

Datasheet for ABIN7127523

Recombinant anti-GZMB antibody

2 Images



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Quantity:	100 μL	
Target:	GZMB	
Reactivity:	Human	
Host:	Rabbit	
Antibody Type:	Recombinant Antibody	
Clonality:	Monoclonal	
Conjugate:	This GZMB antibody is un-conjugated	
Application:	Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF)	
Product Details		
Immunogen:	A synthesized peptide derived from human Granzyme B	
Clone:	6A1	
Isotype:	IgG	
Cross-Reactivity:	Human	
Purification:	Affinity-chromatography	
Target Details		
Target:	GZMB	
Alternative Name:	GZMB (GZMB Products)	
Background:	Background: This enzyme is necessary for target cell lysis in cell-mediated immune responses.	

Target Details

It cleaves after Asp. Seems to be linked to an activation cascade of caspases (aspartate-
specific cysteine proteases) responsible for apoptosis execution. Cleaves caspase-3, -7, -9 and
10 to give rise to active enzymes mediating apoptosis.
Aliases: Granzyme B (EC 3.4.21.79) (C11) (CTLA-1) (Cathepsin G-like 1) (CTSGL1) (Cytotoxic T-
lymphocyte proteinase 2) (Lymphocyte protease) (Fragmentin-2) (Granzyme-2) (Human

lymphocyte protein) (HLP) (SECT) (T-cell serine protease 1-3E), GZMB, CGL1 CSPB CTLA1 GRB

UniProt:

P10144

Pathways:

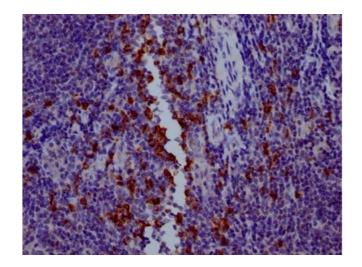
Apoptosis, Caspase Cascade in Apoptosis

Application Details

Application Notes:	Recommended dilution: IHC:1:50-1:200, IF:1:20-1:200,
Restrictions:	For Research Use only

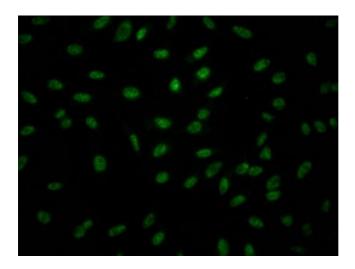
Handling

Format:	Liquid
Buffer:	Rabbit IgG in phosphate buffered saline, pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.



Immunohistochemistry

Image 1. IHC image of ABIN7127523 diluted at 1:100 and staining in paraffin-embedded human tonsil tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10 % normal goat serum 30 min at RT. Then primary antibody (1 % BSA) was incubated at 4 °C overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05 % DAB.



Immunofluorescence

Image 2. Immunofluorescence staining of Hela Cells with ABIN7127523 at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).