



Rabbit Anti-Retinal S antigen antibody

SL11996R

Product Name Retinal S antigen

Chinese Name 视网膜 S 抗原抗体

Alias S-antigen; 48 kDa protein; Arrestin 1; Arrestin; ARRS_HUMAN; DKFZp686I1383; Retinal S antigen (protein); Retinal S-antigen; Rod photoreceptor arrestin; RP47; S AG; S antigen; S antigen retina and pineal gland (arrestin); S antigen retina and pineal gland; S arrestin; S-AG; S-arrestin; SAG.

Research Area Cell biology Neurobiology Signal transduction G protein-coupled receptor G protein signal

Immunogen Species Rabbit

Clonality Polyclonal

React Species (predicted: Human, Mouse, Rat, Dog, Cow, Rabbit, Sheep,)

WB=1:500-2000,IHC-P=1:100-500,IHC-F=1:100-500,ICC/IF=1:100-500,IF=1:100-500,ELISA
(Paraffin sections need antigen repair)

Applications not yet tested in other applications.
optimal dilutions/concentrations should be determined by the end user.

Theoretical molecular weight 45kDa

Cellular localization The cell membrane

Form Liquid

Concentration 1mg/ml

immunogen KLH conjugated synthetic peptide derived from human Retinal S antigen: 285-330/405

Lsotype IgG

Purification affinity purified by Protein A

Buffer Solution 1M TBS(pH7.4) with 1% BSA, 3% Proclin300 and 50% Glycerol.

Storage Shipped at 4°C. Store at -20 °C for one year. Avoid repeated freeze/thaw cycles.

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

PubMed [PubMed](#)

Members of arrestin/beta-arrestin protein family are thought to participate in agonist-mediated deactivation of G-protein-coupled receptors and cause specific dampening of cellular responses to stimuli such as neurotransmitters, or sensory signals. S-arrestin, also known as S-antigen, is a major soluble photoreceptor protein that is involved in desensitization of the photoactivated transduction cascade. It is expressed in the retina and the pineal gland and inhibits coupling of rhodopsin to transducin in vitro. Additionally, S-arrestin is immunogenic, and is capable of inducing experimental autoimmune uveoretinitis. Mutations in this gene are associated with Oguchi disease, a rare autosomal recessive form of night blindness. [provided by RefSeq, Jul 2008]

Function:

Arrestin is one of the major proteins of the ROS (retinal rod outer segments); it binds to photoactivated-phosphorylated rhodopsin, thereby apparently preventing the transducin-mediated activation of phosphodiesterase.

Tissue Specificity:

Retina and pineal gland.

DISEASE:

Defects in SAG are a cause of congenital stationary night blindness Oguchi type 1 (CSNBO1) [MIM:264500], also known as Oguchi disease. Congenital stationary night blindness is a non-progressive retinal disease characterized by impaired night vision. CSNBO is an autosomal recessive form associated with rod dysfunction, retinal discoloration and abnormally slow dark adaptation.

**Product
Detail**

Similarity:

Belongs to the arrestin family.

SWISS:

P10523

Gene ID:

6295

Database links:

[Entrez Gene: 6295](#) Human

[Entrez Gene: 20215](#) Mouse

[Entrez Gene: 280922](#) Cow

[Entrez Gene: 25539](#) Rat

[Omim: 181031](#) Human



[SwissProt: P08168](#) Cow

[SwissProt: P10523](#) Human

[SwissProt: P20443](#) Mouse

[SwissProt: P15887](#) Rat

[Unigene: 32317](#) Cow

[Unigene: 32721](#) Human

[Unigene: 1276](#) Mouse

[Unigene: 9856](#) Rat