

GRK 1 Polyclonal Antibody

Catalog No: YT2063

Reactivity: Human; Mouse; Rat

Applications: WB;IHC;IF;ELISA

Target: GRK1

Fields: >>Chemokine signaling pathway;>>Endocytosis;>>Phototransduction

Gene Name: GRK1

Protein Name: Rhodopsin kinase

Q15835

Q9WVL4

Human Gene Id: 6011

Human Swiss Prot

No:

Mouse Swiss Prot

No:

Rat Gene ld: 81760

Rat Swiss Prot No: Q63651

Immunogen: The antiserum was produced against synthesized peptide derived from human

GRK1. AA range:6-55

Specificity: GRK 1 Polyclonal Antibody detects endogenous levels of GRK 1 protein.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution : WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:40000.. IF 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.



Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 63kD

Cell Pathway: Chemokine; Endocytosis;

Background: 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, Jul 2008],

Function : catalytic activity:ATP + [rhodopsin] = ADP + [rhodopsin]

phosphate., disease: Defects in GRK1 are a cause of congenital stationary night blindness Oguchi type (CSNBO) [MIM:258100]; also known as Oguchi disease.

Congenital stationary night blindness is a non-progressive retinal disorder characterized by impaired night vision. CSNBO is an autosomal recessive form

associated with fundus discoloration and abnormally slow dark adaptation.,function:Phosphorylates rhodopsin thereby initiating its deactivation.,online information:Retina International's Scientific

Newsletter,PTM:Autophosphorylated.,PTM:Farnesylation is required for full activity.,similarity:Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. GPRK subfamily.,similarity:Contains 1 AGC-kinase C-terminal domain.,similarity:Contains 1 protein kinase domain.,similarity:Contains 1 RGS

domain.,tissue specificity:R

Subcellular Location :

Membrane; Lipid-anchor. Cell projection, cilium, photoreceptor outer segment.

Subcellular location is not affected by light or dark conditions. .

Expression: Retinal-specific. Expressed in rods and cones cells.

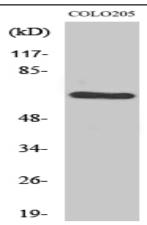
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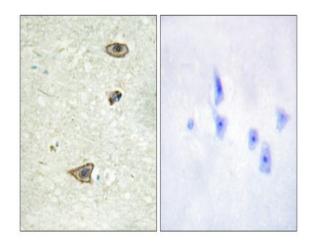
Host: Rabbit

Modifications: Unmodified

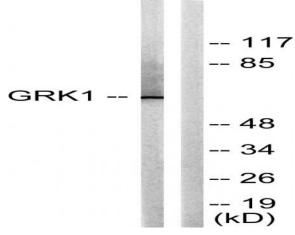
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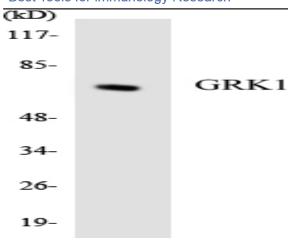
Western Blot analysis of various cells using GRK 1 Polyclonal Antibody



Immunohistochemistry analysis of paraffin-embedded human brain tissue, using GRK1 Antibody. The picture on the right is blocked with the synthesized peptide.



Western blot analysis of lysates from COLO205 cells, using GRK1 Antibody. The lane on the right is blocked with the synthesized peptide.



Western blot analysis of the lysates from Jurkat cells using GRK1 antibody.