### bs-2200R

## [ Primary Antibody ]

# LDL Rabbit pAb



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– DATASHEET ———		400-901-9800
Host: Rabbit	<b>Isotype:</b> IgG	Applications: IHC-P (1:100-500)
Clonality: Polyclonal		<b>IHC-F</b> (1:100-500) <b>IF</b> (1:100-500)
GenelD: LDL		Flow-Cyt (3ug/test)
Target: LDL		Reactivity: Human (predicted: Mouse)
Purification: affinity purified by P	rotein A	Reactivity: Human (predicted. Mouse)
Concentration: 1mg/ml		
<b>Storage:</b> 0.01M TBS (pH7.4) w Glycerol. Shipped at 4°C. Stor freeze/thaw cycles.	ith 1% BSA, 0.02% Proclin300 and 50 e at -20°C for one year. Avoid repeate	% Subcellular Location: <sup>Cell</sup> membrane
Background: LDL (low-density lip- transports cholester tissues. LDL enabless based solution of th synthesis at these si system coordinates component of the pi Study of this system cellular basis of cho focus an important in process of receptor- the juxtamembrano participates in prote density lipoprotein in been shown that the receptors. Endocyto GB virus C/hepatitis was shown to be me	poprotein) is a type of lipoprotein that ol and triglycerides from the liver to fats and cholesterol to move within e blood stream. LDL also regulates ch tes. The low density lipoprotein (LDL the metabolism of cholesterol, an es asma membrane of all mammalian of has led to an enhanced understandi esterol homeostasis. It has also brou nechanism of metabolic regulation	peripheral the water nolesterol ) receptor sential cells. ng of the ught into the est that e low It has s viral cis C virus, rus (BVDV) cells.

### – VALIDATION IMAGES



Tissue/cell: human colon 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-LDL Polyclonal Antibody, Unconjugated(bs-2200R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: Human lung cancer; 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-LDL Polyclonal Antibody, Unconjugated(bs-2200R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control: A549. Primary Antibody (green line): Rabbit Anti-LDL antibody (bs-2200R) Dilution: 3µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-PE Dilution: 1µg /test. Protocol The cells were incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at 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=5.94] Armstrong, Susan M., et al. "A novel assay uncovers an unexpected role for SR-BI in LDL transcytosis." Cardiovascular Research (2015): cvv218. ELISA ;="Human". 26334034 • [IF=3.876] Wang, Xin-Yuan. et al. Accumulation of LDL/ox-LDL in the necrotic region participates in osteonecrosis of the femoral head: a pathological and in vitro study. Lipids Health Dis. 2021 Dec;20(1):1-13 IHC ;Human. 34823555