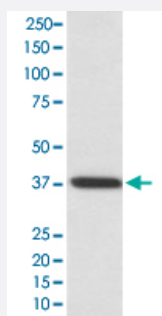


RHO monoclonal antibody, clone AADI-18

Catalog # MAB20675 Size 100 uL

Applications



Western Blot (Tissue lysate)

Western Blot analysis of rat eyeball tissue lysate with RHO monoclonal antibody, clone AADI-18 (Cat # MAB20675).

Specification

Product Description Rabbit monoclonal antibody raised against synthetic peptide of human RHO.

Immunogen A synthetic peptide corresponding to human RHO.

Host Rabbit

Theoretical MW (kDa) 38.893

Reactivity Human

Form Liquid

Purification Affinity purification

Isotype IgG

Recommend Usage
Immunohistochemistry (1:50-1:200)
Western Blot (1:500-1:2000)
The optimal working dilution should be determined by the end user.

Storage Buffer In PBS, 150 mM NaCl, pH 7.4 (50% glycerol, 0.02% sodium azide).

Storage Instruction

Store at -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for a longer time. Avoid repeated freezing and thawing.

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot (Tissue lysate)

Western Blot analysis of rat eyeball tissue lysate with RHO monoclonal antibody, clone AADI-18 (Cat # MAB20675).

- Immunohistochemistry

Gene Info — RHO

Entrez GeneID[6010](#)**Protein Accession#**[P08100](#)**Gene Name**

RHO

Gene Alias

CSNBAD1, MGC138309, MGC138311, OPN2, RP4

Gene Description

rhodopsin

Omim ID[180380](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

Retinitis pigmentosa is an inherited progressive disease which is a major cause of blindness in western communities. It can be inherited as an autosomal dominant, autosomal recessive, or X-linked recessive disorder. In the autosomal dominant form, which comprises about 25% of total cases, approximately 30% of families have mutations in the gene encoding the rod photoreceptor-specific protein rhodopsin. This is the transmembrane protein which, when photoexcited, initiates the visual transduction cascade. Defects in this gene are also one of the causes of congenital stationary night blindness. [provided by RefSeq]

Other Designations

opsin 2, rod pigment|retinitis pigmentosa 4, autosomal dominant|rhodopsin (opsin 2, rod pigment) (retinitis pigmentosa 4, autosomal dominant)

Disease

- [Genetic Diseases](#)

- [Retinal Degeneration](#)
- [Retinal Diseases](#)
- [Retinitis Pigmentosa](#)
- [Vision Disorders](#)