

DESCRIPTION

Source	Mouse myeloma cell line, NS0-derived		
	Human Common β Chain (Met1 - Trp443) Accession # P32927	IEGRMD	Human IgG ₁ (Pro100 - Lys330)
	N-terminus		C-terminus

N-terminal Sequence Analysis Gly23 & Glu25

Structure / Form Disulfide-linked homodimer

Predicted Molecular Mass 74.6 kDa (monomer)

SPECIFICATIONS

SDS-PAGE 85 - 92 kDa, reducing conditions

Activity Measured by its binding ability in a functional ELISA.
When recombinant human (rh) GM-CSF R α is present at 0.5 μ g/mL, it binds to rhCommon β Chain/GM-CSF R β Fc Chimera in the presence of rhGM-CSF. The concentration of rhCommon β Chain/GM-CSF R β Fc Chimera that produces 50% of the optimal binding response was found to be approximately 3–12 ng/mL.

Endotoxin Level <0.10 EU per 1 μ g of the protein by the LAL method.

Purity >90%, by SDS-PAGE under reducing conditions and visualized by silver stain.

Formulation Lyophilized from a 0.2 μ m filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE

Reconstitution Reconstitute at 100 μ g/mL in PBS.

Shipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 3 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

The common β chain (β c) is also known as GM-CSF R β (granulocyte-macrophage colony stimulating factor receptor beta subunit), IL-3 R β , and IL-5 R β (gene name CSF2RB and designated CD131) (1, 2). It is a 120 – 135 kDa type I transmembrane protein that associates with the ligand-specific IL-3 R α on T cells and other cells, IL-5 R α on eosinophils or GM-CSF R α on myeloid cells, to form high affinity receptor complexes (1, 2). The 897 amino acid (aa) human β c contains a 16 aa signal sequence, a 427 aa extracellular domain (ECD) with two fibronectin type III domains, a transmembrane sequence, and a 437 aa cytoplasmic domain (2, 3). Within the ECD, human β c shares 57 – 68% aa sequence identity with mouse, rat, equine, porcine, bovine and canine β c. Complexes of ligand with its specific α subunit then bind pre-formed β c dimers, creating 2:2:2 hexamers (and probably 4:4:4 dodecamers) for signaling (3). β c phosphorylation and binding of Jak2 imparts growth and survival signals. Except for eosinophils, β c is primarily involved when rapid production of leukocytes is needed, rather than for developmental or steady-state cell production (1). β c also associates with other receptors, forming heteroreceptor complexes that allow β c complexes to influence the signaling pathways activated by the associated receptor (1). β c thus enhances angiogenesis (when associated with VEGF R2/KDR/Flt-1 or β 1 integrins), cell protection (with Erythropoietin R), and synergistic growth of stem cells (with SCF R/c-kit) (4 – 9). Defective production of β c in humans is a cause of pulmonary alveolar proteinosis (10).

References:

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