

bsm-52123R**[Primary Antibody]****BioSS**
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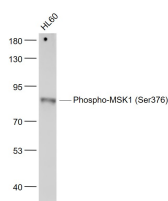
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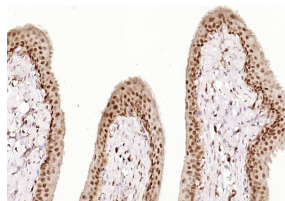
400-901-9800

Phospho-MSK1 (Ser376) Recombinant Rabbit mAb**— DATASHEET —****Host:** Rabbit**Isotype:** IgG**Clonality:** Recombinant**CloneNo.:** 11A1**GeneID:** 9252**SWISS:** O75582**Target:** Phospho-MSK1 (Ser376)**Immunogen:** KLH conjugated synthesised phosphopeptide derived from human MSK1 around the phosphorylation site of Ser376: GY(p-S)FV .**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

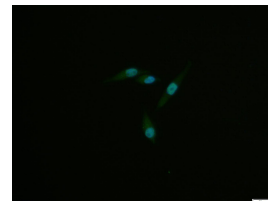
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: MSK1 is a mitogen and stress activated protein kinase 1 which belongs to the AGC family of kinases and is related in structure to the ribosomal p70 S6 kinase subfamily. MSK1 can be activated by ERK1/2 and SAPK2/p38 MAP kinase. It is also known to be required for the phosphorylation of CREB, ATF1 H3 and HMG14 in response to mitogen and stress. Similar to RSK, MSK1 contains two kinase domains (N term and a C term). Once phosphorylated on Thr581 and Ser360 by ERK1/2 and SAPK2/p38, MSK1 autophosphorylates on at least 5 sites. Of these autophosphorylation sites Ser212 and Ser376 get phosphorylated by the C terminal kinase domain of MSK1 which is essential for the catalytic activity of the N terminal kinase domain.**Applications:** WB (1:500-2000)**IHC-P** (1:20-200)**IHC-F** (1:20-200)**IF** (1:20-200)**Flow-Cyt** (2ug/Test)**ICC/IF** (1:100)**Reactivity:** Human, Rat**Predicted MW.:** 90 kDa**Subcellular Location:** Cytoplasm ,Nucleus**— VALIDATION IMAGES —**

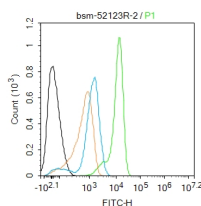
Sample: HL60(Human) Cell Lysate at 30 ug
 Primary: Anti- Phospho-MSK1 (Ser376)
 (bsm-52123R) at 1/1000 dilution Secondary:
 IRDye800CW Goat Anti-Rabbit IgG at 1/20000
 dilution Predicted band size: 90 kD Observed
 band size: 90 kD



Paraformaldehyde-fixed, paraffin embedded (rat bladder); 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-MSK1 (Ser376)) Polyclonal Antibody, Unconjugated (bsm-52123R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



SH-SY5Y cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (Phospho-MSK1 (Ser376)) monoclonal Antibody, Unconjugated (bsm-52123R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control: U-937. Primary Antibody (green line): Rabbit Anti-Phospho-MSK1

Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

(Ser376)antibody (bsm-52123R)

Dilution:2ug/Test; Secondary Antibody : Goat anti-rabbit IgG-FITC Dilution: 0.5ug/Test.

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.