

Basic Information

Product Name	Anti-TNFR1/TNFRSF1A Antibody	
Gene Name	TNFRSF1A	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, IHC	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence in the middle region of human TNF Receptor I.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	55-60 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied.

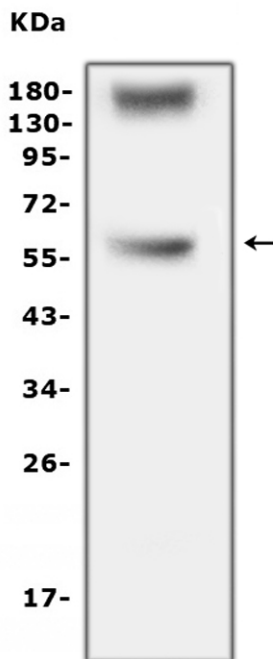
Background Information

Tumor necrosis factor receptor 1 (TNFR1), a potent cytokine, elicits a broad spectrum of biologic responses which are mediated by binding to a cell surface receptor. Its gene is located on 12p13.2. The coding region and the 3-prime untranslated region of TNFR1 are distributed over 10 exons. There are 2 different proteins that serve as major receptors for TNF-alpha, one associated with myeloid cells and one associated with epithelial cells. Additionally, TNFR1 associates with the MADD protein through a death domain-death domain interaction. MADD provides a physical link between TNFR1 and the induction of mitogen-activated protein (MAP) kinase (e.g., ERK2) activation and arachidonic acid release. TNFR1-induced apoptosis involves 2 sequential signaling complexes. Complex I, the initial plasma membrane-bound complex, consists of TNFR1, the adaptor TRADD, the kinase RIP1, and TRAF2 and rapidly signals activation of NF-kappa-B. In a second step, TRADD and RIP1 associate with FADD and caspase-8, forming a cytoplasmic complex, complex II.

Reference

Anti-TNFR1/TNFRSF1A Antibody被引用在1文献中。

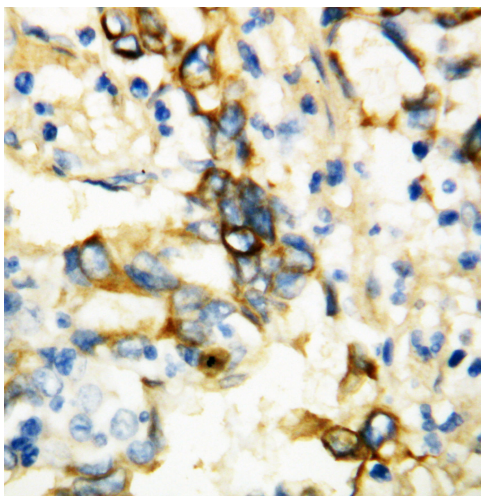
Selected Validation Data



Western blot analysis of TNFR1/TNFRSF1A using anti-TNFR1/TNFRSF1A antibody (BA4891). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human SW579 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-TNFR1/TNFRSF1A antigen affinity purified polyclonal antibody (BA4891) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for TNFR1/TNFRSF1A at approximately 55-60 kDa. The expected band size for TNFR1/TNFRSF1A is at 50 kDa.



IHC analysis of TNFR1/TNFRSF1A using anti-TNFR1/TNFRSF1A antibody (BA4891).

TNFR1/TNFRSF1A was detected in a paraffin-embedded section of human mammary tissue. The tissue section was incubated with rabbit anti-TNFR1/TNFRSF1A Antibody (BA4891) at a dilution of 1:200 and developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB (Catalog # AR1027) as the chromogen.