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InnoCyto Inc.

15375 Barranca Pkwy, Suite I-103
Irvine, CA 92618
Email: info@innocyto.com
Phone: 1-949-418-7543

APC/Cyanine7 Anti-Human CD137 (4-1BB) Antibody

Catalog Number:	108007, 108008
Size:	25 tests, 100 tests
Target Name:	CD137, 4-1BB, ILA, CD-137, TNFRSF9
Regulatory Status:	RUO

PRODUCT DETAILS

Clone:	4B4-1
Application:	Flow Cytometry
Reactivity:	Human
Format:	APC/Cyanine7
Isotype:	Mouse IgG1
Antibody Type:	Monoclonal
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA
Protein Concentration:	Supplied at a lot-specific concentration.
Storage&Handling:	The antibody solution should be stored undiluted between 2°C and 8°C, and protected from prolonged exposure to light. Do not freeze.
Recommended Usage:	For flow cytometric staining, it is recommended to use 5 µL 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. APC/Cyanine7 has an excitation max at 650 nm and an emission max at 774 nm.
Excitation Laser:	Red Laser (633 nm)
Isotype Control:	301405

BACKGROUND INFORMATION

CD137, also known as 4-1BB or TNFRSF9, is a costimulatory receptor that plays an important role in regulating immune responses, particularly those mediated by T lymphocytes and natural killer (NK) cells. CD137 is a member of the tumor necrosis factor receptor (TNFR) superfamily and is expressed primarily on activated immune cells, including CD8+ T cells, CD4+ T cells, NK cells, dendritic cells, and some myeloid cells. Its expression is typically low on resting cells but is rapidly upregulated following antigen recognition or immune activation. CD137 signaling enhances cell survival, proliferation, cytokine production, and cytotoxic activity, making it a key regulator of adaptive and innate immune responses.

Structurally, CD137 is a type I transmembrane glycoprotein consisting of an extracellular domain with several cysteine-rich repeats characteristic of TNFR family members, a single transmembrane region, and a cytoplasmic signaling domain. Unlike receptors that contain intrinsic enzymatic activity, CD137 signals through the recruitment of intracellular adaptor proteins. When activated, the cytoplasmic region interacts with TNF receptor–associated factors (TRAFs), particularly TRAF1 and TRAF2. These interactions trigger

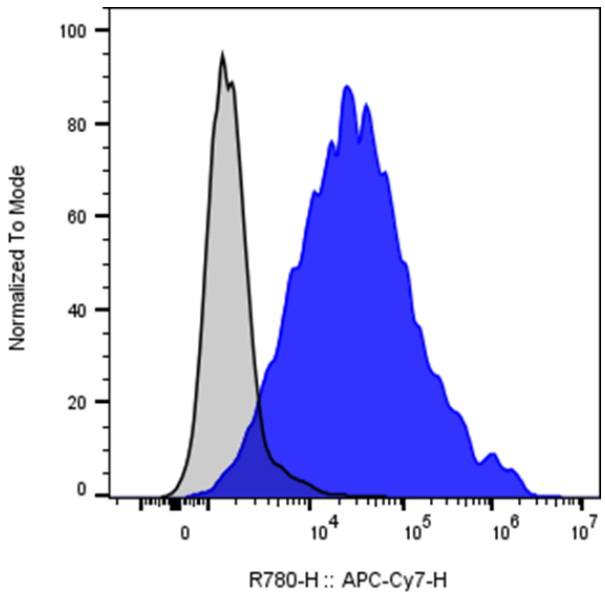
downstream signaling pathways such as NF-κB, MAPK, and PI3K–AKT, which promote cell survival and functional activation.

The natural ligand for CD137 is CD137 ligand (CD137L), also known as 4-1BBL or TNFSF9. CD137L is primarily expressed on antigen-presenting cells including dendritic cells, macrophages, and B cells. Engagement of CD137 by CD137L occurs during immune synapse formation between antigen-presenting cells and activated T cells. This interaction provides a secondary costimulatory signal that strengthens T cell responses following T cell receptor (TCR) engagement. CD137 signaling is particularly important for sustaining the expansion and long-term survival of cytotoxic CD8+ T cells.

CD137 has been implicated in a variety of diseases, especially cancer, chronic infections, and autoimmune conditions. In the context of tumors, CD137 signaling can enhance anti-tumor immunity by boosting the cytotoxic function and persistence of CD8+ T cells and NK cells. However, dysregulated or excessive activation of this pathway may contribute to inflammatory or autoimmune responses. In experimental models, CD137 signaling has also been shown to influence immune regulation in conditions such as viral infections and inflammatory diseases.

Because of its strong immune-activating properties, CD137 has become an important target in immunotherapy. Agonistic antibodies that stimulate CD137 are being investigated as cancer treatments, aiming to amplify anti-tumor T cell responses. Additionally, CD137 signaling domains are commonly incorporated into chimeric antigen receptor (CAR) T cell constructs, where they provide potent costimulatory signals that enhance CAR-T cell persistence and effectiveness. These therapeutic approaches highlight the central role of CD137 in modulating immune activation and improving immune-based therapies.

PRODUCT DATA



Human peripheral blood lymphocytes activated with plate-coated anti-Human CD3/CD28 for three days were stained with APC/Cy7 Anti-Human CD137 clone 4B4-1 (color-filled histogram) or an isotype control (gray histogram).