## SANTA CRUZ BIOTECHNOLOGY, INC.

# G<sub>β</sub> (F-8): sc-136975



#### 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  $\beta$  subunit family. The  $\beta$  subunits of the G proteins are important regulators of G protein  $\alpha$  subunits as well as of certain signal transduction receptors and effectors. In contrast to G<sub> $\beta$ 1-4</sub>, which are at least 83% homologous, G<sub> $\beta$ 5</sub> is only 50% homologous to the other  $\beta$  subunits. Human G<sub> $\beta$ 5</sub> is expressed at high levels in brain, pancreas, kidney, and heart.

### REFERENCES

- Blatt, C., et al. 1988. Chromosomal localization of genes encoding guanine nucleotide-binding protein subunits in mouse and human. Proc. Natl. Acad. Sci. 85: 7642-7646.
- 2. Modi, W.S., et al. 1989. Chromosomal localization of the gene encoding a third form of the  $\beta$  subunit of GTP-binding regulatory proteins. Cytogenet. Cell Genet. 51: 1046.
- 3. Levine, M.A., et al. 1990. Chromosomal localization of the genes encoding two forms of the G-protein  $\beta$  polypeptide,  $\beta$ -1 and  $\beta$ -3, in man. Genomics 8: 380-386.
- Simon, M.I., et al. 1991. Diversity of G proteins in signal transduction. Science 252: 802-808.
- 5. von Weizsäcker, E., et al. 1992. Diversity among the  $\beta$  subunits of heterotrimeric GTP-binding proteins: characterization of a novel  $\beta$  subunit cDNA. Biochem. Biophys. Res. Commun. 183: 350-356.
- 6. Kleuss, C., et al. 1992. Different  $\beta$  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.

#### SOURCE

 $G_{\beta}$  (F-8) is a mouse monoclonal antibody raised against amino acids 1-300 mapping at the N-terminus of transducin  $\beta$ 2 of human origin.

#### PRODUCT

Each vial contains 200  $\mu g$  lgG  $_1$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

#### STORAGE

Store at 4° C, \*\*D0 NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### PROTOCOLS

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

#### APPLICATIONS

 $\rm G_{\beta}$  (F-8) is recommended for detection of  $\rm G_{\beta}$  1,  $\rm G_{\beta}$  2,  $\rm G_{\beta}$  3 and  $\rm G_{\beta}$  4 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffinembedded 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).

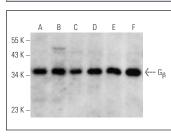
Molecular Weight of  $G_{\beta}$ : 36 kDa.

Positive Controls: Jurkat whole cell lysate: sc-2204, NIH/3T3 whole cell lysate: sc-2210 or AMJ2-C8 whole cell lysate: sc-364366.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850. 4) Immunohistochemistry: use m-IgGκ BP-HRP: sc-516102 with DAB, 50X: sc-24982 and Immunohistomount: sc-45086, or Organo/Limonene Mount: sc-45087.

#### DATA





 ${\rm G}_{\beta}$  (F-8): sc-136975. Western blot analysis of  ${\rm G}_{\beta}$  expression in Jurkat (A), U-251-MG (B), NIH/3T3 (C), AMJ2-C8 (D), C6 (E) and NRK (F) whole cell lysates.

 $G_{\beta}$  (F-8): sc-136975. Immunoperoxidase staining of formalin fixed, paraffin-embedded human urinary bladder tissue showing cytoplasmic and membrane staining of urothelial cells.

#### SELECT PRODUCT CITATIONS

1. Liao, H.R., et al. 2022. Larixol inhibits fMLP-induced superoxide anion production and chemotaxis by targeting the  $\beta\gamma$  subunit of G<sub>i</sub>-protein of fMLP receptor in human neutrophils. Biochem. Pharmacol. 201: 115091.

#### **RESEARCH USE**

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