## bs-6577R

## [ Primary Antibody ]

# Bioss ANTIBODIES

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# S100P binding protein Rabbit pAb

- DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GenelD: 64766 SWISS: Q96BU1

Target: S100P binding protein

Immunogen: KLH conjugated synthetic peptide derived from human S100PBP:

331-408/408.

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:** S100PBP was originally cloned from a pancreatic epithelioid

carcinoma library and encodes a predicted 408 amino acid protein. RT-PCR detected S100PBP expression in brain, breast, spleen, and lung, but not in pancreas and liver. GFP-tagged S100PBP localized

to nuclei of transfected HeLa cells.

**Applications: WB** (1:500-2000)

Flow-Cyt (1ug/Test)

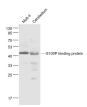
Reactivity: Human, Mouse

(predicted: Rat)

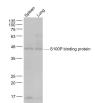
Predicted MW.: 46 kDa

Subcellular Location: Nucleus

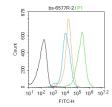
#### VALIDATION IMAGES



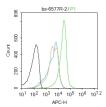
Sample: Molt-4 (Human) Lysate at 40 ug Cerebellum (Mouse) Lysate at 40 ug Primary: Anti- S100P binding protein (bs-6577R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 46 kD Observed band size: 46 kD



Sample: Spleen (Mouse) Lysate at 40 ug Lung (Mouse) Lysate at 40 ug Primary: Anti- S100P binding protein (bs-6577R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 46 kD Observed band size: 46 kD



Blank control: Mouse spleen. Primary Antibody (green line): Rabbit Anti-S100P binding protein antibody (bs-6577R) Dilution: 2µg/10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF488R 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.



Blank control: Mouse spleen. Primary Antibody (green line): Rabbit Anti-S100P binding protein antibody (bs-6577R) Dilution: 2µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit

IgG . Secondary Antibody : Goat anti-rabbit IgG-AF647 Dilution:  $1\mu g$  /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 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=2.677] Yu Chen. et al. Effect of necrostatin-1 on sciatic nerve crush injury in rat models. J ORTHOP SURG RES. 2023 Dec;18(1):1-9 IF; Rat. 36717933