# SANTA CRUZ BIOTECHNOLOGY, INC.

# GOA-1 (cF-17): sc-32686



BACKGROUND

GOA-1, the  $G_{\alpha 0}$  subunit of a heterotrimeric G protein complex, is involved in asymmetric cell division in *C. elegans*. Specifically, GOA-1 assists in the positioning of spindle fibers in one-celled *C. elegans* embryos. Proper spindle fiber positioning also depends upon the interaction of GOA-1 in its GDP bound form to the receptor independent G protein activators GPR-1 and GPR-2, as well as the protein RIC-8, a guanine nucleotide exchange factor (GEF). GOA-1 is localized to the cell cortex and is found in most neurons. Additionally, muscles involved with male mating and egg laying are sites of GOA-1 expression in *C. elegans*. Mutations in the gene coding for GOA-1 are associated with a variety of deficits mimicking serotonin deficiency in *C. elegans*, including male impotence, premature egg laying and hyperactive movement. EGL-10 is an regulator of G protein signaling (RGS) protein that can selectively inhibit GOA-1 function through activation of G<sub>α</sub> GTPase.

#### REFERENCES

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- 2. Mendel, J.E., et al. 1995. Participation of the protein  $G_0$  in multiple aspects of behavior in *C. elegans*. Science 267: 1652-1655.
- 3. Miller, K.G., et al. 1999.  $G_{\alpha\alpha}$  and diacylglycerol kinase negatively regulate the  $G_{\alpha\alpha}$  pathway in *C. elegans*. Neuron 24: 323-333.
- 4. Miller, K.G., et al. 2000. A role for RIC-8 (Synembryn) and GOA-1 ( $G_{o\alpha}$ ) in regulating a subset of centrosome movements during early embryogenesis in *Caenorhabditis elegans*. Genetics 156: 1649-1660.
- 5. van Swinderen, B., et al. 2001.  $G_{0\alpha}$  regulates volatile anesthetic action in *Caenorhabditis elegans*. Genetics 158: 643-655.
- 6. Robatzek, M., et al. 2001. EAT-11 encodes GPB-2, a  $G_{\beta5}$  ortholog that interacts with  $G_{0\alpha}$  and  $G_{q\alpha}$  to regulate *C. elegans* behavior. Curr. Biol. 11: 288-293.
- 7. Gotta, M., et al. 2001. Distinct roles for  $G_{\alpha}$  and  $G_{\beta\gamma}$  in regulating spindle position and orientation in *Caenorhabditis elegans* embryos. Nat. Cell. Biol. 3: 297-300.
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- Manning, D.R., et al. 2003. Evidence mounts for receptor-independent activation of heterotrimeric G proteins normally *in vivo*: positioning of the mitotic spindle in *C. elegans*. Sci. STKE. 2003: 35.

## SOURCE

GOA-1 (cF-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of GOA-1 of *C. elegans* origin.

# STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## PRODUCT

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-32686 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

### **APPLICATIONS**

GOA-1 (cF-17) is recommended for detection of GOA-1 of *Caenorhabditis elegans* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Molecular Weight of GOA-1: 40 kDa.

## **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluo-rescence: use donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

#### PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.