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Human CD30 (TNFRSF8) Protein (C-His)

Catalog Number:	804601, 804602
Size:	25 ug, 100 ug
Target Name:	TNFRSF8, CD30, Ki-1
Regulatory Status:	RUO

PRODUCT DETAILS

Application:	ELISA, BLI
Format:	Liquid, Purified
Expression Host:	CHO
Species:	Human
Sources:	Recombinant Human CD30 protein (phe19-Lys379) with C-terminus His tag is expressed in CHO cells.
Accession Number:	P28908
Molecular Weight:	The protein has a predicted molecular weight of 40 kDa. Under DTT-reducing conditions, it migrates at approximately 50-80 kDa on SDS-PAGE.
Affinity Tag:	C-His
Purity:	>80% based on SDS-PAGE under reducing condition
Formulation:	1xPBS buffer, pH7.4, 0.22 μm filtered
Endotoxin level:	Not tested
Protein Concentration:	25μg size is bottled at 0.2mg/mL concentration. 100 μg size is supplied at a lot-specific concentration.
Storage and Handling:	Briefly centrifuge the vial upon receipt. An unopened vial can be stored at 4°C for up to 2 weeks, or at -20°C or below for up to six months. The protein may be further diluted to 0.1 mg/mL using 0.22 μm-filtered PBS buffer (pH 7.4). For long-term storage, the diluted stock solution should be aliquoted and stored at ≤ –70°C to minimize freeze-thaw cycles. If additional dilution is required, carrier proteins such as FBS or BSA should be added to maintain protein stability.

BACKGROUND INFORMATION

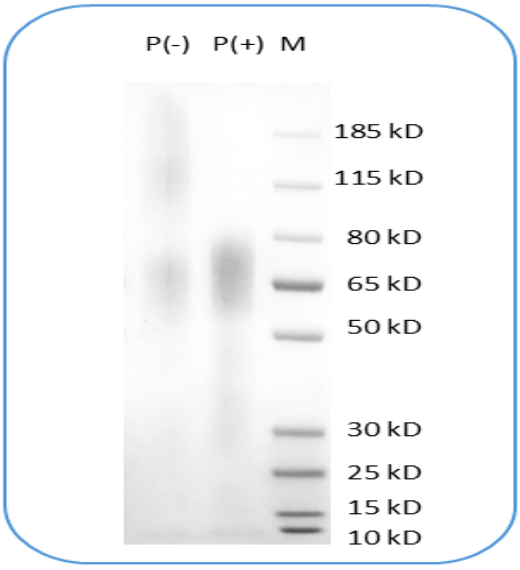
CD30, also known as TNFRSF8, is a type I transmembrane glycoprotein and a member of the tumor necrosis factor receptor (TNFR) superfamily. It is predominantly expressed on activated T and B lymphocytes, and under pathological conditions, on certain lymphoma and carcinoma cells. CD30 functions as a costimulatory receptor involved in regulating cell proliferation, survival, and apoptosis, depending on the context of activation. Its signaling is particularly important in immune cell regulation and inflammatory responses.

Structurally, CD30 consists of three main regions: an extracellular cysteine-rich domain, a single transmembrane domain, and a cytoplasmic tail. The extracellular domain contains multiple cysteine-rich repeats characteristic of TNFR family members, which are essential for ligand binding and receptor oligomerization. The cytoplasmic tail interacts with TNF receptor–associated factor (TRAF) adaptor proteins, initiating downstream signaling cascades such as NF-κB and MAPK pathways. These pathways influence cytokine production and cell fate decisions between survival and programmed cell death.

The primary ligand for CD30 is CD30 ligand (CD30L or CD153), a transmembrane protein belonging to the TNF ligand superfamily. CD30L is expressed mainly on activated T cells, B cells, and some macrophages. Interaction between CD30 and CD30L induces bidirectional signaling: CD30 engagement promotes cell activation or apoptosis depending on cellular context, while reverse signaling through CD30L can modulate immune cell function. The CD30/CD30L axis therefore plays a dual role in immune regulation, contributing to both immune activation and resolution.

In disease contexts, CD30 is clinically significant as a biomarker and therapeutic target. It is highly expressed in certain lymphoid malignancies, including Hodgkin lymphoma and anaplastic large cell lymphoma (ALCL). Its restricted normal expression and high density on tumor cells make CD30 an ideal target for immunotherapy. The most notable example is brentuximab vedotin, an antibody–drug conjugate that binds CD30 and delivers a cytotoxic payload directly to malignant cells. CD30-targeted therapies have achieved remarkable clinical success, leading to durable remissions in relapsed or refractory lymphomas. Moreover, CD30 expression is being explored in non-lymphoid cancers and autoimmune diseases, indicating broader therapeutic potential in modulating immune signaling and inflammation.

PRODUCT DATA



Human CD30 protein (C-His) on SDS-PAGE under reducing condition (P+) and non-reducing condition (P-). The gel was stained for 1 hour with BlinkBlue (catalog 700102). The purity of this protein appears to be greater than 80% based on reducing conditions.

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