

bs-2336R**[Primary Antibody]**

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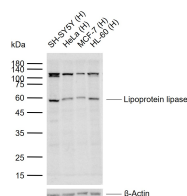
400-901-9800

Lipoprotein lipase Rabbit pAb

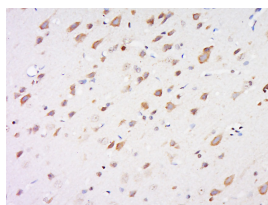
DATASHEET

Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (0.2µg/Test)
Clonality: Polyclonal		
GeneID: 4023	SWISS: P06858	
Target: Lipoprotein lipase		
Immunogen: KLH conjugated synthetic peptide derived from human LPL protein: 301-400/475.		
Purification: affinity purified by Protein A		Reactivity: Human, Rat (predicted: Mouse, Rabbit, Pig, Cow, Dog, Horse)
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: Lipoprotein lipase (LPL) is the central enzyme in plasma triglyceride hydrolysis and is secreted by macrophages in the subendothelial space. Evidence has been provided that LPL produced by macrophages in the vessel wall exerts proatherogenic effects. The atherogenic effects of LPL have been mainly attributed to its ability to favor lipid accumulation within macrophages present in the atherosclerotic lesion. Recently, it has also been shown that LPL promote the development of atherosclerosis through facilitation of monocyte adhesion to endothelial cells, stimulation of tumor necrosis factor alpha (TNF) secretion and induction of vascular smooth muscle cell proliferation.		
		Predicted MW.: 53 kDa
		Subcellular Location: Secreted ,Cell membrane ,Cytoplasm

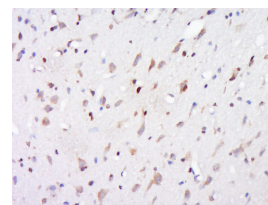
VALIDATION IMAGES



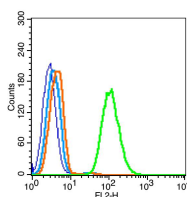
Sample: Lane 1: Human SH-SY5Y cell lysates
 Lane 2: Human HeLa cell lysates Lane 3: Human MCF-7 cell lysates Lane 4: Human HL-60 cell lysates
 Primary: Anti-Lipoprotein lipase (bs-2336R) at 1/500 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution
 Predicted band size: 53 kDa Observed band size: 60 kDa



Paraformaldehyde-fixed, paraffin embedded (rat brain); 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 (Lipoprotein lipase) Polyclonal Antibody, Unconjugated (bs-2336R) at 1:500 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



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Blank control (blue line): rapij(fixed with pre-warmed 4% paraformaldehyde for 30min at 37°C and then permeabilized with 90% ice-cold

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

methanol for 30 min on ice) Primary Antibody
(green line): Rabbit Anti-Lipoprotein lipase
antibody (bs-2336R), Dilution: 0.2µg /10⁶ cells;
Isotype Control Antibody (orange line): Rabbit
IgG . Secondary Antibody (white blue line): Goat
anti-rabbit IgG-PE, Dilution: 1µg /test.

— SELECTED CITATIONS —

- **[IF=2.752]** Xuchun Liu. et al. The Effect of FATP1 on Adipocyte Differentiation in Qinchuan Beef Cattle. Animals-Basel. 2021 Oct;11(10):2789 WB ;Bovine. 10.3390/ani11102789