA range:111-160

# 产品应用 (Application)

**稀释比(Dilution Ratio)** WB 1:500-1:2000,IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:5000-1:20000

**蛋白分子量 (Molecular Weight)** 55kDa

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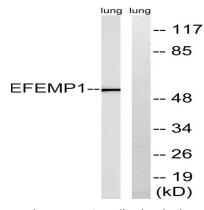
#### 研究背景 (Background)

This gene encodes a member of the fibulin family of extracellular matrix glycoproteins. Like all members of this family, the encoded protein contains tandemly repeated epidermal growth factor-like repeats followed by a C-terminus fibulin-type domain. This gene is upregulated in malignant gliomas and may play a role in the aggressive nature of these tumors. Mutations in this gene are associated with Doyne honeycomb retinal dystrophy. Alternatively spliced transcript variants that encode the same protein have been described.[provided by RefSeq, Nov 2009], alternative products: Experimental confirmation may be lacking for some isoforms, disease: Defects in EFEMP1 are a cause of Doyne honeycomb retinal dystrophy (DHRD) [MIM:126600]; also known as malattia leventinese (MLVT OR ML). DHRD is an autosomal dominant disease characterized by yellow-white deposits known as drusen that accumulate beneath the retinal pigment epithelium., online information: Retina International's Scientific Newsletter, similarity: Belongs to the fibulin family, similarity: Contains 6 EGF-like domains.,

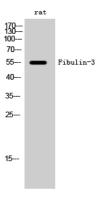
### 研究领域 (Research Area)

Signal; Signaling Pathway; Calcium Signaling; Calcium Binding Proteins; Neuroscience; Sensory System; Visual system

### 图片 (Image Data)



Western blot analysis of lysates from rat lung, using EFEMP1 Antibody. The lane on the right is blocked with the synthesized peptide.



Western Blot analysis of rat cells using Fibulin-3 Polyclonal Antibody

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# 注意事项 (Note)

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