



# Granzyme B Monoclonal Antibody (NGZB), PE-Cyanine7, eBioscience™

<b>Product Details</b>	
Size	100 μg
Species Reactivity	Mouse
Published Species	Mouse
Host/Isotype	Rat / IgG2a, kappa
Recommended Isotype Control	Rat IgG2a kappa Isotype Control (eBR2a), PE-Cyanine7, eBioscience™
Class	Monoclonal
Туре	Antibody
Clone	NGZB
Conjugate	PE-Cyanine7
Form	Liquid
Concentration	0.2 mg/mL
Purification	Affinity chromatography
Storage buffer	PBS, pH 7.2, with 0.1% gelatin
Contains	0.09% sodium azide
Storage conditions	4° C, store in dark, DO NOT FREEZE!
RRID	AB_10853339

Applications	Tested Dilution	Publications
Flow Cytometry (Flow)	0.125 µg/test	27 Publications

## **Product Specific Information**

Description: This NGZB monoclonal antibody reacts with mouse Granzyme B, which is a member of the granzyme serine protease family. Granzyme B is found in the granules of cytotoxic T cells and NK cells. Granzyme B has also been described as CGL1 (cathepsin G-like-1), a serine protease expressed only in cytotoxic T-lymphocytes after cell activation, and CTLA-1 (cytotoxic T lymphocyte-associated serine esterase 1) based on identification of mRNA in various cytotoxic T cells, but not observed in non-cytotoxic lymphoid cells. Granzyme B is crucial for the rapid induction of target cell death by apoptosis, induced by interaction with cytotoxic T cells. The receptor involved has been identified as mannose 6-phosphate receptor. This receptor functions as a death receptor for Granzyme B during cytotoxic T cell-induced apoptosis. This NGZB monoclonal antibody does not crossreact to human Granzyme B nor is staining blocked with GB11, suggesting it recognizes a different epitope.

Applications Reported: This NGZB antibody has been reported for use in intracellular staining and flow cytometric analysis.

Applications Tested: This NGZB antibody has been tested by intracellular staining and flow cytometric analysis of mouse splenocytes using the Intracellular Fixation & Permeabilization Buffer Set (cat. 88-8824) and protocol. Please refer to Best Protocols: Protocol A: Two step protocol for (cytoplasmic) intracellular proteins. This can be used at less than or equal to 0.125 µg per test. A test is defined as the amount (µg) of antibody that will stain a cell sample in a final volume of 100 µL. Cell number should be determined empirically but can range from 10^5 to 10^8 cells/test. It is recommended that the antibody be carefully titrated for optimal performance in the assay of interest.

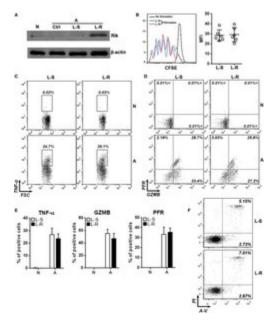
Light sensitivity: This tandem dye is sensitive photo-induced oxidation. Please protect this vial and stained samples from light.

Fixation: Samples can be stored in IC Fixation Buffer (cat. 00-8222) (100 µL cell sample + 100 µL IC Fixation Buffer) or 1-step Fix /Lyse Solution (cat. 00-5333) for up to 3 days in the dark at 4°C with minimal impact on brightness and FRET efficiency /compensation. Some generalizations regarding fluorophore performance after fixation can be made, but clone specific performance should be determined empirically.

Excitation: 488-561 nm; Emission: 775 nm; Laser: Blue Laser, Green Laser, Yellow-Green Laser.

Filtration: 0.2 µm post-manufacturing filtered.

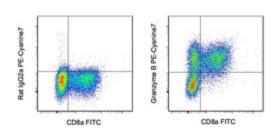
# Advanced Verification Data



## Granzyme B Antibody (25-8898-82)

Figure 3 Overexpression of Rik does not influence CD8 + T cell activation in vitro . (A) Expression of Rik after lentiviral transduction. N: naive CD8 + T cells. A: agonistic antibody-stimulated CD8 + T cells. Ctrl: no transduction. L-S: transduction with lentiviruses containing scramble sequence. L-R: transduction with lentiviruses containing Rik sequence. This is a representative image of two independent experiments. (B) CFSE dilution in lentivirus-transduced CD8 + T cells. Left panel: representative histograms. Right panel: statistics of mean fluorescent intensity. (C,D) Intracellular staining of TNF-alpha (C), granzyme B and perforin (D) in lentivirustransduced CD8 + T cells. Numbers in the plots are proportions of gated cell populations. N: naive CD8 + T cells. A: agonistic antibody-stimulated CD8 + T cells. L-S: transduction with lentiviruses containing scramble sequence. L-R: transduction with lentiviruses containing Rik sequence. (E) Statistical analysis of the proportions of CD8 + T cells expressing TNF-alpha, granzyme B, and perforin. N = 5 per group. (F) CD8 + T cell apoptosis after stimulation and lentiviral transduction. A-V, Annexin V: PI, propidium iodide. This is a representative image of two independent experiments. Cell treatment validation info.

### Product Images For Granzyme B Monoclonal Antibody (NGZB), PE-Cyanine7, eBioscience™



#### Granzyme B Antibody (25-8898-82) in Flow

C57BL/6 mouse splenocytes were stimulated for 72 hours with CD3e and CD28 Monoclonal Antibodies, Functional Grade (Product # 16-0031-85) and (Product # 16-0281-85), followed by Protein Transport Inhibitor Cocktail (Product # 00-4980-03), for an additional 5 hours. Cells were then surface stained with CD8a Monoclonal Antibody, FITC (Product # 11-0081-82) followed by intracellular staining of 0.06  $\mu g$  of Rat IgG2a kappa Isotype Control, PE-Cyanine7 (Product # 25-4321-82) (left) or 0.06  $\mu g$  of Granzyme B Monoclonal Antibody, PE-Cyanine7 (right), using the Intracellular Fixation & Permeabilization Buffer Set (Product # 88-8824-00) and protocol. Cells in the lymphocyte gate were used for analysis.

#### View more figures on thermofisher.com

#### □ 27 References

# Flow Cytometry (27)

**iScience** 

Non-redundant activity of GSK-3 and GSK-3 in T cell-mediated tumor rejection.

"25-8898-82 was used in Flow Cytometry to demonstrate that both isoforms contribute to T cell function to different degrees."

Authors: Steele L, Mannion AJ, Shaw G, Maclennan KA, Cook GP, Rudd CE, Taylor A

**Species** Mouse

Dilution 1:100

**Year** 2021

International journal of nanomedicine

Extracellular Vesicles from *Akkermansia muciniphila* Elicit Antitumor Immunity Against Prostate Cancer via Modulation of CD8<sup>+</sup> T Cells and Macrophages.

"Published figure using Granzyme B monoclonal antibody (Product # 25-8898-82) in Flow Cytometry" Authors: Luo ZW,Xia K,Liu YW,Liu JH,Rao SS,Hu XK,Chen CY,Xu R,Wang ZX,Xie H

Species Not Applicable

**Dilution** Not Cited

**Year** 2021

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# More applications with references on thermofisher.com

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