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Datasheet for ABIN7158426 anti-LSR antibody (AA 112-233)

2 Images



Overview

Quantity:	100 µg
Target:	LSR
Binding Specificity:	AA 112-233
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This LSR antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Lipolysis-stimulated lipoprotein receptor protein (112-233AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

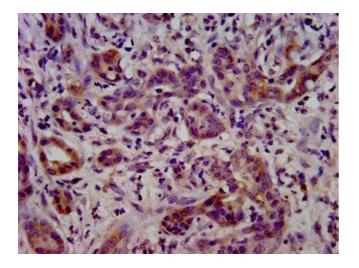
Target:	LSR
Alternative Name:	LSR (LSR Products)
Background:	Background: Probable role in the clearance of triglyceride-rich lipoprotein from blood. Binds
	chylomicrons, LDL and VLDL in presence of free fatty acids and allows their subsequent uptake

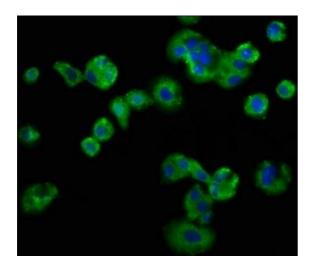
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	in the cells (By similarity).
	Aliases: ILDR3 antibody, Immunoglobulin like domain containing receptor 3 antibody, Lipolysis
	Stimulated Lipoprotein Receptor antibody, Lipolysis stimulated remnant antibody, Lipolysis-
	stimulated lipoprotein receptor antibody, LISCH antibody, LISCH protein antibody, LISCH7
	antibody, Liver specific bHLH Zip transcription factor antibody, LSR antibody, LSR_HUMAN
	antibody, MGC10659 antibody, MGC48312 antibody, MGC48503 antibody
UniProt:	Q86X29

Application Details

Application Notes:	Recommended dilution: IHC:1:200-1:500, IF:1:50-1:200,
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300
	Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be
	handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.





Immunohistochemistry

Image 1. IHC image of ABIN7158426 diluted at 1:400 and staining in paraffin-embedded human pancreatic cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunofluorescence

Image 2. Immunofluorescence staining of HepG2 cells with ABIN7158426 at 1:133, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

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