

## ARHGAP32 Antibody

PACO61622

### Description

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This ARHGAP32 Antibody is supplied as a kit for advanced applications. The kit includes Bradford Reagent to quantify total protein concentration for accurate sample normalization (Optional).

### Product Information

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<b>SKU:</b>	PACO61622
<b>Contents:</b>	50µg Bradford Reagent: 1 vial (2ml)
<b>Category:</b>	-
<b>Synonyms:</b>	ARHGAP32 antibody, GRIT antibody, KIAA0712 antibody, RICS antibody, Rho GTPase-activating protein 32 antibody, Brain-specific Rho GTPase-activating protein antibody, GAB-associated Cdc42/Rac GTPase-activating protein antibody, GC-GAP antibody, GTPase regulator interacting with TrkA antibody, Rho-type GTPase-activating protein 32 antibody, Rho/Cdc42/Rac GTPase-activating protein RICS antibody, RhoGAP involved in the beta-catenin-N-cadherin and NMDA receptor signaling antibody, p200RhoGAP antibody, p250GAP antibody
<b>Clone:</b>	Polyclonal
<b>Applications:</b>	<b>ELISA</b> <b>IF</b>
<b>Conjugation:</b>	Non-conjugated
<b>Reactivity:</b>	Human

### Antibody Data

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<b>Isotype:</b>	IgG
<b>Uniprot:</b>	A7KAX9
<b>Host Species:</b>	Rabbit
<b>Purification:</b>	>95%, Protein G purified
<b>Immunogen:</b>	Recombinant Human Rho GTPase-activating protein 32 protein (130-205AA)
<b>Immunogen Species:</b>	Homo sapiens (Human)

**Manufacturers Statement:** This final kit system is assembled and quality-released by Assay Genie Limited.

**Buffer:** Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

**Form:** Liquid

## Preparation & Storage

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**Storage:** Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.  
Store Bradford Reagent at Room Temperature for 1 Year.

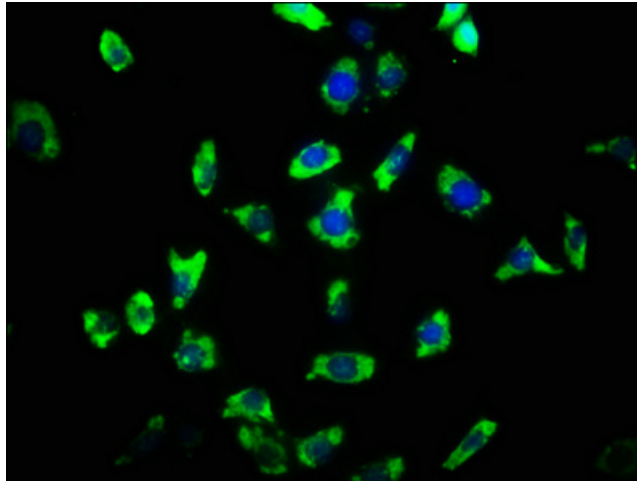
Recommended Dilutions:	Application	Recommended Dilution
	IF	1:20-1:200

**Protein Quantification (Optional):** To quantify total protein levels, use the Bradford Reagent included in this kit. Visit <https://www.assaygenie.com/bradford-protein-assay-protocol/> to view the full protocol

## Validation Data

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**Image**



**Description**

Immunofluorescence staining of SH-SY5Y cells with PACO61622 at 1:33, 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-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).