

# Rap1GAP (D-9): sc-166586

## BACKGROUND

Rap1 GTPase activating protein (Rap1GAP) specifically stimulates GTP hydrolytic activity of the monomeric G protein Rap1. Physical interaction between  $G_{\alpha z}$ , a member of the  $G_i$  family of trimeric G proteins, and Rap1GAP blocks the ability of regulators of G protein signaling to stimulate GTP hydrolysis of the  $\alpha$  subunit, and also attenuates the ability of activated  $G_{\alpha z}$  to inhibit adenylyl cyclase. Rap1GAP is expressed in the brain, kidney and pancreas and may act as a signal integrator to coordinate and/or integrate  $G_2$  signaling and Rap1 signaling in cells. A novel isoform of Rap1 GTPase-activating protein, designated Rap1GAPII, binds specifically to  $G_{\alpha z}$ . Stimulation of the  $G_i$ -coupled M2 Muscarinic receptor translocates Rap1GAPII from the cytosol to the membrane and decreases the amount of GTP-bound Rap1, resulting in the activation of ERK/MAPK.

## REFERENCES

1. Janoueix-Lerosey, I., et al. 1994. Phosphorylation of Rap1GAP during the cell cycle. *Biochem. Biophys. Res. Commun.* 202: 967-975.
2. Kurachi, H., et al. 1997. Human SPA-1 gene product selectively expressed in lymphoid tissues is a specific GTPase-activating protein for Rap1 and Rap2. Segregate expression profiles from a Rap1GAP gene product. *J. Biol. Chem.* 272: 28081-28088.

## CHROMOSOMAL LOCATION

Genetic locus: RAP1GAP (human) mapping to 1p36.12; Rap1gap (mouse) mapping to 4 D3.

## SOURCE

Rap1GAP (D-9) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 15-50 near the N-terminus of Rap1GAP of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g IgG<sub>2a</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Rap1GAP (D-9) is available conjugated to agarose (sc-166586 AC), 500  $\mu$ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-166586 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-166586 PE), fluorescein (sc-166586 FITC), Alexa Fluor<sup>®</sup> 488 (sc-166586 AF488), Alexa Fluor<sup>®</sup> 546 (sc-166586 AF546), Alexa Fluor<sup>®</sup> 594 (sc-166586 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-166586 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-166586 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-166586 AF790), 200  $\mu$ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-166586 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

Alexa Fluor<sup>®</sup> is a trademark of Molecular Probes, Inc., Oregon, USA

## STORAGE

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

## APPLICATIONS

Rap1GAP (D-9) is recommended for detection of Rap1GAP of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

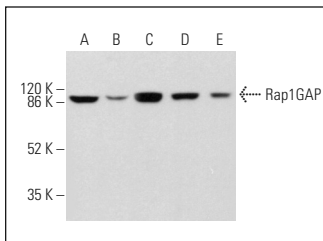
Rap1GAP (D-9) is also recommended for detection of Rap1GAP in additional species, including canine and bovine.

Suitable for use as control antibody for Rap1GAP siRNA (h): sc-36388, Rap1GAP siRNA (m): sc-155959, Rap1GAP siRNA (r): sc-270196, Rap1GAP shRNA Plasmid (h): sc-36388-SH, Rap1GAP shRNA Plasmid (m): sc-155959-SH, Rap1GAP shRNA Plasmid (r): sc-270196-SH, Rap1GAP shRNA (h) Lentiviral Particles: sc-36388-V, Rap1GAP shRNA (m) Lentiviral Particles: sc-155959-V and Rap1GAP shRNA (r) Lentiviral Particles: sc-270196-V.

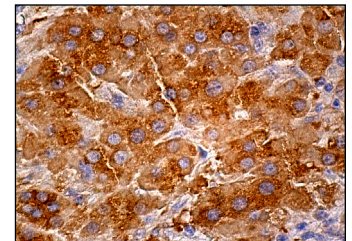
Molecular Weight of Rap1GAP: 89 kDa.

Positive Controls: SK-N-SH cell lysate: sc-2410, Jurkat whole cell lysate: sc-2204 or K-562 whole cell lysate: sc-2203.

## DATA



Rap1GAP (D-9): sc-166586. Western blot analysis of Rap1GAP expression in K-562 (A), SK-N-SH (B), Jurkat (C), HeLa (D) and A549 (E) whole cell lysates. Detection reagent used: m-IgG $\kappa$  BP-HRP: sc-516102.



Rap1GAP (D-9): sc-166586. Immunoperoxidase staining of formalin fixed, paraffin-embedded human adrenal gland tissue showing cytoplasmic staining of glandular cells.

## SELECT PRODUCT CITATIONS

1. Zhou, B., et al. 2017. Mitochondrial activity and oxidative stress functions are influenced by the activation of AhR-induced CYP1A1 overexpression in cardiomyocytes. *Mol. Med. Rep.* 16: 174-180.
2. Gao, W.L., et al. 2018. The downregulation of Rap1 GTPase-activating protein is associated with a poor prognosis in colorectal cancer and may impact on tumor progression. *Oncol. Lett.* 15: 7661-7668.
3. Wu, J., et al. 2019. Novel compound cedrelone inhibits hepatocellular carcinoma progression via PBLD and Ras/Rap1. *Exp. Ther. Med.* 18: 4209-4220.
4. Hoy, J.J., et al. 2020. Protein kinase A inhibitor proteins (PKIs) divert GPCR- $G_{\alpha s}$ -cAMP signaling toward EPAC and ERK activation and are involved in tumor growth. *FASEB J.* E-published.

## RESEARCH USE

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