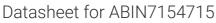
antibodies -online.com





anti-GNAT2 antibody (AA 2-139)





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Overview

Quantity:	100 μg
Target:	GNAT2
Binding Specificity:	AA 2-139
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GNAT2 antibody is un-conjugated
Application:	ELISA, Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Guanine nucleotide-binding protein G(t) subunit alpha-2 protein (2-139AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

Target:	GNAT2
Alternative Name:	GNAT2 (GNAT2 Products)
Background:	Background: 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.

Aliases: ACHM4 antibody, Cone type transducin alpha subunit antibody, GNAT 2 antibody, GNAT C antibody, Gnat2 antibody, GNAT2_HUMAN antibody, GNATC antibody, Guanine nucleotide binding protein (G protein) alpha transducing activity polypeptide 2 antibody, Guanine nucleotide binding protein G t subunit alpha 2 antibody, Guanine nucleotide-binding protein G(t) subunit alpha-2 antibody, Transducin alpha 2 antibody, Transducin alpha-2 chain antibody, Transducin alpha-2 antibody, Transducin cone specific alpha polypeptide antibody

UniProt: P19087

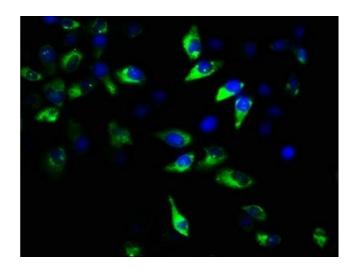
Pathways: G-protein mediated Events, Phototransduction

Application Details

Application Notes:	Recommended dilution: IF:1:50-1:200,
Restrictions:	For Research Use only

Handling

Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.



Immunofluorescence

Image 1. Immunofluorescence staining of Hela cells with ABIN7154715 at 1:167, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).