

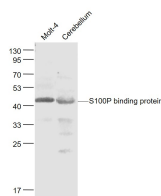
**bs-6577R****[ Primary Antibody ]****S100P binding protein Rabbit pAb****Bioss**  
**ANTIBODIES**

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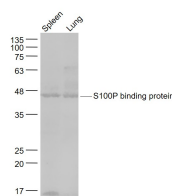
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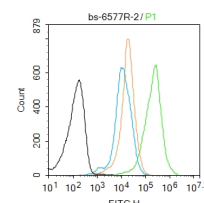
400-901-9800

**DATASHEET****Host:** Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 64766**SWISS:** Q96BU1**Target:** S100P binding protein**Immunogen:** KLH conjugated synthetic peptide derived from human S100BP: 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:** S100BPB was originally cloned from a pancreatic epithelioid carcinoma library and encodes a predicted 408 amino acid protein. RT-PCR detected S100BPB expression in brain, breast, spleen, and lung, but not in pancreas and liver. GFP-tagged S100BPB 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**

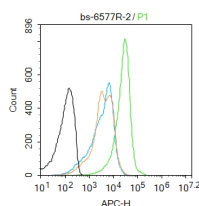
Sample: Molt-4 (Human) Lysate at 40 ug  
Cerebellum (Mouse) Lysate at 40 ug  
Primary: Anti- S100P binding protein (bs-6577R) at 1/1000  
dilution  
Secondary: IRDye800CW Goat Anti-  
Rabbit IgG at 1/20000 dilution  
Predicted band size: 46 kD  
Observed band size: 46 kD



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<sup>6</sup> 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 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.



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

Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

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 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=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