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# Datasheet for ABIN7151234 anti-TRAF7 antibody (AA 1-90)

2 Images



#### Overview

Quantity:	100 µg
Target:	TRAF7
Binding Specificity:	AA 1-90
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This TRAF7 antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)

#### Product Details

Immunogen:	Recombinant Human E3 ubiquitin-protein ligase TRAF7 protein (1-90AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

## Target Details

Target:	TRAF7
Alternative Name:	TRAF7 (TRAF7 Products)
Background:	Background: E3 ubiquitin ligase capable of auto-ubiquitination, following phosphorylation by
	MAP3K3. Potentiates MEKK3-mediated activation of the NF-kappa-B, JUN/AP1 and DDIT3

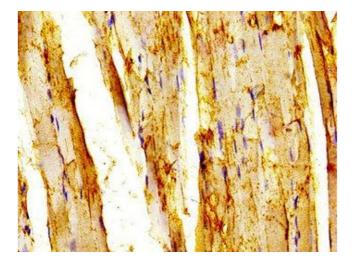
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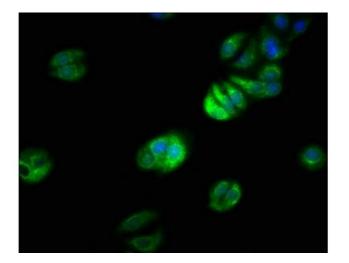
## Target Details

	transcriptional regulators. Induces apoptosis when overexpressed.
	Aliases: E3 ubiquitin protein ligase TRAF7 antibody, E3 ubiquitin-protein ligase TRAF7 antibody,
	RFWD1 antibody, Ring finger and WD repeat domain 1 antibody, RING finger and WD repeat-
	containing protein 1 antibody, RING finger protein 119 antibody, RNF119 antibody, TNF
	receptor associated factor 7 antibody, TNF receptor-associated factor 7 antibody, TRAF7
	antibody, TRAF7_HUMAN antibody
UniProt:	Q6Q0C0

## Application Details

Application Notes:	Recommended dilution: IHC:1:200-1:500, 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.





#### Immunohistochemistry

**Image 1.** IHC image of ABIN7151234 diluted at 1:200 and staining in paraffin-embedded human skeletal muscle 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.

#### Immunofluorescence

**Image 2.** Immunofluorescence staining of HepG2 cells with ABIN7151234 at 1:66, 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).

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