

bs-3567R**[Primary Antibody]****Bioss**
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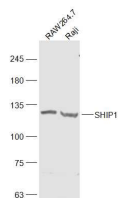
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SHIP1 Rabbit pAb**— DATASHEET —**

Host: Rabbit Clonality: Polyclonal GeneID: 3635 Target: SHIP1 Immunogen: KLH conjugated synthetic peptide derived from human SHIP1: 701-800/1189. 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: SHIP1 is a member of the inositol polyphosphate-5-phosphatase (INPP5) family and contains an N-terminal SH2 domain, an inositol phosphatase domain, and two C-terminal protein interaction domains. Expression of this protein is restricted to hematopoietic cells where its movement from the cytosol to the plasma membrane is mediated by tyrosine phosphorylation in response to multiple cytokine and B and T cell receptor activation. At the plasma membrane, the protein hydrolyzes the 5' phosphate from phosphatidylinositol (3,4,5)-trisphosphate and inositol-1,3,4,5-tetrakisphosphate, thereby affecting multiple signaling pathways. Overall the protein functions as a negative regulator of myeloid cell proliferation and survival.	Isotype: IgG SWISS: Q92835	Applications: WB (1:500-2000) Reactivity: Human, Mouse (predicted: Rat, Pig, Cow, Horse) Predicted MW.: 133 kDa Subcellular Location: Cell membrane ,Cytoplasm
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— VALIDATION IMAGES —

Sample: RAW264.7(Mouse) Cell Lysate at 30 ug
Raji(Human) Cell Lysate at 30 ug Primary: Anti-SHIP1 (bs-3567R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 133 kD Observed band size: 133 kD

— SELECTED CITATIONS —

- **[IF=6.376]** Wang, Wenxin. et al. Preparation of a miR-155-activating nucleic acid nanoflower to study the molecular mechanism of miR-155 in inflammation. MOL MED. 2022 Dec;28(1):1-17 WB ;Human. 35715753
- **[IF=4.784]** Wan B et al. Porcine FcγRIIb mediated PRRSV ADE infection through inhibiting IFN-β by cytoplasmic inhibitory signal transduction. Int J Biol Macromol. 2019 Jul 5;138:198-206. WB ;Monkey. 31284005
- **[IF=4.9]** Limei Wen. et al. Synergistic and toxicity-reducing effects of acteoside as an adjuvant therapy of oxaliplatin against hepatocellular carcinoma. INT J ONCOL. 2025 Jun;66(6):1-18 WB ;Mouse. 40341416

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