– DATASHEET –

[Primary Antibody]

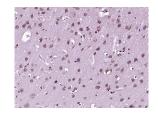
phospho-FoxO1 (Ser256) Rabbit pAb



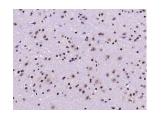
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- DATASHEET			
Host: Rabbit	Isotype: IgG	Applications: IHC-P (1:100-500)	
Clonality: Polyclonal		IHC-F (1:100-500) IF (1:100-500)	
GenelD: 2308	SWISS: Q12778	Flow-Cyt (1µg /Test)	
Target: FoxO1	(Ser256)	ELISA (1:5000-10000)	
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human FOXO1 around the phosphorylation site of Ser256: AA(p-S)MD.		Reactivity: Human, Mouse, Rat, Rabbit (predicted: Pig, Cow, Chicken, Dog, Horse)	
Purification: affinity purified by Protein A			
Concentration: 1mg/ml		Predicted	
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		Predicted MW.: ^{72 kDa} Subcellular Location: ^{Cytoplasm} , Nucleus	
freeze,			
which specifi howev Translo	ene belongs to the forkhead family of transcription factors are characterized by a distinct forkhead domain. The c function of this gene has not yet been determined; er, it may play a role in myogenic growth and differentiatio ocation of this gene with PAX3 has been associated with ar rhabdomyosarcoma. [provided by RefSeq].	m.	

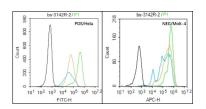
– VALIDATION IMAGES



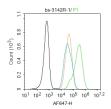
Paraformaldehyde-fixed, paraffin embedded (Rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-FoxO1 (Ser256)) Polyclonal Antibody, Unconjugated (bs-3142R) 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 (mouse brain); 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-FoxO1 (Ser256)) Polyclonal Antibody, Unconjugated (bs-3142R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Black line : Positive blank control (Hela); Negative blank control (Molt4) Green line : Primary Antibody (Rabbit Anti-Phospho-FoxO1 (Ser256) antibody (bs-3142R)) Orange line: Isotype Control Antibody (Rabbit IgG). Blue line : Secondary Antibody (Goat anti-rabbit IgG-AF488) Hela (Positive) and Molt4 (Negative control) cells (black) were fixed with 4% PFA for 10min at room temperature, permeabilized with 90% ice-cold methanol for 20 min at -20°C, and incubated in 5% BSA blocking buffer for 30 min at room temperature. Cells were then stained with Phospho-FoxO1 (Ser256) Antibody(bs-3142R)at 1:50 dilution in blocking buffer and incubated for 30 min at room temperature, washed twice with 2% BSA in PBS, followed by secondary antibody(blue) incubation for 40 min at room temperature. Acquisitions of 20,000 events were performed. Cells stained with primary antibody (green), and isotype control (orange).



Blank control: HepG2. Primary Antibody (green line): Rabbit Anti-Phospho-FoxO1 (Ser256) antibody (bs-3142R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF647 Dilution: 1µg /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 nonspecific 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 -

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