$G_{\beta 1}$ (F-2): sc-515764



The Power to Question

BACKGROUND

Heterotrimeric G proteins function to relay information from cell surface receptors to intracellular effectors. Each of a very broad range of receptors specifically detects an extracellular stimulus (i.e., a photon, pheromone, odorant, hormone or neurotransmitter), while the effectors (e.g., adenyl cyclase), which act to generate one or more intracellular messengers, are less numerous. Each subunit of the G protein complex is encoded by a member of one of three corresponding gene families $(\alpha,\,\beta,\,\gamma)$. In mammals, there are five different members of the β -subunit family. The β subunits of the G proteins are important regulators of G protein a subunits as well as of certain signal transduction receptors and effectors. In contrast to $G_{\beta\ 1-4}$, which are at least 83% homologous, $G_{\beta\ 5}$ is only 50% homologous to the other β subunits. Human $G_{\beta\ 5}$ is expressed at high levels in brain, pancreas, kidney, and heart.

REFERENCES

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- 2. Gautam, N., et al. 1990. G protein diversity is increased by associations with a variety of γ subunits. Proc. Natl. Acad. Sci. USA 87: 7973-7977.
- Simon, M.I., et al. 1991. Diversity of G proteins in signal transduction. Science 252: 802-808.
- 4. von Weizsäcker, E., et al. 1992. Diversity among the β subunits of hetero-trimeric GTP-binding proteins: characterization of a novel β -subunit cDNA. Biochem. Biophys. Res. Commun. 183: 350-356.
- 5. Kleuss, C., et al. 1992. Different β subunits determine G protein interaction with transmembrane receptors. Nature 358: 424-426.
- Blank, J.L., et al. 1992. Activation of cytosolic phosphoinositide phospholipase C by G protein βγ subunits. J. Biol. Chem. 267: 23069-23075.
- 7. Hurowitz, E.H., et al. 2000. Genomic characterization of the human heterotrimeric G protein α , β and γ subunit genes. DNA Res. 7: 111-120.

CHROMOSOMAL LOCATION

Genetic locus: GNB1 (human) mapping to 1p36.33; Gnb1 (mouse) mapping to 4 E2.

SOURCE

 $\text{G}_{\beta\,1}$ (F-2) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 189-201 within an internal region of $\text{G}_{\beta\,1}$ of human origin.

PRODUCT

Each vial contains 200 μ g IgM kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

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

APPLICATIONS

 $G_{\beta,1}$ (F-2) is recommended for detection of $G_{\beta,1}$ of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], 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).

Suitable for use as control antibody for G $_{\beta\,1}$ siRNA (h): sc-41762, G $_{\beta\,1}$ siRNA (m): sc-41763, G $_{\beta\,1}$ shRNA Plasmid (h): sc-41762-SH, G $_{\beta\,1}$ shRNA Plasmid (m): sc-41763-SH, G $_{\beta\,1}$ shRNA (h) Lentiviral Particles: sc-41762-V and G $_{\beta\,1}$ shRNA (m) Lentiviral Particles: sc-41763-V.

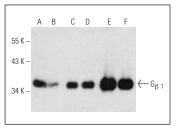
Molecular Weight of G_{B 1}: 36 kDa.

Positive Controls: NCI-H226 whole cell lysate: sc-364256, HeLa whole cell lysate: sc-2200 or Jurkat whole cell lysate: sc-2204.

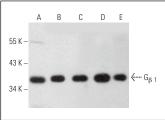
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein L-Agarose: sc-2336 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA







 $G_{\beta,1}$ (F-2): sc-515764. Western blot analysis of G_{β} expression in HeLa (A), Jurkat (B) and NCI-H226 (C whole cell lysates and human hippocampus (D) and human eye (E) tissue extracts.

SELECT PRODUCT CITATIONS

 Zhang, Y., et al. 2022. Paeoniflorin-6'0-benzene sulfonate suppresses fibroblast-like synoviocytes proliferation and migration in rheumatoid arthritis through regulating GRK2-Gβγ interaction. Exp. Ther. Med. 24: 523.

RESEARCH USE

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