Rim2 (C-14): sc-98109



The Power to Question

BACKGROUND

Rab3, a neural/neuroendocrine-specific member of the Rab family, is involved in Ca²⁺-regulated exocytosis. Rab3 functions in an inhibitory capacity by controlling the recruitment of secretory vesicles into a releasable pool at the plasma membrane. Rim (rab3 interacting molecule), a putative effector protein for Rab3s, is composed of an N-terminal zinc-finger motif and C-terminal PDZ and C2 domains. Rim exists as two variants, Rim1 and Rim2, produced by alternative splicing. The 3'-end of the Rim2 gene produces an independent mRNA that encodes a smaller protein referred to as Nim2, which like Rim, also regulates exocytosis. Rim serves as a Rab3-dependent regulator of synapticvesicle fusion by forming a GTP-dependent complex between synaptic plasma membranes and docked synaptic vesicles. Both Rim1 and Rim2 can bind to cAMP-GEFII, which is a direct target of cAMP in regulated exocytosis and is responsible for cAMP-dependent, PKA-dependent exocytosis. Rim also localizes on the plasma membrane of INS-1E cells and pancreatic β-cells. Rab3 binding domain of Rim enhances glucose-stimulated secretion in intact cells and Ca²⁺-stimulated exocytosis in permeabilized cells, suggesting that Rim may also play a regulatory role in Insulin secretion.

REFERENCES

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- Coppola, T., et al. 1999. Disruption of Rab3-calmodulin interaction, but not other effector interactions, prevents Rab3 inhibition of exocytosis. EMBO J. 18: 5885-5891.
- Ozaki, N., et al. 2000. cAMP-GEFII is a target of cAMP in regulated exocytosis. Nat. Cell Biol. 2: 805-811.
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- 5. lezzi, M., et al. 2000. The Rab3-interacting molecule RIM is expressed in pancreatic beta-cells and is implicated in Insulin exocytosis. FEBS Letts. 474: 66-70.
- Haynes, L.P., et al. 2001. A direct inhibitory role for the Rab3-specific effector, Noc2, in Ca²⁺-regulated exocytosis in neuroendocrine cells. J. Biol. Chem. 27: 9726-9732.

CHROMOSOMAL LOCATION

Genetic locus: RIMS2 (human) mapping to 8g22.3.

SOURCE

Rim2 (C-14) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of Rim2 of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-98109 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

Rim2 (C-14) is recommended for detection of Rim2 of human 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); non cross-reactive with isoforms 2, RimL3a or RimL3c.

Suitable for use as control antibody for Rim2 siRNA (h): sc-77790, Rim2 shRNA Plasmid (h): sc-77790-SH and Rim2 shRNA (h) Lentiviral Particles: sc-77790-V.

Molecular Weight of Rim2: 160 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) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

STORAGE

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

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.



Try **Rim2 (63-M7): sc-100842**, our highly recommended monoclonal alternative to Rim2 (C-14).

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