SANTA CRUZ BIOTECHNOLOGY, INC.

GM-CSFRa (8G6 BOT): sc-80649



BACKGROUND

The human IL-3, IL-5 and GM-CSF receptors are each composed of both unique α subunits and a common β subunit. The α subunits are low affinity ligand binding proteins while the β subunits do not themselves bind ligand, but are required for high affinity binding by the α subunits. In contrast, the mouse IL-3 receptor has two distinct β subunits, one that functions only in IL-3 mediated cell signaling and a second that is shared with IL-5 and GM-CSF. The murine β subunits are 91% homologous at the amino acid level but only 56% homologous to the human β subunit. Although neither the murine nor the human β subunit contains tyrosine kinase domains, both activate tyrosine phosphorylation mediated signaling pathways.

REFERENCES

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- 4. Sakamaki, K., et al. 1992. Critical cytoplasmic domains of the common β subunit of the human GM-CSF, IL-3, and IL-5 receptors for growth signal transduction and tyrosine phosphorylation. EMBO J. 11: 3541-3549.
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- Rao, P., et al. 1995. Human IL-3 receptor signaling: rapid induction of phosphatidylcholine hydrolysis is independent of protein kinase C but dependent on tyrosine phosphorylation in transfected NIH 3T3 cells. J. Immunol. 154: 1664-1674.

CHROMOSOMAL LOCATION

Genetic locus: CSF2RA (human) mapping to Xp22.33/Yp11.32; Csf2ra (mouse) mapping to 19 D3.

SOURCE

 $GM\text{-}CSFR\alpha$ (8G6 BOT) is a mouse monoclonal antibody raised against GM-CSFR α of human origin.

PRODUCT

Each vial contains 200 μg IgG_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

GM-CSFR α (8G6 BOT) is recommended for detection of GM-CSFR α of mouse, rat and human origin by Western Blotting (starting dilution 1:200, 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) and flow cytometry (1 µg per 1 x 10⁶ cells).

Suitable for use as control antibody for GM-CSFR α siRNA (h): sc-35501, GM-CSFR α siRNA (m): sc-40057, GM-CSFR α shRNA Plasmid (h): sc-35501-SH, GM-CSFR α shRNA Plasmid (m): sc-40057-SH, GM-CSFR α shRNA (h) Lentiviral Particles: sc-35501-V and GM-CSFR α shRNA (m) Lentiviral Particles: sc-40057-V.

Molecular Weight of GM-CSFR α : 80 kDa.

Positive Controls: GM-CSFR (h2): 293T Lysate: sc-159461 or HL-60 + DMSO cell lysate: sc-24703.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG 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 A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgG κ BP-FITC: sc-516140 or m-IgG 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





GM-CSFR α (8G6 BOT): sc-80649. Western blot analysis of GM-CSFR α expression in non-transfected: sc-117752 (A) and human GM-CSFR α transfected: sc-159461 (B) 293T whole cell lysates.

GM-CSFRa (866 BOT): sc-80649. Western blot analysis of GM-CSFRa expression in non-transfected sc-117752 (**A**) and human GM-CSFRa transfected: sc-159381 (**B**) 293T whole cell lysates.

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.

PROTOCOLS

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