

## Basic Information

<b>Product Name</b>	Anti-DR4/TNFRSF10A Antibody	
<b>Gene Name</b>	TNFRSF10A	
<b>Source</b>	Rabbit	
<b>Isotype</b>	IgG	
<b>Species Reactivity</b>	human,mouse,rat	
<b>Tested Application</b>	WB,ICC/IF,FCM	
<b>Contents</b>	500 ug/ml antibody with PBS , 0.02% NaN <sub>3</sub> , 1 mg BSA and 50% glycerol.	
<b>Immunogen</b>	A synthetic peptide corresponding to a sequence at the N-terminus of human DR4 (99-131aa VLLQVVPSSAATIKLHDQSIGTQQWEHSPLGEL).	
<b>concentration</b>	500 ug/ml	
<b>Purification</b>	Immunogen affinity purified.	
<b>Observed MW</b>	50KD	
<b>Dilution Ratios</b>	Western blot (WB): 1:500-2000 Immunocytochemistry/Immunofluorescence (ICC/IF):1:50-400 Flow cytometry (FCM): 1-3 µg/1x10 <sup>6</sup> cells	

## 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

TNFRSF10A (Tumor Necrosis Factor Receptor Subfamily Member 10A), also known as APO2, DR4 or TRAILR1, is a protein that in humans is encoded by the TNFRSF10A gene. The protein encoded by this gene is a member of the TNF-receptor superfamily. This receptor is activated by tumor necrosis factor-related apoptosis inducing ligand (TNFSF10/TRAIL), and thus transduces cell death signal and induces cell apoptosis. Studies with FADD-deficient mice suggested that FADD, a death domain containing adaptor protein, is required for the apoptosis mediated by this protein.

## Selected Validation Data



Figure 1. Western blot analysis of anti-TNFRSF10A antibody (A02152). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: rat spleen tissue lysates,

Lane 2: mouse spleen tissue lysates,

Lane 3: human MCF-7 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-TNFRSF10A antigen affinity purified polyclonal antibody (A02152) 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 TNFRSF10A at approximately 50 kDa. The expected band size for TNFRSF10A is at 50 kDa.

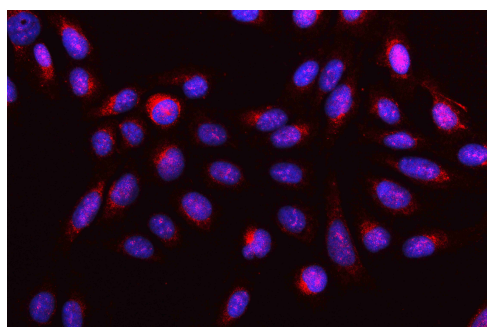


Figure 2. IF analysis of TNFRSF10A using anti-TNFRSF10A antibody (A02152).

TNFRSF10A was detected in an immunocytochemical section of U2OS cells. Cy3-conjugated Anti-rabbit IgG Secondary Antibody (red)(Catalog#BA1032) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).

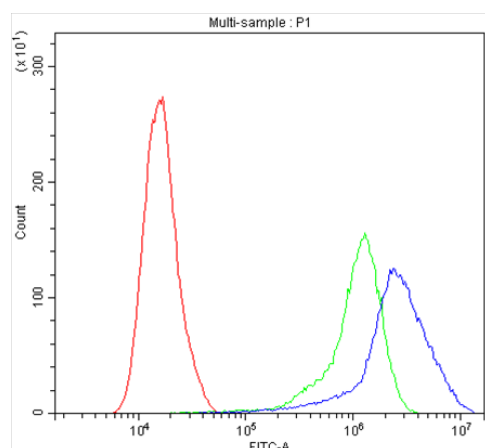


Figure 3. Flow Cytometry analysis of A549 cells using anti-TNFRSF10A antibody (A02152).

Overlay histogram showing A549 cells stained with A02152 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-TNFRSF10A Antibody (A02152, 1  $\mu$ g/ $1 \times 10^6$  cells). DyLight®488 conjugated goat anti-rabbit IgG (BA1127, 5-10  $\mu$ g/ $1 \times 10^6$  cells) was used as secondary antibody. Isotype control antibody (Green line) was rabbit IgG (Catalog # BA1045) (1  $\mu$ g/ $1 \times 10^6$ ) used under the same conditions. Unlabelled sample (Red line) was also used as a control.