

**bs-17221R****[ Primary Antibody ]****phospho-TRIM28 (Ser473) Rabbit pAb****Bioss**  
**ANTIBODIES**

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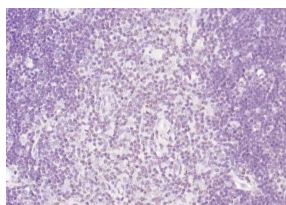
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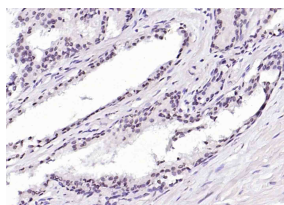
400-901-9800

**— DATASHEET —**

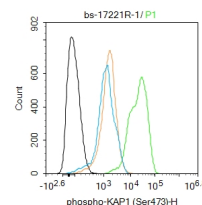
<b>Host:</b> Rabbit	<b>Isotype:</b> IgG	<b>Applications:</b> IHC-P (1:100-500)
<b>Clonality:</b> Polyclonal		<b>IHC-F</b> (1:100-500)
<b>GeneID:</b> 10155	<b>SWISS:</b> Q13263	<b>IF</b> (1:100-500)
<b>Target:</b> TRIM28 (Ser473)		<b>Flow-Cyt</b> (1ug/Test)
<b>Immunogen:</b> KLH conjugated synthesised phosphopeptide derived from human TRIM28 around the phosphorylation site of Ser473: SR(p-S)GE.		<b>Reactivity:</b> Human, Mouse, Rat (predicted: Rabbit, Macaque Monkey, Gorilla)
<b>Purification:</b> affinity purified by Protein A		<b>Predicted MW.:</b> 88 kDa
<b>Concentration:</b> 1mg/ml		<b>Subcellular Location:</b> Nucleus
<b>Storage:</b> 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.		
<b>Background:</b> The protein encoded by this gene mediates transcriptional control by interaction with the Kruppel-associated box repression domain found in many transcription factors. The protein localizes to the nucleus and is thought to associate with specific chromatin regions. The protein is a member of the tripartite motif family. This tripartite motif includes three zinc-binding domains, a RING, a B-box type 1 and a B-box type 2, and a coiled-coil region. [provided by RefSeq, Jul 2008]		

**— VALIDATION IMAGES —**

Paraformaldehyde-fixed, paraffin embedded (mouse thymus); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (phospho-KAP1 (Ser473)) Polyclonal Antibody, Unconjugated (bs-17221R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human prostate); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (phospho-KAP1 (Ser473)) Polyclonal Antibody, Unconjugated (bs-17221R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control (black line) :A431. Primary Antibody (green line): Rabbit Anti-phospho-KAP1 (Ser473) antibody (bs-17221R) Dilution:1ug/Test; Secondary Antibody (white/blue line) : Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line) : Normal Rabbit IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C. The cells were then incubated in 5% BSA to block non-specific protein-protein interactions for 30 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

**— SELECTED CITATIONS —**

- **[IF=8.2]** Zhao Huakan. et al. Single-cell analysis of post-translational modifications identifies immunosuppressive macrophage subtypes in the HBV-positive hepatocellular carcinoma microenvironment. CANCER IMMUNOL RES. 2025 Jun.; WB ;Human. 40455063