

Product datasheet

info@arigobio.com

ARG23652 anti-LPL / Lipoprotein Lipase antibody [5D2]

Package: 100 μg Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [5D2] recognizes LPL / Lipoprotein Lipase

Tested Reactivity Hu, Rat, Bov, Chk, Gpig

Species Does Not React With Ms

Tested Application ELISA, IP, WB

Host Mouse

Clonality Monoclonal

Clone 5D2 Isotype IgG1

Target Name LPL / Lipoprotein Lipase

Species Bovine

Immunogen Purified Bovine milk lipoprotein lipase.

Conjugation Un-conjugated

Alternate Names EC 3.1.1.34; LPL; Lipoprotein lipase; LIPD; HDLCQ11

Application Instructions

Application table	Application	Dilution
	ELISA	1:500 - 1:5000
	IP	Assay-dependent
	WB	Assay-dependent
Application Note	WB: This product detects a band of \sim 53 kDa using partially purified LPL from postheparin plasma. ELISA: This product can be used in an indirect ELISA, or as the capture antibody in a sandwich ELISA. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid	
Purification	Purification with Protein G.	
Buffer	PBS and 0.09% Sodium azide.	
Preservative	0.09% Sodium azide	
Concentration	1 mg/ml	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed	

before use.

Note

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol LPL

Gene Full Name lipoprotein lipase

Background LPL encodes lipoprotein lipase, which is expressed in heart, muscle, and adipose tissue. LPL functions as

a homodimer, and has the dual functions of triglyceride hydrolase and ligand/bridging factor for receptor-mediated lipoprotein uptake. Severe mutations that cause LPL deficiency result in type I hyperlipoproteinemia, while less extreme mutations in LPL are linked to many disorders of lipoprotein

metabolism. [provided by RefSeq, Jul 2008]

Function The primary function of this lipase is the hydrolysis of triglycerides of circulating chylomicrons and very

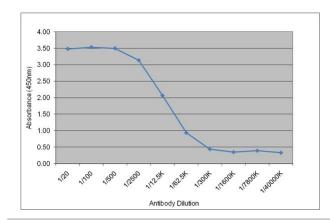
low density lipoproteins (VLDL). Binding to heparin sulfate proteogylcans at the cell surface is vital to the function. The apolipoprotein, APOC2, acts as a coactivator of LPL activity in the presence of lipids on

the luminal surface of vascular endothelium (By similarity). [UniProt]

Calculated Mw 53 kDa

PTM Tyrosine nitration after lipopolysaccharide (LPS) challenge down-regulates the lipase activity. [UniProt]

Images



ARG23652 anti-LPL / Lipoprotein Lipase antibody [5D2] ELISA image

ELISA: Bovine lipoprotein lipase detected with ARG23652 anti-LPL / Lipoprotein Lipase antibody [5D2] followed by Rabbit F(ab')2 anti-Mouse IgG (HRP).