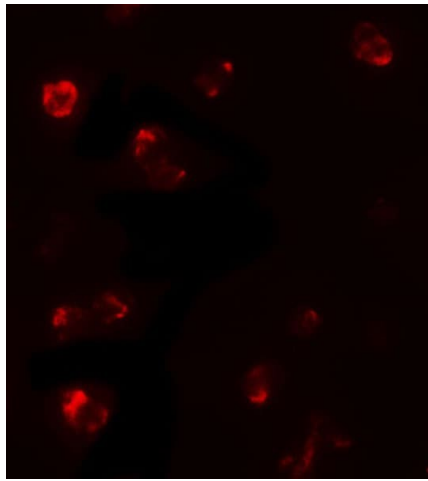
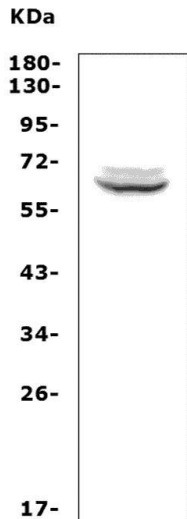


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<b>Product name:</b>	TBX21 /T-bet
<b>Cat number:</b>	AB-84321
<b>Size:</b>	100ug
<b>Clone:</b>	POLY
<b>Concentration:</b>	1mg/ml
<b>Host:</b>	Rb
<b>Isotype:</b>	IgG
<b>Immunogen:</b>	A synthetic peptide corresponding to a sequence of human Tbet/Tbx21(NNVTQMIVLQSLHKYQPR LHIVEVNDGE).
<b>Reactivity:</b>	Hu, Ms, Rt
<b>Applications:</b>	Western blot: 0.25-0.5ug/ml Flow Cytometry: 1-3ug/1x10 <sup>6</sup> cells,
<b>Molecular Weight:</b>	58kDa
<b>Purification:</b>	Aff. Pur.
<b>Form:</b>	Liquid
<b>Buffer:</b>	Each vial contains 4mg Trehalose, 0.9mg NaCl, 0.2mg Na <sub>2</sub> HPO <sub>4</sub> , 0.05mg Na <sub>3</sub> N.
<b>Storage:</b>	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.
<b>Background:</b>	T-box transcription factor TBX21 is a protein that in humans is encoded by the TBX21 gene. It is mapped to 17q21.32. This gene is a member of a phylogenetically conserved family of genes that share a common DNA-binding domain, the T-box. T-box genes encode transcription factors involved in the regulation of developmental processes. This gene is the human ortholog of mouse Tbx21/Tbet gene. Studies in mouse show that Tbx21 protein is a Th1 cell-specific transcription factor that controls the expression of the hallmark Th1 cytokine, interferon-gamma (IFNG). Expression of the human ortholog also correlates with IFNG expression in Th1 and natural killer cells, suggesting a role for this gene in initiating Th1 lineage development from naive Th precursor cells.



Immunofluorescent analysis of 293 cells labeling T-bet / Tbx21 with AB-84247 at 5 µg/ml.

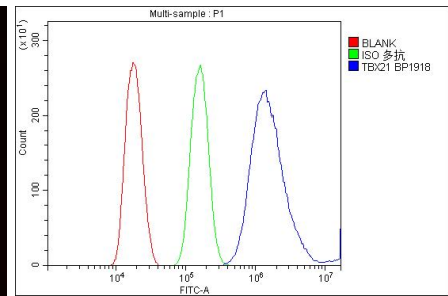


Figure 2. Flow Cytometry analysis of HEP2A1-6 cells using anti-TBX21 antibody. Overlay histogram showing HEP2A1-6 cells stained with (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-TBX21 Antibody 1µg/1x10<sup>6</sup> cells for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (5-10µg/1x10<sup>6</sup> cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1µg/1x10<sup>6</sup>) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

Figure 1. Western blot analysis of TBX21 using anti-TBX21 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 50µg of sample under reducing conditions. Lane 1: mouse RAW246.7 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-TBX21 antigen affinity purified polyclonal antibody 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 1:5000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit with Tanon 5200 system. A specific band was detected for TBX21 at approximately 65KD. The expected band size for TBX21 is at 58KD.

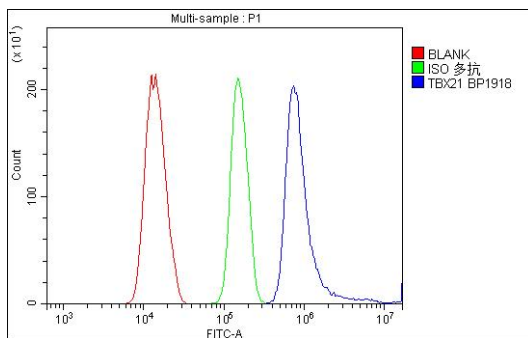


Figure 3. Flow Cytometry analysis of NRK cells using anti-TBX21 antibody. Overlay histogram showing NRK cells stained with (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-TBX21 Antibody (1µg/1x10<sup>6</sup> cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (5-10µg/1x10<sup>6</sup> cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1µg/1x10<sup>6</sup>) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

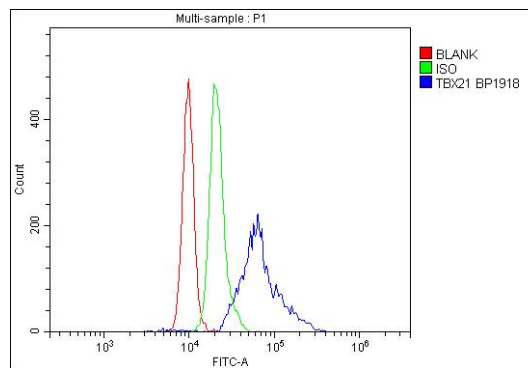


Figure 4. Flow Cytometry analysis of mouse PBMC cells using anti-TBX21 antibody. Overlay histogram showing mouse PBMC cells stained with (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-TBX21 Antibody (1µg/1x10<sup>6</sup> cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (5-10µg/1x10<sup>6</sup> cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1µg/1x10<sup>6</sup>) used under the same conditions. Unlabelled sample (Red line) was also used as a control.