## BACKGROUND

Rho GTPases are molecular switches that regulate many essential cellular processes, including Actin dynamics, cell adhesion, cell-cycle progression and transcription. The Rho-type guanosine triphosphatase (GTPase), Cdc42, has been implicated in a variety of functions in the yeast life cycle, including septin organization for cytokinesis, pheromone response, haploid invasive growth and the establishment and maintenance of cell polarity. In yeast, the role of Cdc42 in polarization of cell growth includes polarization of the Actin cytoskeleton, which delivers secretory vesicles to growth sites at the plasma membrane. A group of proteins (Rga1, Rga2 and Bem3) called GTPase-activating proteins (GAPs) catalyze the hydrolysis of GTP to GDP, thereby inactivating Cdc42. Phosphorylation states of Cdc42 regulate its interaction with Rho GDP dissociation inhibitor and its extraction from biological membranes. Yeast Cdc42 functions at a late step in exocytosis, specifically during polarized growth of the emerging bud.

## REFERENCES

1. Adamo, J.E., Moskow, J.J., Gladfelter, A.S., Viterbo, D., Lew, D.J. and Brennwald, P.J. 2001. Yeast Cdc42 functions at a late step in exocytosis, specifically during polarized growth of the emerging bud. J. Cell Biol. 155: 581-592.
2. Forget, M.A., Desrosiers, R.R., Gingras, D. and Beliveau, R. 2002. Phosphorylation states of Cdc42 and RhoA regulate their interactions with Rho GDP dissociation inhibitor and their extraction from biological membranes. Biochem. J. 361: 243-254.
3. Smith, G.R., Givan, S.A., Cullen, P. and Sprague, G.F., Jr. 2002. GTPaseactivating proteins for Cdc42. Eukaryot. Cell 1: 469-480.
4. Hazan, I. and Liu, H. 2002. Hyphal tip-associated localization of Cdc42 is F-Actin dependent in Candida albicans. Eukaryot. Cell 1: 856-864.
5. Choi, S.C. and Han, J.K. 2002. Xenopus Cdc42 regulates convergent extension movements during gastrulation through $\mathrm{Wnt} / \mathrm{Ca}^{2+}$ signaling pathway. Dev. Biol. 244: 342-357.

## SOURCE

Cdc42 (yC-19) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C -terminus of Cdc42 of Saccharomyces cerevisiae origin.

## PRODUCT

Each vial contains $200 \mu \mathrm{ggG}$ in 1.0 ml of PBS with < $0.1 \%$ sodium azide and $0.1 \%$ gelatin.
Blocking peptide available for competition studies, sc-6793 P, ( $100 \mu \mathrm{~g}$ peptide in 0.5 ml PBS containing $<0.1 \%$ sodium azide and $0.2 \%$ BSA).

## STORAGE

Store at $4^{\circ} \mathrm{C},{ }^{* *}$ 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.

## APPLICATIONS

Cdc42 (yC-19) is recommended for detection of Cdc42 of Saccharomyces cerevisiae origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [ $1-2 \mu \mathrm{~g}$ per 100-500 $\mu \mathrm{g}$ of total protein ( 1 ml of cell lysate)) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).
Molecular Weight of Cdc42: 25 kDa .
Positive Controls: Cdc42 (h): 293T Lysate: sc-110467.

## 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 MarkerTM 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) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 ( 0.5 ml agarose/2.0 ml).

## DATA



Cdc42 (yC-19): sc-6793. Western blot analysis of Cdc42 expression in non-transfected: sc-117752 (A and human Cdc42 transfected: sc-110467 (B) 293T whole cell lysates and yeast recombinant $\operatorname{Cdc} 42$ (C)

## PROTOCOLS

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

