antibodies

Datasheet for ABIN7143386 anti-ADGRE2 antibody (AA 151-323)

3 Images



Overview

Quantity:	100 µg
Target:	ADGRE2
Binding Specificity:	AA 151-323
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This ADGRE2 antibody is un-conjugated
Application:	ELISA, Immunofluorescence (IF), Immunohistochemistry (IHC)

Product Details

Immunogen:	Recombinant Human Adhesion G protein-coupled receptor E2 protein (151-323AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

Target:	ADGRE2
Alternative Name:	ADGRE2 (ADGRE2 Products)
Background:	Background: Cell surface receptor that binds to the chondroitin sulfate moiety of
	glycosaminoglycan chains and promotes cell attachment. Promotes granulocyte chemotaxis,

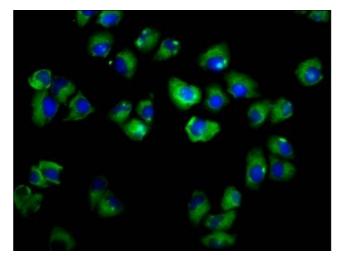
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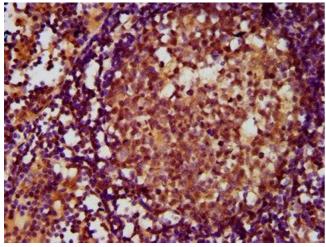
Target Details

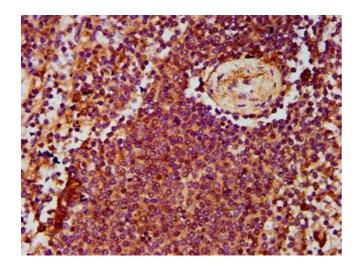
	degranulation and adhesion. In macrophages, promotes the release of inflammatory cytokines,
	including IL8 and TNF. Signals probably through G-proteins. Is a regulator of mast cell
	degranulation (PubMed:26841242).
	Aliases: CD 312 antibody, CD 312 antigen antibody, CD312 antibody, CD312 antigen antibody,
	Egf like module containing mucin like hormone receptor like 2 antibody, Egf like module
	containing mucin like hormone receptor like sequence 2 antibody, EGF like module EMR2
	antibody, EGF-like module receptor 2 antibody, EGF-like module-containing mucin-like hormone
	receptor-like 2 antibody, EMR2 antibody, EMR2_HUMAN antibody
UniProt:	Q9UHX3
Application Details	
Application Notes:	Recommended dilution: IHC:1:500-1:1000, IF:1:50-1:200,
Restrictions:	For Research Use only

Handling

Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: 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.







Immunofluorescence

Image 1. Immunofluorescence staining of Hela cells with ABIN7143386 at 1:166, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

Immunohistochemistry

Image 2. IHC image of ABIN7143386 diluted at 1:500 and staining in paraffin-embedded human lymph node 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 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunohistochemistry

Image 3. IHC image of ABIN7143386 diluted at 1:500 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 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

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