

bs-16678R**[Primary Antibody]****phospho-HSF1 (Thr142) Rabbit pAb****Bioss**
ANTIBODIES

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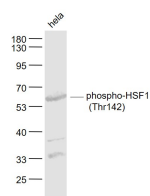
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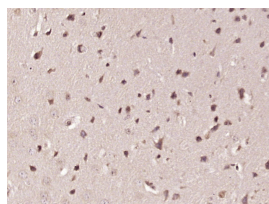
400-901-9800

— DATASHEET —

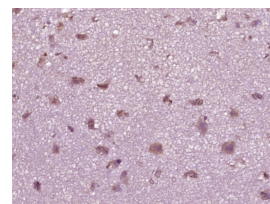
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (2ug/Test)
Clonality: Polyclonal		
GeneID: 3297	SWISS: Q00613	
Target: HSF1 (Thr142)		
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human HSF1 around the phosphorylation site of Thr142: LL(p-T)DV.		Reactivity: Human, Mouse (predicted: Rat, Pig, Cow, Dog)
Purification: affinity purified by Protein A		
Concentration: 1mg/ml		Predicted MW.: 57 kDa
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.		Subcellular Location: Cytoplasm ,Nucleus
Background: The product of this gene is a heat-shock transcription factor. Transcription of heat-shock genes is rapidly induced after temperature stress. Hsp90, by itself and/or associated with multichaperone complexes, is a major repressor of this gene. [provided by RefSeq, Jul 2008].		

— VALIDATION IMAGES —

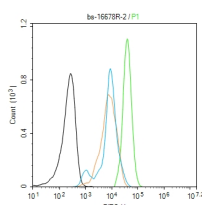
Sample: HeLa(Human) Cell Lysate at 30 ug
Primary: Anti- phospho-HSF1 (Thr142)
(bs-16678R) at 1/1000 dilution Secondary:
IRDye800CW Goat Anti-Rabbit IgG at 1/20000
dilution Predicted band size: 57 kD Observed
band size: 57 kD



Paraformaldehyde-fixed, paraffin embedded
(mouse brain tissue); 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 (HSF1 (Thr142)) Polyclonal
Antibody, Unconjugated (bs-16678R) at 1:400
overnight at 4°C, followed by operating
according to SP Kit(Rabbit) (sp-0023)
instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded
(human brain glioma); 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 (HSF1 (Thr142))
Polyclonal Antibody, Unconjugated (bs-16678R)
at 1:400 overnight at 4°C, followed by operating
according to SP Kit(Rabbit) (sp-0023)
instructions and DAB staining.



Blank control (black line) :HeLa. Primary
Antibody (green line): Rabbit Anti-phospho-HSF1
(Thr142) antibody (bs-16678R) Dilution:2ug/Test;
Secondary Antibody (white/blue line) : Goat
anti-rabbit IgG-FITC 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

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

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.