## bs-1133R

# [ Primary Antibody ]

www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

# SPARC Rabbit pAb

DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

**GenelD:** 6678 **SWISS:** P09486

Target: SPARC

Immunogen: KLH conjugated synthetic peptide derived from human SPARC:

101-200/303.

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: This gene encodes a cysteine-rich acidic matrix-associated protein.

The encoded protein is required for the collagen in bone to become calcified but is also involved in extracellular matrix synthesis and promotion of changes to cell shape. The gene product has been associated 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

2011].

Applications: WB (1:500-2000)

**IHC-P** (1:100-500) IHC-F (1:100-500) **IF** (1:100-500) Flow-Cyt (2ug/Test)

Reactivity: Human, Mouse, Rat

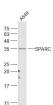
(predicted: Rabbit, Pig, Cow, Dog, Horse)

Predicted MW.: 33 kDa

Subcellular Secreted ,Extracellular

Location: matrix, Nucleus

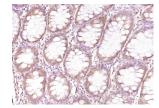
## VALIDATION IMAGES



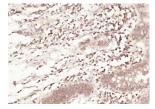
Sample: A549(Human) Cell Lysate at 30 ug Primary: Anti-SPARC (bs-1133R) at 1/1000 dilution Secondary: IRDve800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 33 kD Observed band size: 35 kD



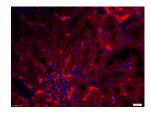
Sample: Brain (Mouse) Lysate at 40 ug Primary: Anti-SPARC (bs-1133R) at 1/300 dilution Secondary: IRDve800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 33 kD Observed band size: 33 kD



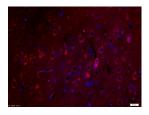
Paraformaldehyde-fixed, paraffin embedded (human colon carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (SPARC) Polyclonal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

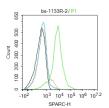


Tissue/cell: rat brain tissue:4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-SPARC

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) instructions and 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(5ug/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 -

- [IF=7.097] Liting Cheng. et al. Bioresponsive micro-to-nano albumin-based systems for targeted drug delivery against complex fungal infections. Acta Pharm Sin B. 2021 May;: IF; Mouse. 10.1016/j.apsb.2021.04.020
- [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