## bsm-52141R

## [ Primary Antibody ]

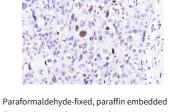
## phospho-c-Jun (Ser63) Recombinant Rabbit mAb A N T | B



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- DATASHEET		400-901-9800
Host: Rabbit	<b>Isotype:</b> IgG	Applications: WB (1:500-2000)
Clonality: Recombinant	CloneNo.: 6C2	<b>IHC-P</b> (1:50-100) <b>IHC-F</b> (1:50-100)
GenelD: 3725	SWISS: P05412	<b>IF</b> (1:50-200)
Target: c-Jun (Ser63)		ICC/IF (1:50-200)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human c-Jun around the phosphorylation site of Ser63: LT(p-S)PD.		<b>Reactivity:</b> Human, Mouse (predicted: Rat)
Purification: affinity purified by Pro	otein A	
Concentration: 1mg/ml Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%		Predicted MW.: <sup>36 kDa</sup>
Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Location: Nucleus
factor AP-1 that binds elements and appears SAPK/JNK signaling p is regulated by phosp signals including grov UV irradiation stimula activate c-Jun depend renders c-Jun nonres signaling pathways. T to the N-terminal regi	jun) is a component of the transcription and activates transcription at TRE/AP-1 s to be a major downstream target of the athway. The transcriptional activity of c-Jun horylation at Ser63 and Ser73. Extracellular <i>t</i> th factors, transforming oncoproteins and te phosphorylation of c-Jun at Ser63/73 and dent transcription. Mutation of Ser63/73 ponsive to mitogenic and stress induced he MAP kinase homologue, SAPK/JNK, binds on of c-Jun and phosphorylates c-Jun at the activity of SAPK/JNK is stimulated by the vate c-Jun.	
- VALIDATION IMAGES		

Sample: Lane 1: NIH/3T3 (Mouse) Cell Lysate at 30 ug Lane 2: Hela (Human) Cell Lysate at 30 ug Lane 3: SH-SY5Y (Human) Cell Lysate at 30 ug Lane 4: U251 (Human) Cell Lysate at 30 ug Primary: Anti-phospho-c-Jun (Ser63) (bsm-52141R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 36/43 kD Observed band size: 43 kD



(human laryngeal carcinoma); 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-c-Jun (Ser63)) Monoclonal Antibody, Unconjugated (bsm-52141R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



by operating according to SP Kit(Rabbit)

(sp-0023) instructionsand DAB staining.

## - SELECTED CITATIONS -

• [IF=5.3] Caixia Tan. et al.Mcl-1 downregulation enhances BCG treatment efficacy in bladder cancer by promoting macrophage polarization.Cancer Cell International.2025 Feb 15;25(1):48.; 39955585