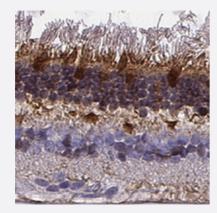


GUCA1A polyclonal antibody

Catalog # PAB31073 Size 100 uL

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemical staining of human retina shows strong cytoplasmic positivity in photoreceptor layer and outer plexiform layer.

Specification	
Product Description	Rabbit polyclonal antibody raised against partial recombinant human GUCA1A.
Immunogen	Recombinant protein corresponding to human GUCA1A.
Sequence	EYVAALSLVLKGKVEQKLRWYFKLYDVDGNGCIDRDELLTIIQAIRAINPCSDTTMTAEEFTDTVFSK IDVNGDGELSLEEFIEGVQKDQMLLDTLTRSLDLTRIVRRLQNGEQDEEGADEAAEAAG
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Antigen affinity purification
Isotype	lgG
Recommend Usage	Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (1:5000-10000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide).



Product Information

Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

• Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human retina shows strong cytoplasmic positivity in photoreceptor layer and outer plexiform layer.

Gene Info — GUCA1A	
Entrez GenelD	<u>2978</u>
Protein Accession#	<u>P43080</u>
Gene Name	GUCA1A
Gene Alias	COD3, GCAP, GCAP1, GUCA, GUCA1
Gene Description	guanylate cyclase activator 1A (retina)
Omim ID	<u>600364</u> <u>602093</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene plays a role in the recovery of retinal photoreceptors from photobleaching. In the recove ry phase, the phototransduction messeneger cGMP is replenished by retinal guanylyl cyclase-1 (G C1). GC1 is activated by decreasing Ca(2+) concentrations following photobleaching. The protein encoded by this gene, guanylyl cyclase activating protein 1 (GCAP1), mediates the sensitivity of GC1 to Ca(2+) concentrations. GCAP1 promotes activity of GC1 at low Ca(2+) concentrations and inhibits GC1 activity at high Ca(2+) concentrations. Mutations in this gene cause autosomal do minant cone dystrophy (COD3); a disease characterized by reduced visual acuity associated with progressive loss of color vision. Mutations in this gene prohibit the inactivation of RetGC1 at high Ca(2+) concentrations; causing the constitutive activation of RetGC1 and, presumably, increased cell death. This gene is expressed in retina and spermatagonia. [provided by RefSeq
Other Designations	OTTHUMP0000016397 OTTHUMP00000196466

Pathway

Olfactory transduction



Disease

- Retinal Degeneration
- Retinal Diseases