bs-1408R

- DATASHEET -

[Primary Antibody]

SMARCA4 Rabbit pAb



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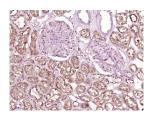
> IHC-F (1:100-500) **IF** (1:100-500) Flow-Cyt (3ug/test)

(predicted: Mouse, Dog)

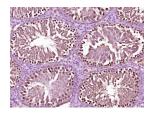
DATASITL	- 1			
Host:	Rabbit	lsotype: IgG	Applications	IHC-P (1:100-500)
Clonality:	Polyclonal			IHC-F (1:100-500) IF (1:100-500)
GenelD:	6597	SWISS: P51532		Flow-Cyt (3ug/test
Target: SMARCA4			Reactivity	Human, Rat
Immunogen:	KLH conjugated syn 51-167/1647.	thetic peptide derived from human SMARC		(predicted: Mouse,
Purification :	affinity purified by P	Protein A		
Concentration :	1mg/ml	Predicted MW.:	185 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:	Nucleus
Background:	family of proteins ar Drosophila. Member activities and are the by altering the chron encoded protein is p remodeling complex	d by this gene is a member of the SWI/SNF nd is similar to the brahma protein of rs of this family have helicase and ATPase ought to regulate transcription of certain g matin structure around those genes. The part of the large ATP-dependent chromatin x SNF/SWI, which is required for transcripti	onal	

activation of genes normally repressed by chromatin. In addition, this protein can bind BRCA1, as well as regulate the expression of the tumorigenic protein CD44. Alternatively spliced transcripts have been found for this gene, but their full-length natures have not been determined.

- VALIDATION IMAGES



Paraformaldehyde-fixed, paraffin embedded (Human kidney); 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 (BRG-1) Polyclonal Antibody, Unconjugated (bs-1408R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Rat testis); 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 (BRG-1) Polyclonal Antibody, Unconjugated (bs-1408R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.

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Blank control:A549. Primary Antibody (green line): Rabbit Anti-BRG-1 antibody (bs-1408R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-PE Dilution: 3µg /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 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=5.08] Chen, Luyue, et al. "SNORD76, a box C/D snoRNA, acts as a tumor suppressor in glioblastoma." Scientific reports 5 (2015). WB ;Human. 25715874