### bs-1133R

## [ Primary Antibody ]

# SPARC Rabbit pAb

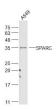


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- DATASHEET		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)
GenelD: 6678	SWISS: P09486	<b>IF</b> (1:100-500)
Target: SPARC		Flow-Cyt (2ug/Test)
Immunogen: KLH conjugated synthetic peptide derived from human SPARC: 101-200/303.		<b>Reactivity:</b> Human, Mouse, Rat (predicted: Rabbit, Pig,
Purification: affinity purified by F	rotein A	Cow, Dog, Horse)
Concentration: 1mg/ml		Predicted MW.: <sup>33 kDa</sup>
<b>Storage:</b> 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 Secreted ,Extracellular Location: matrix ,Nucleus
The encoded protei become calcified bu synthesis and prom	o cysteine-rich acidic matrix-associated protein. n is required for the collagen in bone to t is also involved in extracellular matrix otion of changes to cell shape. The gene asociated with tumor suppression but has also	

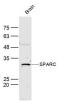
been correlated with metastasis based on changes to cell shape which can promote tumor cell invasion. [provided by RefSeq, Dec

### - VALIDATION IMAGES



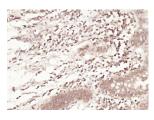
2011].

Sample: A549(Human) Cell Lysate at 30 ug Primary: Anti-SPARC (bs-1133R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 33 kD Observed band size: 35 kD

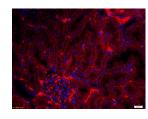


Sample: Brain (Mouse) Lysate at 40 ug Primary: Anti-SPARC (bs-1133R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 33 kD Observed band size: 33 kD Paraformaldehyde-fixed, paraffin embedded (human colon carcinoma): Antiren retrieval b

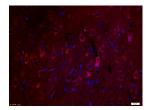
(human colon carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (SPARC) Polyclonal Antibody, Unconjugated (bs-1133R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



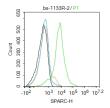
Paraformaldehyde-fixed, paraffin embedded (human rectal carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min;



Tissue/cell: mouse kidney tissue;4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-SPARC



Tissue/cell: rat brain tissue;4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-SPARC Antibody incubation with (SPARC) Polyclonal Antibody, Unconjugated (bs-1133R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining. Polyclonal Antibody, Unconjugated(bs-1133R) 1:200, overnight at 4°C; The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C. DAPI(Sug/ml,blue,C-0033) was used to stain the cell nuclei Polyclonal Antibody, Unconjugated(bs-1133R) 1:200, overnight at 4°C; The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C. DAPI(5ug/ml,blue,C-0033) was used to stain the cell nuclei



Blank control:U87MG. Primary Antibody (green line): Rabbit Anti-SPARC antibody (bs-1133R) Dilution: 2ug/Test; Secondary Antibody : Goat anti-rabbit IgG-FITC Dilution: 0.5ug/Test. Protocol The cells were 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.

### - SELECTED CITATIONS -

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- [IF=4.547] Zheng, Qingbo. et al. Construction of transcriptome atlas of white yak hair follicle during anagen and catagen using single-cell RNA sequencing. BMC GENOMICS. 2022 Dec;23(1):1-14 IHC ;Yak. 36482306
- [IF=3.73] Chen, Jie, et al. "SPARC is a key regulator of proliferation, apoptosis and invasion in human ovarian cancer." PLoS One 7.8 (2012): e42413. IHC ;="Human". 22879971
- [IF=4.2] Camila Machado Baldavira. et al. Matrisome analysis of NSCLC unveils clinically-important cancer-associated extracellular matrix changes. BBA-MOL BASIS DIS. 2025 Apr;1871:167709 IHC ;Human. 39938232
- [IF=1.5] Kurtul, Neslihan, et al. "Prognostic Value of SPARC Expression in Unresectable NSCLC Treated with Concurrent Chemoradiotherapy." Asian Pacific Journal of Cancer Prevention 15.20 (2014): 8911-8916. IHC ;="Human". 25374228