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In Vivo Star Anti-Mouse CD3e Antibody

Catalog Number:	509301, 509302, 509303
Size:	1 mg, 5 mg, 25 mg
Target Name:	CD3, T3, CD3ε
Regulatory Status:	RUO

PRODUCT DETAILS

Clone:	17A2-m2a
Application:	ELISA, WB, Flow cytometry, IHC, ICC, animal model study
Reactivity:	Mouse
Format:	Liquid
Product Description:	In vivo Grade Recombinant Anti-mouse CD3 Monoclonal Antibody
Isotype:	Mouse IgG2a Kappa
Antibody Type:	Recombinant
Purity:	>95% by reducing SDS-PAGE
Endotoxin:	< 1 EU per 1 mg of the protein by the LAL method.
Storage Conditions:	4ºC
Grade:	In vivo
Recommended Usage:	This product is suitable for in vivo animal use. Optimal amounts need to be determined empirically for each experiment.
Hidden Synonyms:	InVivoMab, InVivoPlus, GoInVivo, In Vivo Gold

BACKGROUND INFORMATION

CD3 is a critical component of the T-cell receptor (TCR) complex, essential for T-cell activation and the adaptive immune response. It is present on nearly all mature T lymphocytes and serves as a marker for their identification. Functionally, CD3 transduces activation signals from the TCR upon antigen recognition, triggering cascades that lead to cytokine production, cell proliferation, and differentiation. This process enables T-cells to respond effectively to pathogens and abnormal cells, anchoring immune defense mechanisms.

Structurally, CD3 is not a single protein but a multimeric complex composed of four distinct polypeptide chains: CD3γ, CD3δ, CD3ε, and CD3ζ. These chains associate non-covalently with the antigen-specific α and β chains of the TCR. Each CD3 subunit contains one or more immunoreceptor tyrosine-based activation motifs (ITAMs) in their cytoplasmic tails. Upon TCR engagement with an antigen–major histocompatibility complex (MHC), the ITAMs become phosphorylated, serving as docking sites for signaling molecules such as ZAP-70, which propagate downstream signaling required for full T-cell activation. CD3 itself does not bind directly to antigens; instead, it functions as a signaling adaptor within the TCR complex. Its effective operation relies on interactions between the TCR and peptide-MHC complexes presented by antigen-presenting cells. The physical and functional association

between CD3 and the TCR ensures that antigen recognition is tightly coupled to intracellular signaling pathways that determine T-cell fate and function.

Alterations in CD3 expression or signaling are implicated in various immune-related diseases. Mutations in CD3 genes can cause severe combined immunodeficiency (SCID), characterized by defective T-cell development. Dysregulated signaling may also contribute to autoimmune conditions, where inappropriate T-cell activation leads to tissue damage. Additionally, CD3 surface expression levels are sometimes altered in chronic infections and leukemias.

Therapeutically, CD3 is an important target in immunomodulation. Monoclonal antibodies such as muromonab-CD3 (OKT3) were among the first biologics used to prevent organ transplant rejection by depleting or inactivating T-cells. More recently, bispecific T-cell engagers (BiTEs) like blinatumomab exploit CD3 to redirect cytotoxic T-cells toward tumor cells expressing specific antigens, offering potent anti-cancer activity. By harnessing or modulating CD3-mediated signaling, modern immunotherapies continue to expand the therapeutic potential of T-cell-based treatments.

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