

## Anti-Human TCR $\gamma/\delta$ Antibody

<b>Catalog Number:</b>	111901, 111902
<b>Size:</b>	25ug, 100 ug
<b>Target Name:</b>	T cell receptor $\gamma/\delta$ , $\gamma/\delta$ TCR, TCR- $\gamma/\delta$
<b>Regulatory Status:</b>	RUO

### PRODUCT DETAILS

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<b>Clone:</b>	B1
<b>Application:</b>	Flow Cytometry
<b>Reactivity:</b>	Human
<b>Format:</b>	Purified
<b>Isotype:</b>	Mouse IgG1
<b>Antibody Type:</b>	Monoclonal
<b>Formulation:</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide
<b>Protein Concentration:</b>	0.5 mg/mL
<b>Storage&amp;Handling:</b>	The antibody solution should be stored between 2°C and 8°C
<b>Recommended Usage:</b>	For flow cytometric staining, it is recommended to use less than 0.2 ug of this reagent per 0.5-1.0 million cells in a 100 $\mu$ L volume. Optimal reagent performance should be determined by titration for each specific application
<b>Isotype Control:</b>	301401

### BACKGROUND INFORMATION

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Human TCR  $\gamma/\delta$  refers to the T cell receptor expressed on  $\gamma/\delta$  T cells, a distinct subset of T lymphocytes that bridge innate and adaptive immunity. Unlike conventional  $\alpha\beta$  T cells,  $\gamma/\delta$  T cells respond rapidly to stress signals, infection, and cellular transformation without requiring classical antigen presentation. They are enriched in epithelial and mucosal tissues, where they contribute to immune surveillance, tissue repair, and early defense against pathogens.

Structurally, the  $\gamma/\delta$  T cell receptor is a heterodimer composed of one gamma ( $\gamma$ ) chain and one delta ( $\delta$ ) chain, each containing variable (V), diversity (D, for  $\delta$ ), joining (J), and constant (C) regions generated through somatic recombination. The receptor associates with the CD3 signaling complex, which transduces activation signals into the cell. Compared to  $\alpha\beta$  TCRs,  $\gamma/\delta$  TCRs exhibit more limited diversity but greater flexibility in antigen recognition.

The ligands recognized by  $\gamma/\delta$  TCRs are diverse and not restricted to peptide antigens presented by classical major histocompatibility complex (MHC) molecules. Instead, they detect non-peptide phosphoantigens, lipid antigens, and stress-induced molecules such as MICA/MICB and butyrophilin family members (e.g., BTN3A1). This allows  $\gamma/\delta$  T cells to sense metabolic dysregulation and cellular stress commonly associated with infection or malignancy.

In disease,  $\gamma/\delta$  T cells play complex roles. They contribute to protective immunity against infections and tumors by producing

cytokines and exerting cytotoxic activity. However, they can also promote inflammation in autoimmune diseases such as psoriasis and multiple sclerosis. Dysregulation of  $\gamma\delta$  T cell responses may therefore contribute to both protective and pathological outcomes.

Therapeutically,  $\gamma\delta$  T cells are being actively explored in immunotherapy. Their ability to recognize tumor cells in an MHC-independent manner makes them attractive candidates for cancer treatment. Approaches include adoptive transfer of expanded  $\gamma\delta$  T cells, stimulation with phosphoantigens, and engineering  $\gamma\delta$  TCR-based therapies. These strategies aim to harness their rapid response and broad reactivity to enhance anti-tumor immunity while minimizing the risk of immune escape.

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