

Recombinant Human CD40 Ligand/TNFSF5

Catalog Number: 6420-CLB

DESCRIPTION

Source Human embryonic kidney cell, HEK293-derived human CD40 Ligand/TNFSF5 protein

YPYDVPDYA GCN4-IZ GGGSGGGSGGS (Met113-Leu261)
Accession # P29965.1

N-terminus C-terminus

N-terminal Sequence Tyr (of HA tag)

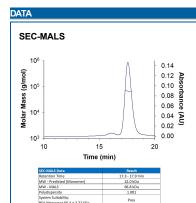
Analysis

Predicted Molecular 22 kDa

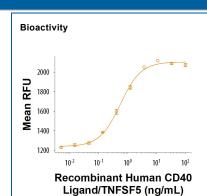
Mass

SPECIFICATIONS	
SDS-PAGE	20-27 kDa, under reducing conditions.
Activity	Measured in a cell proliferation assay using human B cells. The ED ₅₀ for this effect is 0.250-3.00 ng/mL in the presence of Recombinant Human IL-4 (Catalog # 204-IL) and a cross linking HA Tag Antibody (Catalog # MAB060).
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.
Purity	>95%, by SDS-PAGE under reducing conditions and visualized by silver stain.
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS and EDTA with Trehalose. See Certificate of Analysis for details.

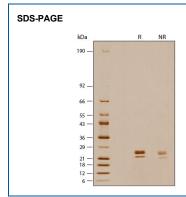
PREPARATION AND STORAGE	
Reconstitution	Reconstitute at 500 μg/mL in PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months 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.



Recombinant Human CD40 Ligand/TNFSF5 Protein SEC-MALS. Recombinant Human CD40 Ligand/TNFSF5 (Catalog # 6420-CLB) has a molecular weight (MW) of 66.8 kDa as analyzed by SEC-MALS, suggesting that this protein is a homotrimer. MW may differ from predicted MW due to post-translational modifications (PTMs) present (i.e. Glycosylation).



Recombinant Human CD40 Ligand/TNFSF5 (HEK293-expressed) Bioactivity. Recombinant Human CD40 Ligand/TNFSF5 (HEK293-expressed) (Catalog # 6420-CLB) stimulates the proliferation of human B cells in the presence of Recombinant Human IL-4 (Catalog # 204-IL) and a cross linking HA Tag Antibody (Catalog # MAB060). The ED₅₀ for this effect is 0.250-3.00 ng/mL.



Recombinant Human CD40 Ligand/TNFSF5 (HEK293expressed) Protein SDS-PAGE. 1 µg/lane of Recombinant Human CD40 Ligand/TNFSF5 (HEK293expressed) Protein (Catalog # 6420-CLB) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by silver staining, showing bands at 20-27 kDa.

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BACKGROUND

CD40 Ligand, also known as TNFSF, CD154, TRAP, and gp39, is a 34-39 kDa type II transmembrane glycoprotein that belongs to the TNF superfamily (1-3). Mature human CD40 Ligand consists of a 22 amino acid (aa) cytoplasmic domain, a transmembrane segment, and an 215 aa extracellular region (4, 5). The extracellular domain of human CD40 Ligand shares 74% and 76% aa sequence identity with mouse and rat CD40 Ligand, respectively. Similar to other TNF superfamily members, CD40 Ligand forms a bioactive homotrimer, both as membrane bound and soluble forms (6-9). The 18 kDa soluble form (aa 113-261) arises from proteolytic processing. Mutation and alternative splicing generate additional forms of CD40 Ligand that are often truncated or non-trimerizable (8). CD40 Ligand is expressed on platelets, as well as on activated T cells and B cells, basophils, eosinophils, fibroblasts, mast cells, monocytes, natural killer cells, vascular endothelial cells, and smooth muscle cells. CD40 Ligand binds to CD40, which is expressed on the surface of B cells, dendritic cells, macrophages, monocytes, platelets, endothelial, and epithelial cells (10). The interaction of CD40 Ligand with CD40 initiates signaling in both CD40 and CD40 Ligand expressing cells (11). CD40 ligation by CD40 Ligand promotes B cell activation and T cell-dependent humoral responses (12, 13). CD40 Ligand dysregulation on T cells and antigen presenting cells contributes to the immune deficiency associated with HIV infection and AIDS (14, 15). It is also implicated in the pathology of multiple cardiovascular diseases including atherosclerosis, atherothrombosis, and restenosis (16, 17).

References:

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