# Rhodopsin Rabbit mAb

Catalog No: #49524

Package Size: #49524-1 50ul #49524-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

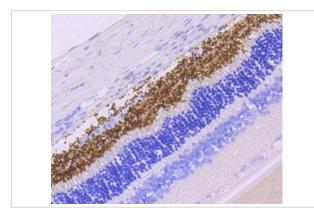
## Description

Product Name	Rhodopsin Rabbit mAb
Host Species	Recombinant Rabbit
Clonality	Monoclonal
Clone No.	JA10-91
Purification	ProA affinity purified
Applications	WB, IHC
Species Reactivity	Hu, Rt
Immunogen Description	recombinant protein
Conjugates	Unconjugated
Other Names	CSNBAD1 antibody MGC138309 antibody MGC138311 antibody OPN 2 antibody OPN2 antibody opsd
	antibody OPSD_HUMAN antibody opsin 2 antibody Opsin 2 rod pigment antibody Opsin-2 antibody Opsin-2
	antibody Retinitis Pigmentosa 4 antibody Retinitis pigmentosa 4 autosomal dominant antibody RHO
	antibody Rhodopsin antibody RP 4 antibody RP4 antibody
Accession No.	Swiss-Prot#:P08100
Calculated MW	39 kDa
Formulation	1*TBS (pH7.4), 1%BSA, 40%Glycerol. Preservative: 0.05% Sodium Azide.
Storage	Store at -20°C

## **Application Details**

WB: 1:500-1:1,000 IHC: 1:50-1:200

#### **Images**



Immunohistochemical analysis of paraffin-embedded rat retina tissue using anti-Rhodopsin antibody. Counter stained with hematoxylin.

#### Background

In aerobic respiration reactions, succinate dehydrogenase (SDH) catalyzes the oxidation of succinate and ubiquinone to fumarate and ubiquinol. Four subunits comprise the SDH protein complex: a flavochrome subunit (SDHA), an iron-sulfur protein (SDHB), and two membrane-bound subunits

(SDHC and SDHD) anchored to the inner mitochondrial membrane. Mutations to these subunits cause mitochondrial dysfunction, corresponding to several distinct disorders. Mutations in the membrane bound components may cause hereditary paraganglioma, while SDHA mutations are associated with juvenile encephalopathy as well as Leigh Syndrome, a severe neurological disorder. Inactivating mutations in SDHB correlate with inherited, but not necessarily sporadic, cases of pheochromocytoma.

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Note: This product is for in vitro research use only and is not intended for use in humans or animals.