IL-4R α (S-20): sc-686



The Power to Question

BACKGROUND

The IL-2 receptor is a multicomponent complex consisting of three subunits, α , β and γ , each of which is required for high affinity binding of IL-2. The α chain functions primarily in binding IL-2, whereas the β and γ chains contribute to IL-2 binding and are essential to IL-2-induced activation of signaling pathways leading to T cell growth. Both IL-4R and IL-7R were initially described as single chain, high-affinity ligand-binding cytokine receptors. However, it is now well established that the IL-2R γ chain functions as a second subunit of the high affinity IL-4R and IL-7R receptors. Consequently, the originally described subunits of these latter receptors are now referred to as IL-4R α and IL-7R α , respectively, while the common subunit is referred to as γ_c . Although the common γ chain enhances ligand binding in these three cytokine receptors, it has no capacity to bind these ligands on its own. There is evidence that the γ_c chain is also a subunit of IL-13R.

CHROMOSOMAL LOCATION

Genetic locus: II4ra (mouse) mapping to 7 F3.

SOURCE

IL-4R α (S-20) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the C-terminus of IL-4R α of mouse origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-686 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

IL-4R α (S-20) is recommended for detection of IL-4R α of mouse and rat 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 solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for IL-4R α siRNA (m): sc-35662, IL-4R α shRNA Plasmid (m): sc-35662-SH and IL-4R α shRNA (m) Lentiviral Particles: sc-35662-V.

Molecular Weight of IL-4Rα: 140 kDa.

Molecular Weight of IL-4Rα glycoprotein: 130 kDa.

Positive Controls: WEHI-231 whole cell lysate: sc-2213 or NIH/3T3 whole cell lysate: sc-2210.

RESEARCH USE

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

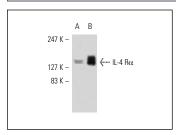
PROTOCOLS

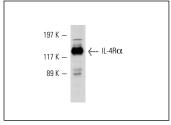
See our web site at www.scbt.com or our catalog for detailed protocols and support products.

STORAGE

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

DATA





IL-4R α (S-20): sc-686. Western blot analysis of IL-4R α expression in untreated NIH/3T3 (**A**) and IL-4 induced NIH/3T3 (**B**) whole cell lysates.

IL-4R α (S-20): sc-686. Western blot analysis of IL-4R α expression in WEHI-231 whole cell lysate.

SELECT PRODUCT CITATIONS

- Webb, D.C., et al. 2004. Polymorphisms in IL-4Rα correlate with airways hyperreactivity, eosinophilia, and Ym protein expression in allergic IL-13-/mice. J. Immunol. 172: 1092-1098.
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- 5. Hocke, A.C., et al. 2006. Regulation of interleukin IL-4, IL-13, IL-10, and their downstream components in lipopolysaccharide-exposed rat lungs. Comparison of the constitutive expression between rats and humans. Cytokine 33: 199-211.
- MacKinnon, A.C., et al. 2008. Regulation of alternative macrophage activation by Galectin-3. J. Immunol. 180: 2650-2658.
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- 8. Shimizu, E., et al. 2008. IL-4-induced selective clearance of oligomeric β -amyloid peptide by rat primary type 2 microglia. J. Immunol. 181: 6503-6513.
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- Kawahara, K., et al. 2012. Intracerebral microinjection of interleukin-4/ interleukin-13 reduces β-amyloid accumulation in the ipsilateral side and improves cognitive deficits in young amyloid precursor protein 23 mice. Neuroscience 207: 243-260.



Try IL-4R α (H-4): sc-28361 or IL-4R α (E-1): sc-165974, our highly recommended monoclonal aternatives to IL-4R α (S-20). Also, for AC, HRP, FITC, PE, α SAlexa Fluor α 488 and Alexa Fluor α 647 conjugates, see IL-4R α (H-4): sc-28361.