bsm-54494R

[Primary Antibody]

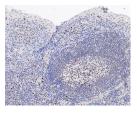
phospho-V-Myb+C-Myb (Ser11) Recombinant Rabbit mAb



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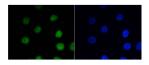
- DATASHEET -			400-901-9800
Host: Rab	bit Isoty	pe: IgG	Applications: WB (1:100-500)
Clonality: Rec	ombinant		IHC-P (1:100-500) IHC-F (1:400-800)
GenelD: 460	2 SWI	SS: P10242	IF (1:50-200)
Target: V-Myb+C-Myb (Ser11)			ICC/IF (1:50-200)
Purification: affinity purified by Protein A			Reactivity: Human
Concentration: 1mg/ml Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%			
Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.			Predicted MW.: ^{84 kDa}
Background: Transcriptional activator; DNA-binding protein that specifically recognize the sequence 5'-YAAC[GT]G-3'. Plays an important role in the control of proliferation and differentiation of hematopoietic			Subcellular Location:

– VALIDATION IMAGES



progenitor cells.

Immunohistochemical analysis of paraffinembedded human tonsil tissue using anti-P-V-Myb+C-Myb(S11) antibody. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the primary antibody (bsm-54494R, 1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.



ICC staining of P-V-Myb+C-Myb(S11) in AGS cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (bsm-54494R, 1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).