## bs-23550R

DATACHEET

## [ Primary Antibody ]

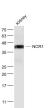
# NCR1 Rabbit pAb



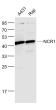
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

– DATASHEFT ––––––		
Host: Rabbit	<b>Isotype:</b> IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)
<b>GenelD:</b> 9437	SWISS: 076036	<b>IF</b> (1:100-500)
Target: NCR1		Flow-Cyt (2ug/Test)
Immunogen: KLH conjugated synthetic peptide derived from human NCR1: 1-100/304. < Extracellular >		Reactivity: Human, Mouse
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.		Predicted MW.: <sup>31 kDa</sup>
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Location: Cell membrane
characterized fam be major triggerin glycoprotein that i a ~40 amino acid r domain, and a sho represent a novel	xicity receptors (NCRs) are a recently ly of Ig-like activation receptors that appear to g receptors in tumor cell recognition. NCR1 is a has two extracellular Ig-like domains followed by esidue stalk region, a type I transmembrane rt cytoplasmic tail. NCR1 has been shown to NK cell-specific molecule involved in human NK R1 has been implicated in NK cell-mediated lysis	

#### - VALIDATION IMAGES

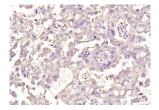


Sample: Kidney (Mouse) Lysate at 40 ug Primary: Anti- NCR1 (bs-23550R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 31 kD Observed band size: 46 kD

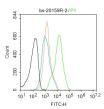


of several autologous tumor cells and pathogen-infected cell lines.

Sample: A431 (Human) Cell Lysate at 30 ug Raji (Human) Cell Lysate at 30 ug Primary: Anti- NCR1 (bs-23550R) at 1/500 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 31 kD Observed band size: 46 kD



Paraformaldehyde-fixed, paraffin embedded (human skin cancer); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (NCR1) Polyclonal Antibody, Unconjugated (bs-23550R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control:Mouse spleen. Primary Antibody (green line): Rabbit Anti-NCR1 antibody (bs-23550R) Dilution: 2µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF488  $\underbrace{\frac{1}{10^{2}}}_{FTCH} \underbrace{\frac{1}{10^{2}}}_{FTCH} \underbrace{\frac{1}{10^{2}}}_{FTC$ 

Blank control:Mouse kidney. Primary Antibody (green line): Rabbit Anti-NCR1 antibody (bs-23550R) Dilution: 2µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF488 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. 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 -----

- [IF=7.658] Xin Fang. et al. IDO1 can impair NK cells function against non-small cell lung cancer by downregulation of NKG2D Ligand via ADAM10. Pharmacol Res. 2022 Mar;177:106132 IHC ;MOUSE. 10.1016/j.phrs.2022.106132
- [IF=5.6] Na Qu. et al. Methionine enkephalin inhibited cervical cancer migration as well as invasion and activated CD11b+ NCR1+ NKs of tumor microenvironment. INT IMMUNOPHARMACOL. 2023 Nov;124:110967 IF ;MOUSE. 37741126
- [IF=2.1] Yating Ge. et al.Development of a Diagnostic Model for Focal Segmental Glomerulosclerosis: Integrating Machine Learning on Activated Pathways and Clinical Validation..Int J Gen Med.2025 Feb 26:18:1127-1142. IHC ;Human. 40026806