

Anti-Human CD14 Antibody

Catalog Number:	106101, 106102
Size:	100 ug, 500 ug
Target Name:	CD14, LPS receptor
Regulatory Status:	RUO

PRODUCT DETAILS

Clone:	63D3
Application:	Flow Cytometry
Reactivity:	Human
Format:	Purified
Isotype:	Mouse IgG1
Antibody Type:	Monoclonal
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide
Protein Concentration:	0.5 mg/mL
Storage and Handling:	The antibody solution should be stored between 2°C and 8°C
Recommended Usage:	For flow cytometric staining, it is recommended to use less than 0.2 µg of this reagent per 0.5-1.0 million cells in a 100 µL volume. Optimal reagent performance should be determined by titration for each specific application
Isotype Control:	301401

BACKGROUND INFORMATION

CD14 is a pattern recognition receptor that plays a central role in innate immune sensing of microbial products, particularly components of bacterial cell walls. It is primarily expressed on cells of the myeloid lineage, including monocytes, macrophages, neutrophils, and to a lesser extent dendritic cells. CD14 functions as a coreceptor in the recognition of pathogen-associated molecular patterns (PAMPs) and is especially important for initiating inflammatory responses to Gram-negative bacterial infections.

Structurally, CD14 is a glycoprotein composed largely of leucine-rich repeat (LRR) motifs that form a curved solenoid structure optimized for ligand binding. CD14 exists in two main forms: a membrane-bound form (mCD14), which is attached to the cell surface via a glycosylphosphatidylinositol (GPI) anchor, and a soluble form (sCD14), which is present in plasma and other body fluids. Unlike many immune receptors, CD14 lacks a transmembrane and intracellular signaling domain, meaning it cannot signal independently and must cooperate with other receptors to transmit activation signals.

Functionally, CD14 acts as a high-affinity binding protein for lipopolysaccharide (LPS), a major component of the outer membrane of Gram-negative bacteria. In conjunction with LPS-binding protein (LBP), CD14 transfers LPS to the Toll-like receptor 4 (TLR4)-MD-2 complex, which then initiates downstream signaling pathways leading to activation of NF-κB and production of pro-inflammatory cytokines such as TNF-α, IL-1β, and IL-6. Beyond LPS, CD14 can bind a range of microbial ligands, including lipoteichoic acid from

Gram-positive bacteria, peptidoglycan, and certain viral and fungal components, broadening its role in innate immune surveillance.

CD14 is strongly implicated in disease, particularly in conditions driven by excessive inflammation. Overactivation of CD14-dependent pathways contributes to sepsis and septic shock, where uncontrolled cytokine release can lead to tissue damage, organ failure, and death. Elevated levels of soluble CD14 are associated with chronic inflammatory states, including atherosclerosis, metabolic disease, and HIV infection, and are often used as biomarkers of monocyte activation and systemic inflammation.

Therapeutically, CD14 is an attractive target for modulating innate immune responses. Monoclonal antibodies and other inhibitors targeting CD14 have been explored as strategies to dampen harmful inflammation in sepsis and other inflammatory disorders by limiting LPS-induced TLR4 activation. In diagnostics and research, CD14 is widely used as a defining marker of monocytes and macrophages in flow cytometry and immunohistochemistry, underscoring its importance in both clinical and experimental immunology.

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