



Gα t1 Monoclonal Antibody

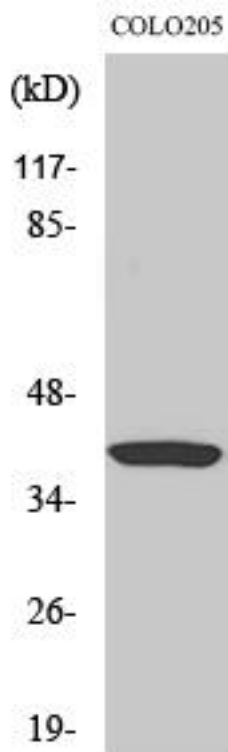
Catalog No	YP-mAb-16174
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	GNAT1
Immunogen	The antiserum was produced against synthesized peptide derived from human GNAT1. AA range:71-120
Specificity	Gα t1 Monoclonal Antibody detects endogenous levels of Gα t1 protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	GDNF family receptor alpha 1;GDNF family receptor alpha-1;GDNF R;GDNF RA; GDNF receptor alpha;GDNF receptor alpha-1;GDNFR alpha 1;GDNFR alpha; GDNFR;GDNFR-alpha-1;GDNFRA;GDNFRalpha;GFR alpha 1;GFR alpha1; GFR-alpha-1;GFRA 1;Gfra1;GFRA1_HUMAN;GFRalpha1;Glial cell line derived neurotrophic factor receptor alpha;GPI linked anchor protein;MGC2345;PI linked cell surface accessory protein;RET 1L;RET ligand 1;RET1L;RETL 1;RETL1;TGF beta related neurotrophic factor receptor 1;TGF-beta-related neurotrophic factor receptor 1;TRNR 1;TRNR1;Gα t1;GNAT1;Guanine nucleotide-binding protein G(t) subunit alpha-1;GNATR;Transducin alpha-1 chain
Observed Band	36kD
Calculated Molecular Weight	36kD
Cell Pathway	Cell projection, cilium, photoreceptor outer segment . Membrane ; Peripheral membrane protein . Photoreceptor inner segment . Localizes mainly in the outer segment in the dark-adapted state, whereas is translocated to the inner part of the photoreceptors in the light-adapted state. During dark-adapted conditions, in the presence of UNC119 mislocalizes from the outer segment to the inner part of rod photoreceptors which leads to decreased photoreceptor damage caused by light. .



Tissue Specificity	Rod photoreceptor cells (PubMed:1614872). Predominantly expressed in the retina followed by the ciliary body, iris and retinal pigment epithelium (PubMed:22190596).
Function	disease:Defects in GNAT1 are the cause of congenital stationary night blindness autosomal dominant type 3 (CSNBAD3) [MIM:610444]; also known as congenital stationary night blindness Nougaret type. Congenital stationary night blindness is a non-progressive retinal disorder characterized by impaired night vision.,function:Guanine nucleotide-binding proteins (G proteins) are involved as modulators or transducers in various transmembrane signaling systems. Transducin is an amplifier and one of the transducers of a visual impulse that performs the coupling between rhodopsin and cGMP-phosphodiesterase.,online information:Retina International's Scientific Newsletter,similarity:Belongs to the G-alpha family. G(i/o/t/z) subfamily.,subunit:G proteins are composed of 3 units; alpha, beta and gamma. The alpha chain contains the guanine nucleotide binding site.,tissue specificity:Rod.,
Background	Transducin is a 3-subunit guanine nucleotide-binding protein (G protein) which stimulates the coupling of rhodopsin and cGMP-phosphodiesterase during visual impulses. The transducin alpha subunits in rods and cones are encoded by separate genes. This gene encodes the alpha subunit in rods. This gene is also expressed in other cells, and has been implicated in bitter taste transduction in rat taste cells. Mutations in this gene result in autosomal dominant congenital stationary night blindness. Multiple alternatively spliced variants, encoding the same protein, have been identified. [provided by RefSeq, Feb 2009],
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.



Products Images



Western Blot analysis of various cells using Ga t1
Monoclonal Antibody