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APC Human CD200 (OX-2) Protein (C-Fc)

Catalog Number:	815103, 815104
Size:	25 ug, 100 ug
Target Name:	CD200, MOX1, MOX2, MRC, OX-2, My033
Regulatory Status:	RUO

PRODUCT DETAILS

Application:	Flow Cytometry
Format:	Liquid, APC
Expression Host:	CHO
Species:	Human
Sources:	Recombinant Human CD200 (Gln31-Gly232) with C-terminus Fc-tag is expressed in CHO cell and conjugated to APC.
Accession Number:	P41217
Molecular Weight:	The protein has a predicted molecular weight of 48.7 kDa. Under DTT-reducing conditions, it migrates at approximately 65 kDa on SDS-PAGE prior to conjugation.
Affinity Tag:	C-Fc
Formulation:	1xPBS buffer, pH7.4, 0.09% NaN3 with a carrier protein
Endotoxin level:	Not tested
Protein Concentration:	25μg size is bottled at 0.2 mg/mL concentration. 100 μg size is bottled at lot specific concentration.
Storage and Handling:	Briefly centrifuge the vial upon receipt. An unopened vial may be stored at 2–8°C for up to six months.

BACKGROUND INFORMATION

CD200, also known as OX-2, is an immunoregulatory cell surface glycoprotein that plays a key role in maintaining immune tolerance and limiting inflammatory responses. It is broadly expressed on a variety of cell types, including thymocytes, B cells, activated T cells, dendritic cells, endothelial cells, neurons, and certain tumor cells. CD200 functions primarily by delivering inhibitory signals to myeloid lineage cells, thereby suppressing excessive immune activation and protecting tissues from immune-mediated damage.

Structurally, CD200 is a type I transmembrane protein belonging to the immunoglobulin superfamily. It consists of two extracellular immunoglobulin-like domains (one variable-like and one constant-like), a single transmembrane helix, and a short cytoplasmic tail that lacks known signaling motifs. Unlike many immune receptors, CD200 does not signal intracellularly through its own cytoplasmic domain. Instead, its biological effects are mediated through engagement of its receptor, CD200R, which is expressed predominantly on macrophages, monocytes, dendritic cells, mast cells, and some T cell subsets.

The primary ligand for CD200 is CD200R (CD200 receptor), an inhibitory receptor containing cytoplasmic signaling motifs that recruit adaptor proteins and downstream inhibitory pathways. Binding of CD200 to CD200R suppresses pro-inflammatory cytokine production, reduces antigen presentation capacity, and promotes an anti-inflammatory or tolerogenic phenotype in myeloid cells. This interaction is particularly important in immune-privileged sites such as the central nervous system, where CD200 expression on neurons helps restrain microglial activation.

Dysregulation of CD200 signaling has been implicated in several diseases. Overexpression of CD200 is observed in various malignancies, including chronic lymphocytic leukemia (CLL), multiple myeloma, and certain solid tumors, where it contributes to immune evasion by suppressing antitumor immunity. Conversely, insufficient CD200 signaling may exacerbate autoimmune or inflammatory conditions due to unchecked myeloid activation.

Therapeutically, CD200 is being explored as a target in oncology and immune modulation. Blocking antibodies against CD200 aim to restore antitumor immune responses by relieving myeloid suppression. In contrast, agonistic strategies enhancing CD200–CD200R signaling are being investigated for inflammatory and autoimmune diseases. By modulating innate immune checkpoints, CD200 represents a promising target for rebalancing immune responses in diverse clinical settings.

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