

G_β (H-1): sc-166123



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., adenylyl 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 (α , β and γ). 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 α subunits as well as of certain signal transduction receptors and effectors. In contrast to G _{β 1-4}, which are at least 83% homologous, G _{β 5} is only 50% homologous to the other β subunits. Human G _{β 5} is expressed at high levels in brain, pancreas, kidney, and heart.

SOURCE

G _{β} (H-1) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 302-340 at the C-terminus of G _{β} of mouse origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

G _{β} (H-1) is available conjugated to agarose (sc-166123 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-166123 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-166123 PE), fluorescein (sc-166123 FITC), Alexa Fluor[®] 488 (sc-166123 AF488), Alexa Fluor[®] 546 (sc-166123 AF546), Alexa Fluor[®] 594 (sc-166123 AF594) or Alexa Fluor[®] 647 (sc-166123 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-166123 AF680) or Alexa Fluor[®] 790 (sc-166123 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-166123 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

APPLICATIONS

G _{β} (H-1) is recommended for detection of G _{β 1}, G _{β 2}, G _{β 3} and G _{β 4} 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), 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).

G _{β} (H-1) is also recommended for detection of G _{β 1}, G _{β 2}, G _{β 3} and G _{β 4} in additional species, including equine, canine, bovine, porcine and avian.

Molecular Weight of G _{β} : 36 kDa.

Positive Controls: MCF7 whole cell lysate: sc-2206, C6 whole cell lysate: sc-364373 or U-251-MG whole cell lysate: sc-364176.

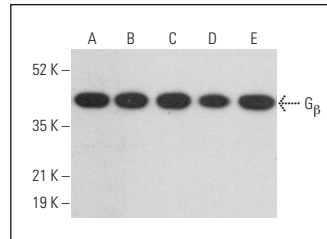
STORAGE

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

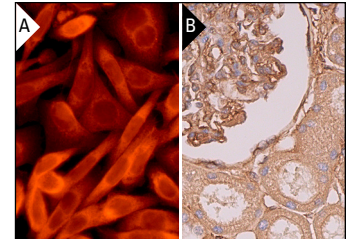
RESEARCH USE

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

DATA



G _{β} (H-1): sc-166123. Western blot analysis of G _{β} expression in U-251-MG (A), NIH/3T3 (B), AMJ2-C8 (C), C6 (D) and MCF7 (E) whole cell lysates. Detection reagent used: m-IgGκ BP-HRP: sc-516102.



G _{β} (H-1) Alexa Fluor[®] 546: sc-166123 AF546. Direct immunofluorescence staining of formalin-fixed SW480 cells showing cytoplasmic and membrane localization. Blocked with UltraCruz[®] Blocking Reagent: sc-516214 (A). G _{β} (H-1): sc-166123. Immunoperoxidase staining of formalin fixed, paraffin-embedded human kidney tissue showing cytoplasmic staining of cells in glomeruli and cells in tubules (B).

SELECT PRODUCT CITATIONS

- May, V., et al. 2010. Pituitary adenylate cyclase-activating polypeptide (PACAP)/PAC1HOP1 receptor activation coordinates multiple neurotrophic signaling pathways: Akt activation through phosphatidylinositol 3-kinase γ and vesicle endocytosis for neuronal survival. *J. Biol. Chem.* 285: 9749-9761.
- Onken, M.D., et al. 2018. Targeting nucleotide exchange to inhibit constitutively active G protein α subunits in cancer cells. *Sci. Signal.* 11: eaao6852.
- Ruiz-Velasco, A., et al. 2020. Targeting mir128-3p alleviates myocardial Insulin resistance and prevents ischemia-induced heart failure. *Elife* 9: e54298.
- Kawakami, K., et al. 2022. Heterotrimeric G _{α} proteins act as a switch for GRK5/6 selectivity underlying β -arrestin transducer bias. *Nat. Commun.* 13: 487.
- Kaur, N., et al. 2022. Paracrine signal emanating from stressed cardiomyocytes aggravates inflammatory microenvironment in diabetic cardiomyopathy. *iScience* 25: 103973.
- Zhang, Y., et al. 2022. Paeoniflorin-6'-O-benzene sulfonate suppresses fibroblast-like synoviocytes proliferation and migration in rheumatoid arthritis through regulating GRK2-G β γ interaction. *Exp. Ther. Med.* 24: 523.
- Gavid, M., et al. 2023. Technique of flat-mount immunostaining for mapping the olfactory epithelium and counting the olfactory sensory neurons. *PLoS ONE* 18: e0280497.

PROTOCOLS

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

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