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Datasheet for ABIN5518884 anti-TNFRSF11A antibody (Middle Region)

3 Images



Overview

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Quantity:	100 µg
Target:	TNFRSF11A
Binding Specificity:	AA 235-262, Middle Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))
Product Details	
Purpose:	Rabbit IgG polyclonal antibody for Tumor necrosis factor receptor superfamily member
	11A(TNFRSF11A) detection. Tested with WB, IHC-P in Human,Mouse,Rat.
Immunogen:	A synthetic peptide corresponding to a sequence in the middle region of human RANK (235-
	262aa YRKKGKALTANLWHWINEACGRLSGDKE), different from the related mouse sequence by seven amino acids.
Sequence:	YRKKGKALTA NLWHWINEAC GRLSGDKE
lsotype:	lgG
Cross-Reactivity (Details):	No cross reactivity with other proteins.
Characteristics:	Rabbit IgG polyclonal antibody for Tumor necrosis factor receptor superfamily member
	11A(TNFRSF11A) detection. Tested with WB, IHC-P in Human,Mouse,Rat.
	Gene Name: tumor necrosis factor receptor superfamily, member 11a, NFKB activator
	Protein Name: Tumor necrosis factor receptor superfamily member 11A

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Product Details

Purification:

Immunogen affinity purified.

Target Details

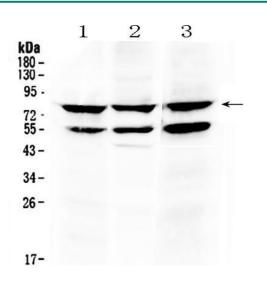
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Target:	TNFRSF11A
Alternative Name:	TNFRSF11A (TNFRSF11A Products)
Background:	Receptor Activator of Nuclear Factor к В (RANK), also known as TRANCE Receptor, is a type I
	membrane protein that is expressed on the surface of osteoclasts and is involved in their
	activation upon ligand binding. RANK is also expressed on dendritic cells and facilitates
	immune signaling. It is found on the surface of stromal cells, osteoblasts, and T cells. By
	analysis of somatic cell and radiation hybrid panels, this gene is mapped to 18q22.1.
	Synonyms: Tumor necrosis factor receptor superfamily member 11A Osteoclast
	differentiation factor receptor ODFR Receptor activator of NF-KB CD265 Tnfrsf11a Rank
	Q9Y6Q6
Gene ID:	8792
UniProt:	Q9Y6Q6
Pathways:	NF-kappaB Signaling
Application Details	
Application Notes:	WB: Concentration: 0.1-0.5 µg/mL, Tested Species: Human, Mouse, Rat
	IHC-P: Concentration: 0.5-1 μ g/mL, Tested Species: Human, Epitope Retrieval by Heat: Boiling
	the paraffin sections in 10 mM citrate buffer, pH 6.0, for 20 mins is required for the staining of
	formalin/paraffin sections.
	Notes: Tested Species: Species with positive results. Other applications have not been tested.
	Optimal dilutions should be determined by end users.
Comment:	Antibody can be supported by chemiluminescence kit ABIN921124 in WB, supported by
	ABIN921231 in IHC(P).
Restrictions:	For Research Use only
Handling	
Handling Format:	Lyophilized

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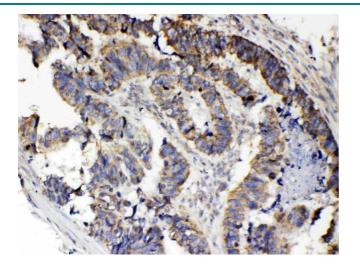
Concentration:	500 μg/mL
Buffer:	Each vial contains 5 mg BSA, 0.9 mg NaCl, 0.2 mg Na2HPO4, 0.05 mg Sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	At -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20 °C for a longer time. Avoid repeated freezing and thawing.

Images



Western Blotting

Image 1. Western blot analysis of RANK using anti- RANK antibody. Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each Lane was loaded with 50ug of sample under reducing conditions. Lane 1: rat thymus tissue lysates, Lane 2: mouse thymus tissue lysates, Lane 3: HEPG2 whole Cell lysates. After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti- RANK antigen affinity purified polyclonal antibody (Catalog #) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002) with Tanon 5200 system. A specific band was detected for RANK at approximately 80KD. The expected band size for RANK is at 66KD.



Immunohistochemistry

Image 2. IHC analysis of RANK using anti- RANK antibody . RANK was detected in paraffin-embedded section of human intestinal cancer tissues. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1µg/ml rabbit anti- RANK Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.

Immunohistochemistry

Image 3. IHC analysis of RANK using anti- RANK antibody . RANK was detected in paraffin-embedded section of human lung cancer tissues. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1µg/ml rabbit anti- RANK Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.

