# biotechne

# Human Granzyme K Antibody

Recombinant Monoclonal Rabbit IgG Clone # 2471A Catalog Number: MAB10216

# **R**Dsystems

DESCRIPTION	
Species Reactivity	Human
Specificity	Detects human Granzyme K in direct ELISAs.
Source	Recombinant Monoclonal Rabbit IgG Clone # 2471A
Purification	Protein A or G purified from cell culture supernatant
Immunogen	Synthetic peptide containing human Granzyme K Accession # P49863
Formulation	Lyophilized from a 0.2 μm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 μm filtered solution in PBS.

## APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	0.1 μg/mL	See Below
Flow Cytometry	25 μg/mL	Human peripheral blood mononuclear cells (PBMCs)
Immunohistochemistry	5-25 μg/mL	See Below

## DATA



#### Detection of Human Granzyme K by Western Blot. Western blot shows lysates of human NK cells, U2OS human osteosarcoma cell line, and RT-4 human bladder carcinoma cell line. PVDF membrane was probed with 0.1 µg/mL of Rabbit Anti-Human Granzyme K Monoclonal Antibody (Catalog # MAB10216) followed by HRP-conjugated Anti-Rabbit IgG Secondary Antibody (Catalog #HAF008). A specific band was detected for Granzyme K at approximately 30 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

### Immunohistochemistry



Granzyme K in Human Tonsil. Granzyme K was detected in immersion fixed paraffinembedded sections of human tonsil using Rabbit Anti-Human Granzyme K Monoclonal Antibody (Catalog # MAB10216) at 5 µg/mL for 1 hour at room temperature followed by incubation with the Anti-Rabbit IgG VisUCyte™ HRP Polymer Antibody (Catalog # VC003). Before incubation with the primary antibody, tissue was subjected to heat-induced epitope retrieval using Antigen Retrieval Reagent-Basic (Catalog # CTS013). Tissue was stained using DAB (brown) and counterstained with hematoxylin (blue). Specific staining was localized to cytoplasm in lymphocytes. View our protocol for IHC Staining with VisUCyte HRP Polymer Detection Reagents.

Rev. 10/10/2022 Page 1 of 2



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# **R**DSYSTEMS

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Flow Cytometry			
Number Nu	Detection of Granzyme K in expanded Human NK cells by      Flow Cytometry. Detection of      Granzyme K in expanded Human      NK cells by Flow Cytometry. NK      cells were expanded from Human      peripheral blood mononuclear      cells (PBMCs) for 14 days using      ExCellerate™ Human NK Cell      Expansion Media (Catalog #      CCM032). NK cells were stained      with (A) Mouse Anti-Human      Granzyme K Monoclonal Antibody      (Catalog # MAB10216), followed      by PE-conjugated anti-Rabbit IgG      secondary antibody (Catalog #      F0110) and Rabbit Anti-Human      CD56 Alexa Fluor® 647-      conjugated Monoclonal Antibody      (Catalog # FAB24086R). (B)      Quadrant markers were set based      on control antibody staining      (Catalog # MAB1050). To      facilitate intracellular staining,      cells were fixed and permeabilized      with FlowX FoxP3 Fixation &      Permeabilization Buffer Kit      (Catalog # FC012). View our      protocol for Staining Intracellular		
PREPARATION AND S	STORAGE		
Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.		
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw control of the second seco	cycles.	

- I month, 2 to 8 C under sterile conditions after reconstitution.
  C months, 20 to 70 °C under sterile conditions offer reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.

## BACKGROUND

Granzymes are released by cytoplasmic granules within NK and cytotoxic T cells. They are serine proteases that induce apoptosis in the target cell. Granzymes have also been found to help initiate the inflammatory response by activating macrophages and mast cells when in an extracellular state. Granzymes have also been found to protect the body against the formation of different kinds of lymphomas.

#### References:

- 1. Bots, M. and JP Medema (2006). J.Cell Sci. 119:5011.
- 2. Walch, M. et al. (2014). Cell. 157:1309.
- 3. Cullen, SP. et al. (2010). Cell Death Differ. 17:616.

Rev. 10/10/2022 Page 2 of 2



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