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anti-CARD11 antibody (Internal Region)

2 Images



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Overview	
Quantity:	100 μL
Target:	CARD11
Binding Specificity:	Internal Region
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CARD11 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human CARD11, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	CARD11 Antibody detects endogenous levels of total CARD11.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	CARD11

Target Details

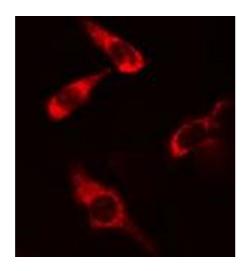
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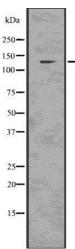
Expiry Date:

rarget Details	
Alternative Name:	CARD11 (CARD11 Products)
Background:	Description: Involved in the costimulatory signal essential for T-cell receptor (TCR)-mediated T-
	cell activation. Its binding to DPP4 induces T-cell proliferation and NF-kappa-B activation in a T-
	cell receptor/CD3-dependent manner. Activates NF-kappa-B via BCL10 and IKK. Stimulates the
	phosphorylation of BCL10. Also activates the TORC1 signaling pathway.
	Gene: CARD11
Molecular Weight:	133 kDa
Gene ID:	84433
UniProt:	Q9BXL7
Pathways:	TCR Signaling, Fc-epsilon Receptor Signaling Pathway, BCR Signaling
Application Details	
Application Notes:	WB 1:1000-3000, IF/ICC 1:100-1:500, 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

Store at -20 °C. Stable for 12 months from date of receipt.

12 months





Immunofluorescence (fixed cells)

Image 1. ABIN6277817 staining COLO205 cells 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) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod

Western Blotting

Image 2. Western blot analysis of CARD11 using HepG2 whole lysates.