

# Rabbit Anti-GRK1 antibody

# SL1082R

GRK1
G蛋白偶合受体激酶1抗体
G-protein coupled receptor kinase 1; GRK1; GPRK1; RK; Grk1; Rhok; RHODOPSIN KINASE; RK_HUMAN.
Rabbit
Polyclonal
Human, Mouse, Rat, Chicken, Cow,
WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
62kDa
The cell membrane
Lyophilized or Liquid
lmg/ml
KLH conjugated synthetic peptide derived from human GRK1:201-300/563
IgG
affinity purified by Protein A
0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Store at -20 °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 °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 °C.
<u>PubMed</u>
This gene encodes a member of the guanine nucleotide-binding protein (G protein)-coupled receptor kinase subfamily of the Ser/Thr protein kinase family. The protein phosphorylates rhodopsin and initiates its deactivation. Defects in GRK1 are known to cause Oguchi disease 2 (also known as stationary night blindness Oguchi type-2). [provided by RefSeq]

#### **Function:**

Phosphorylates rhodopsin thereby initiating its deactivation. This rapid desensitization is essential for scotopic vision and permits rapid adaptation to changes in illumination.

## **Subcellular Location:**

Membrane.

# Tissue Specificity:

Retina and pineal gland.

#### Post-translational modifications:

Autophosphorylated.

Farnesylation is required for full activity.

#### **DISEASE:**

Defects in GRK1 are a cause of congenital stationary night blindness Oguchi type 2 (CSNBO2) [MIM:613411]. It is non-progressive retinal disorder characterized by impaired night vision, often associated with nystagmus and myopia. Congenital stationary night blindness Oguchi type is associated with fundus discoloration and abnormally slow dark adaptation.

## Similarity:

Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. GPRK subfamily.

Contains 1 AGC-kinase C-terminal domain.

Contains 1 protein kinase domain.

Contains 1 RGS domain.

## **SWISS:**

O15835

#### Gene ID:

6011

#### Database links:

Entrez Gene: 6011 Human

Entrez Gene: 24013 Mouse

Omim: 180381 Human

SwissProt: Q15835 Human

SwissProt: O9WVL4 Mouse

Unigene: 103501 Human

Unigene: 721727 Human

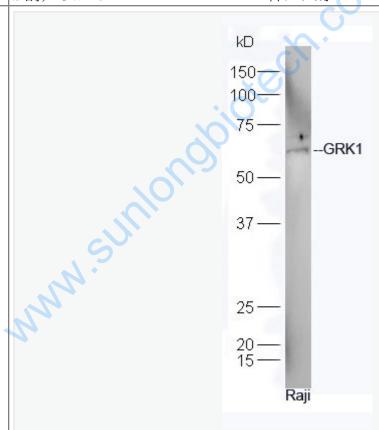
Unigene: 257501 Mouse

# **Important Note:**

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

# G protein-coupled

receptor激酶1(GRK1)是催化激动剂诱导的GPCR磷酸化以及启动GPCR脱敏的关键激酶,主要用于GRKThe cell membrane转位和对GRK功能的影响.



Picture:

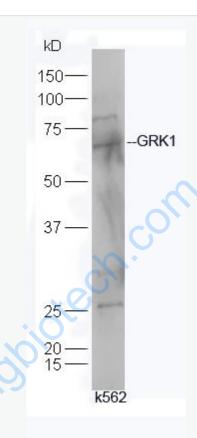
Sample: Raji Cell Lysate at 40 ug

Primary: Anti-GRK1 (SL1082R) at 1:300 dilution;

Secondary: HRP conjugated Goat-Anti-rabbit IgG(SL1082R) at 1: 5000 dilution;

Predicted band size: 62 kD





Sample:

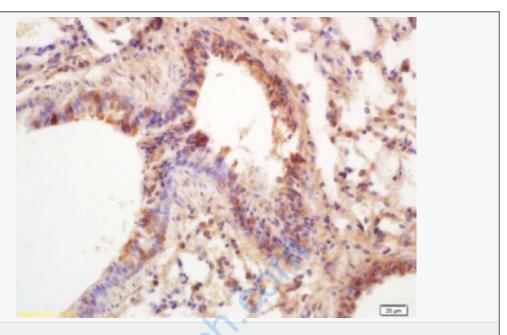
K562 Cell (Human) Lysate at 30 ug

Primary: Anti-GRK1 (SL1082R) at 1/300 dilution

Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution

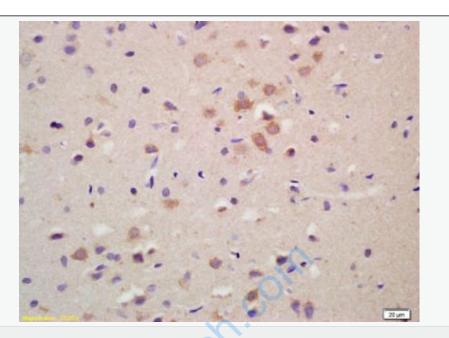
Predicted band size: 62 kD

Observed band size: 65 kD



Tissue/cell: mouse lung tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min;

Incubation: Anti-GRK1 Polyclonal Antibody, Unconjugated(SL1082R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0 ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min;

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