

Basic Information

Product Name	Anti-TRIM33 Antibody	
Gene Name	TRIM33	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, FCM, ICC/IF	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	E.coli-derived human TIF1 gamma recombinant protein (Position: M1001-K1127). Human TIF1 gamma shares 96.1% amino acid (aa) sequence identity with mouse TIF1 gamma.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	150 kDa	
Dilution Ratios	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-200
	Immunocytochemistry/Immunofluorescence (ICC/IF)	1:50-400
	Flow Cytometry (Fixed):	1:50-200
	(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.	

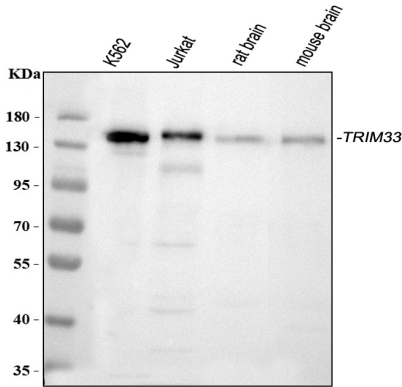
Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

Tripartite motif-containing 33 (TRIM33), also known as transcriptional intermediary factor 1 gamma (TIF1- γ), is a human gene. The TRIM33 gene is mapped to chromosome 1p13 by FISH. The protein encoded by this gene is thought to be a transcriptional corepressor. However, molecules that interact with this protein have not yet been identified. The protein is a member of the tripartite motif family. This motif includes three zinc-binding domains, a RING, a B-box type 1 and a B-box type 2, and a coiled-coil region. Three alternatively spliced transcript variants for this gene have been described; however, the full-length nature of one variant has not been determined.

Selected Validation Data



Western blot analysis of TRIM33 using anti-TRIM33 antibody (PB9836). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: 22RV1 whole cell lysates,

Lane 2: SW620 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-TRIM33 antigen affinity purified polyclonal antibody (PB9836) at a dilution of 1:1000

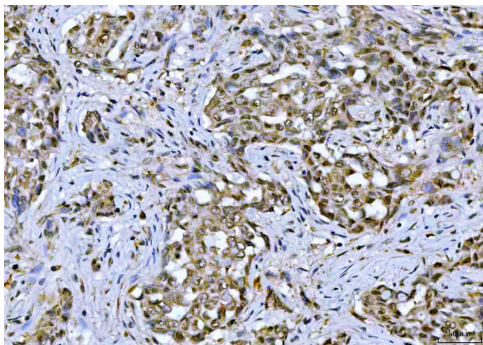
and probed with a goat anti-rabbit IgG-HRP secondary antibody

(Catalog # BA1054). The signal is developed using ECL Plus

Western Blotting Substrate (Catalog # AR1197). A specific band

was detected for TRIM33 at approximately 150 kDa. The expected

band size for TRIM33 is at 123 kDa.



IHC analysis of TRIM33 using anti-TRIM33 antibody (PB9836).

TRIM33 was detected in a paraffin-embedded section of human lung

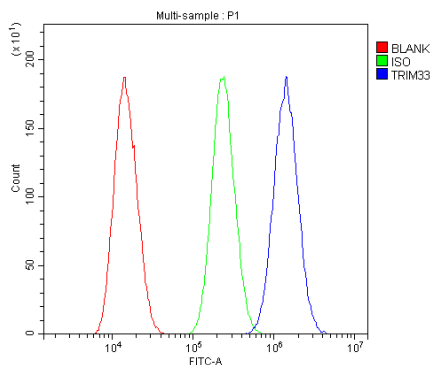
adenocarcinoma tissue. The tissue section was incubated with

rabbit anti-TRIM33 Antibody (PB9836) 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.



Flow Cytometry analysis of U87 cells using anti-TRIM33 antibody (PB9836).

Overlay histogram showing U87 cells stained with PB9836 (Blue line).

To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer.

The cells were blocked with 10% normal goat serum. And then

incubated with rabbit anti-TRIM33 Antibody (PB9836) at 1:100

dilution for 30 min at 20°C. DyLight®488 conjugated goat anti-

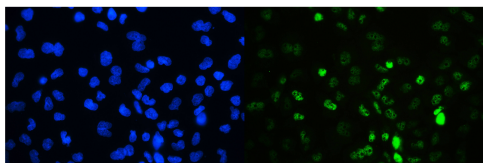
rabbit IgG (BA1127) was used as secondary antibody at 1:100

dilution for 30 minutes at 20°C. Isotype control antibody (Green

line) was rabbit IgG at 1:100 dilution used under the same

conditions. Unlabelled sample without incubation with primary

antibody and secondary antibody (Red line) was used as a blank control.



IF analysis of TRIM33 using anti-TRIM33 antibody (PB9836). TRIM33 was detected in an immunocytochemical section of A431 cells. The section was incubated with rabbit anti-TRIM33 Antibody (PB9836) at a dilution of 1:100. DyLight®488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).