

ARHGEF37 (P-20): sc-246912

BACKGROUND

Rho GTPases, which play fundamental roles in numerous cellular processes, are initiated by external stimuli that signal through G-protein coupled receptors. ARHGEF37 (Rho guanine nucleotide exchange factor (GEF) 37) is a 675 amino acid protein that contains one BAR domain, one DH (DBL-homology) domain and 2 SH3 domains. Conserved in chimpanzee, canine, bovine, mouse, rat, chicken and zebrafish, ARHGEF37 is encoded by a gene that maps to human chromosome 5q32. Chromosome 5 makes up approximately 6% of the human genome and contains 181 million base pairs, which encode 1,000 genes. Chromosome 5 is associated with Cockayne syndrome, familial adenomatous polyposis and Treacher Collins syndrome. Deletion of 5q, or chromosome 5 altogether, is common in myelodysplastic syndrome and acute myelogenous leukemias.

REFERENCES

- Dixon, M.J., et al. 1991. The gene for Treacher Collins syndrome maps to the long arm of chromosome 5. *Am. J. Hum. Genet.* 49: 17-22.
- Saltman, D.L., et al. 1993. A physical map of 15 loci on human chromosome 5q23-q33 by two-color fluorescence *in situ* hybridization. *Genomics* 16: 726-732.
- Kadmon, M., et al. 2001. Duodenal adenomatosis in familial adenomatous polyposis coli. A review of the literature and results from the Heidelberg polyposis register. *Int. J. Colorectal Dis.* 16: 63-75.
- Marklund, L., et al. 2006. Adult-onset autosomal dominant leukodystrophy with autonomic symptoms restricted to 1.5 Mbp on chromosome 5q23. *Am. J. Med. Genet. B Neuropsychiatr. Genet.* 141B: 608-614.
- Aretz, S., et al. 2007. Somatic APC mosaicism: a frequent cause of familial adenomatous polyposis (FAP). *Hum. Mutat.* 28: 985-992.
- Cleaver, J.E., et al. 2007. Cockayne syndrome exhibits dysregulation of p21 and other gene products that may be independent of transcription-coupled repair. *Neuroscience* 145: 1300-1308.
- Mullighan, C.G., et al. 2008. Genomic analysis of the clonal origins of relapsed acute lymphoblastic leukemia. *Science* 322: 1377-1380.

CHROMOSOMAL LOCATION

Genetic locus: ARHGEF37 (human) mapping to 5q32; Arhgef37 (mouse) mapping to 18 E1.

SOURCE

ARHGEF37 (P-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of ARHGEF37 of human origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-246912 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

ARHGEF37 (P-20) is recommended for detection of ARHGEF37 of mouse, rat and 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).

ARHGEF37 (P-20) is also recommended for detection of ARHGEF37 in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for ARHGEF37 siRNA (h): sc-91898, ARHGEF37 siRNA (m): sc-140320, ARHGEF37 shRNA Plasmid (h): sc-91898-SH, ARHGEF37 shRNA Plasmid (m): sc-140320-SH, ARHGEF37 shRNA (h) Lentiviral Particles: sc-91898-V and ARHGEF37 shRNA (m) Lentiviral Particles: sc-140320-V.

Molecular Weight of ARHGEF37: 76 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) or donkey anti-goat IgG-TR: sc-2783 (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.