

bs-6023R**[Primary Antibody]**

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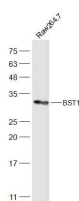
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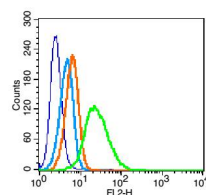
400-901-9800

BST1 Rabbit pAb**— DATASHEET —**

Host: Rabbit Clonality: Polyclonal Target: BST1 Immunogen: KLH conjugated synthetic peptide derived from human BST1/CD157: 51-150/318. 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: Bone marrow stromal cell antigen 1 (BST1) is a pleiotropic ectoenzyme which belongs to the CD38 family and to the growing number of leukocyte surface molecules known to act independently as both receptors and enzymes. The BST1 molecule displays two distinct domains in its extracellular component. The first is implicated in the enzymic activities of the molecule (it synthesizes cyclic ADP-ribose, a second messenger that elicits calcium release from intracellular stores) and the second domain has adhesion/signalling properties. Bone marrow stromal cell antigen 1 facilitates pre-B-cell growth. The deduced amino acid sequence exhibits 33% similarity with CD38. BST1 expression is enhanced in bone marrow stromal cell lines derived from patients with rheumatoid arthritis. The polyclonal B-cell abnormalities in rheumatoid arthritis may be, at least in part, attributed to BST1 overexpression in the stromal cell population.	Isotype: IgG GeneID: 683	Applications: WB (1:500-2000) Flow-Cyt (1µg/Test)
		Reactivity: Human, Mouse (predicted: Rat, Rabbit, Pig, Cow, Dog)
		Predicted MW: 33 kDa
		Subcellular Location: Cell membrane

— VALIDATION IMAGES —

Sample: Raw264.7(Mouse) Cell Lysate at 30 ug
 Primary: Anti-BST1 (bs-6023R) at 1/500 dilution
 Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 33 kD
 Observed band size: 33 kD



Blank control: U937(blue). Primary Antibody: Rabbit Anti-BST1 antibody(bs-6023R), Dilution: 1µg in 100 µL 1X PBS containing 0.5% BSA; Isotype Control Antibody: Rabbit IgG (orange), used under the same conditions. Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA. Protocol The cells were fixed with 2% paraformaldehyde (10 min). Primary antibody (bs-6023R, 1µg /1x10⁶ cells) were incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 10% goat serum (15 min) to block non-specific protein-protein interactions. Then the Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned above to react with the primary antibody at 1/200 dilution for 30 min on ice. Acquisition of 20,000 events was performed.