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Recombinant anti-ELANE antibody

Images



Overview

Quantity:	100 μL
Target:	ELANE
Reactivity:	Human
Host:	Rabbit
Antibody Type:	Recombinant Antibody
Clonality:	Monoclonal
Conjugate:	This ELANE antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC), Flow Cytometry (FACS)

Product Details

Immunogen:	A synthesized peptide derived from human Neutrophil Elastase
Clone:	2H12
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	Affinity-chromatography

Target Details

Target:	ELANE
Alternative Name:	ELANE (ELANE Products)
Background:	Background: Modifies the functions of natural killer cells, monocytes and granulocytes. Inhibits

Target Details

	C5a-dependent neutrophil enzyme release and chemotaxis.
	Aliases: Neutrophil elastase (EC 3.4.21.37) (Bone marrow serine protease) (Elastase-2) (Human
	leukocyte elastase) (HLE) (Medullasin) (PMN elastase), ELANE, ELA2
UniProt:	P08246

Application Details	
Application Notes:	Recommended dilution: IHC:1:50-1:200, FC: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

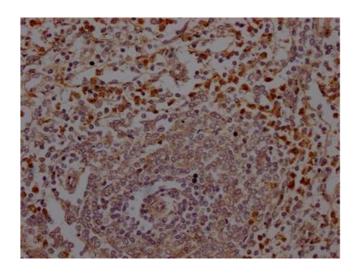
should be handled by trained staff only.

-20 °C,-80 °C

Storage Comment: Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

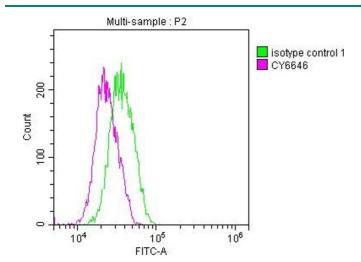
Images

Storage:



Immunohistochemistry

Image 1. IHC image of ABIN7127640 diluted at 1:100 and staining in paraffin-embedded human spleen 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.



Flow Cytometry

Image 2. Overlay histogram showing Jurkat cells stained with ABIN7127640 (red line) at 1:50. The cells were fixed with 70 % Ethylalcohol (18h) and then incubated in 10 % normal goat serum to block non-specific protein-protein interactions followedby the antibody (1 μ g/1*106cells) for 1 h at 4 °C.The secondary antibody used was FITC-conjugated goat anti-rabbit IgG (H+L) at 1/200 dilution for 30 min at 4 °C. Control antibody (green line) was Rabbit IgG (1 μ g/1*106cells) used under the same conditions. Acquisition of >10,000 events was performed.