### bs-2782R

# [ Primary Antibody ]

# CD44v5 Rabbit pAb

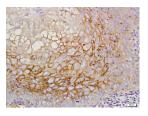


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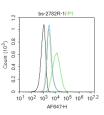
– DATASHEET –––––		400-901-9800
Host: Rabbit	<b>lsotype:</b> lgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		<b>IHC-P</b> (1:100-500) <b>IHC-F</b> (1:100-500)
<b>GenelD:</b> 960	SWISS: P16070	<b>IF</b> (1:100-500)
Target: CD44v5		Flow-Cyt (1µg /Test)
Immunogen: KLH conjugated synthetic peptide derived from human CD44v5: 61-139/139.		Reactivity: Human
Purification: affinity purified by	Protein A	
Concentration: 1mg/ml		Predicted
<b>Storage:</b> 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.		Predicted MW.: <sup>85 kDa</sup> Subcellular Location: <sup>Cell</sup> membrane
involved in cell-cell receptor for hyaluro ligands, such as ost metalloproteinases variety of cellular fu recirculation and h Transcripts for this results in many fun length nature of so Alternative splicing	ed by this gene is a cell-surface glycoprotein interactions, cell adhesion and migration. It is a pric acid (HA) and can also interact with other ecopontin, collagens, and matrix (MMPs). This protein participates in a wide unctions including lymphocyte activation, oming, hematopoiesis, and tumor metastasis. gene undergo complex alternative splicing that ctionally distinct isoforms, however, the full me of these variants has not been determined. is the basis for the structural and functional tein, and may be related to tumor metastasis. q, Jul 2008].	
- VALIDATION IMAGES		



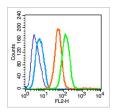
Sample: Lane 1: Huvec (Human) Cell Lysate at 30 ug Lane 2: A549 (Human) Cell Lysate at 30 ug Primary: Anti-CD44v5 (bs-2782R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 110-180 kD Observed band size: 110-180 kD



Tissue/cell: human laryngo carcinoma; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0 ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-CD44v5 Polyclonal Antibody, Unconjugated(bs-2782R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control: HELA. Primary Antibody (green line): Rabbit Anti-CD44v5 antibody (bs-2782R) Dilution: 1µg /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 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(blue): U251 (fixed with Ice-cold 70% ethanol overnight at 4 °C). Primary Antibody:Rabbit Anti-CD44v5 antibody (bs-2782R,Green); 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 antirabbit IgG-FITC(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA.

## - SELECTED CITATIONS -

• [IF=3.2] Hao Chen. et al.CD44v6-mediated regulation of gastric cancer stem cells: a potential therapeutic target.clinical and experimental medicine.2025 Mar 12;25(1):80. IHC ;Human. 40069421