

## 丝氨酸蛋白酶 56 抗体

产品货号: mlR19444

英文名称: PRSS56

中文名称: 丝氨酸蛋白酶 56 抗体

别 名: MCOP6; Protease serine 56; PRS56\_HUMAN; Putative serine protease 56; Serine protease 56.

研究领域: 细胞生物 发育生物学 神经生物学 干细胞

抗体来源: Rabbit

克隆类型: Polyclonal

交叉反应: Human,

产品应用: ELISA=1:500-1000 IHC-P=1:400-800 IHC-F=1:400-800 ICC=1:100-500 IF=1:100-500 (石蜡切片需

做抗原修复)

not yet tested in other applications.

optimal dilutions/concentrations should be determined by the end user.

分子量: 63kDa

细胞定位: 细胞浆

性 状: Lyophilized or Liquid

浓 度: 1mg/ml

免疫原: KLH conjugated synthetic peptide derived from human PRSS56:401-500/603



亚型: IgG

纯化方法: affinity purified by Protein A

储存液: 0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.

保存条件: Store at -20  $^{\circ}$  C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 $^{\circ}$  C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4  $^{\circ}$  C.

PubMed: PubMed

产品介绍: This gene encodes a protein that contains a peptidase S1 domain and possesses trypsin-like serine protease activity. The encoded protein may play a role in eye development, and mutations in this gene are a cause of autosomal recessive posterior microphthalmos. [provided by RefSeq, Dec 2011]

## **Subcellular Location:**

endoplasmic reticulum

## **Tissue Specificity:**

Expressed neural retina, cornea, sclera and optic nerve.

## DISEASE:

A developmental ocular disorder characterized by small malformed eyes. Clinical features are extreme hyperopia due to short axial length with essentially normal anterior segment, steep corneal curvatures, shallow anterior chamber, thick lenses, and thickened scleral wall. Palpebral fissures appear narrow because of relatively deep-set eyes, visual acuity is mildly to moderately reduced, and anisometropic or strabismic amblyopia is common. The fundus of the eye shows crowded optical disks, tortuous vessels, and an abnormal foveal avascular zone.

SWISS:



applications.

POCW18		
Gene ID:		
646960		
Important Note:		

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic