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Datasheet for ABIN108721 anti-GZMK antibody

3 Images



Overview

Quantity:	100 µg
Target:	GZMK
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This GZMK antibody is un-conjugated
Application:	ELISA, Flow Cytometry (FACS), Cell-ELISA (cELISA)

Product Details

Immunogen:	genetic immunisation with cDNA encoding human Granzyme K
Clone:	GM-26E7
Isotype:	lgG1
Specificity:	The antibodies do not cross-react with the human granzymes A, B or M.
Purification:	Protein G

Target Details

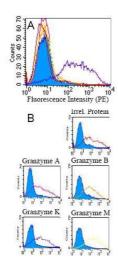
Target:	GZMK
Alternative Name:	Granzyme K (GZMK Products)
Background:	Granzymes are exogenous serine proteases that are stored in the cytotoxic granules of
	activated T cells and NK cells. GM24C3 was generated by genetic immunisation and reacts

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Target Details

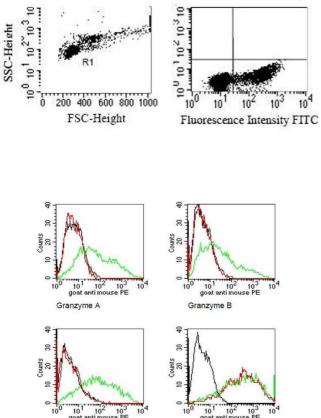
with human granzyme K (GrK), a 28 kDa serine protease with trypsin-like specificity.
P49863
Flow cytometry: 1.2 µg/10^6 cells
ELISA: 1:200 - 1:400
CELISA: 1:200 - 1:400
For each application a titration should be performed to determine the optimal concentration.
ELISA pair with GM6C3 (detector)
For Research Use only
PBS, pH 7.2
Avoid repeated freezing and thawing.
4 °C
short term: 2 °C - 8 °C, long term: -20 °C

Images



Flow Cytometry

Image 1. Specificity testing of GM24C3, GM26E7 andGM6C3. BOSC cells weretransiently transfected withexpression vectors for gran-zyme A, B, K and M as well as an irrelevant protein. Ex-pression of the constructs wastested with an anti-tag mono-clonal antibody (B)



Granzyme M

Granzyme K

Image 2. Intracellular detection of granzyme K in humanPBMC. Mononuclear cells from peripheral blood (PBMC)were seperated by Ficoll-Hypaque, fixed andpermeabilised. Cells incubated with were hybridomasupernatant of GM24C3, GM26E7 or GM6C3for 30minutes. Bound anti

Flow Cytometry

Image 3. BOSC cells were transiently transfected with expression vectors for Granzyme A, B, K, or M. Expression of the constructs was tested with an anti-myc monoclonal antibody (green curves), an irrelevant monoclonal antibody served as negative control (black curves). For specificity testing, E7 hybridoma supernatant was tested on all transfectants. A positive signal was obtained only with Granzyme K transfected cells (red curves).

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