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## anti-ADGRE2 antibody (N-Term)

**Images** 



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Quantity:	100 μL	
Target:	ADGRE2	
Binding Specificity:	N-Term	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This ADGRE2 antibody is un-conjugated	
Application:	ELISA, Western Blotting (WB), Immunofluorescence (IF), Immunocytochemistry (ICC), Immunohistochemistry (IHC)	

### **Product Details**

Immunogen:	A synthesized peptide derived from human EMR2, corresponding to a region within N-terminal amino acids.
Isotype:	IgG
Specificity:	EMR2 Antibody detects endogenous levels of total EMR2.
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

## **Target Details**

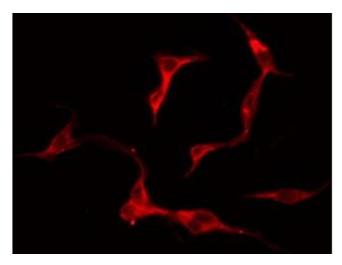
Target:	ADGRE2	
Alternative Name:	ADGRE2 (ADGRE2 Products)	

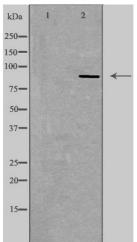
## **Target Details**

Expiry Date:

12 months

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Background:	Description: Cell surface receptor that binds to the chondroitin sulfate moiety of glycosaminoglycan chains and promotes cell attachment. Promotes granulocyte chemotaxis, 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).  Gene: ADGRE2
Molecular Weight:	85 kDa
Gene ID:	30817
UniProt:	Q9UHX3
Application Details	
Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
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
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.





#### Immunofluorescence (fixed cells)

**Image 1.** ABIN6275937 staining COLO205 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod

#### **Western Blotting**

**Image 2.** Western blot analysis of extracts from COLO cells, using EMR2 antibody. The lane on the left is treated with the antigen-specific peptide.