



LPL Monoclonal Antibody

Catalog No	YP-mAb-02331
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	LPL
Immunogen	Purified recombinant fragment of LPL expressed in E. Coli.
Specificity	LPL Monoclonal Antibody detects endogenous levels of LPL protein.
Formulation	Ascitic fluid containing 0.03% sodium azide,0.5% BSA, 50%glycerol.
Source	Monoclonal, Mouse
Purification	Affinity purification
Dilution	Western Blot: 1/500 - 1/2000.
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	HDLCQ11;LIPD;lipoprotein lipase;LPL
Observed Band	53kD
Calculated Molecular Weight	53kD
Cell Pathway	Cell membrane ; Peripheral membrane protein ; Extracellular side . Secreted . Secreted, extracellular space, extracellular matrix . Newly synthesized LPL binds to cell surface heparan proteoglycans and is then released by heparanase. Subsequently, it becomes attached to heparan proteoglycan on endothelial cells (PubMed:27811232). Locates to the plasma membrane of microvilli of hepatocytes with triglyceride-rich lipoproteins (TRL). Some of the bound LPL is then internalized and located inside non-coated endocytic vesicles (By similarity). .
Tissue Specificity	Detected in blood plasma (PubMed:2340307, PubMed:11893776, PubMed:12641539). Detected in milk (at protein level) (PubMed:2340307).
Function	catalytic activity:Triacylglycerol + H(2)O = diacylglycerol + a carboxylate., disease:Defects in LPL are a cause of familial chylomicronemia [MIM:238600]; also known as hyperlipoproteinemia type I. Familial chylomicronemia is a recessive disorder usually manifesting in childhood. On a normal diet, patients often present with abdominal pain, hepatosplenomegaly, lipemia retinalis, eruptive xanthomata, and massive hypertriglyceridemia, sometimes complicated



with acute pancreatitis.,disease:Defects in LPL are the cause of lipoprotein lipase deficiency (LPL deficiency) [MIM:238600]. LPL deficiency leads to hypertriglyceridemia.,function:The primary function of this lipase is the hydrolysis of triglycerides of circulating chylomicrons and very low density lipoproteins (VLDL). The enzyme functions in the presence of apolipoprotein C-2 on the luminal surface of vascular endothelium.,online inform

Background

lipoprotein lipase(LPL) Homo sapiens 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],

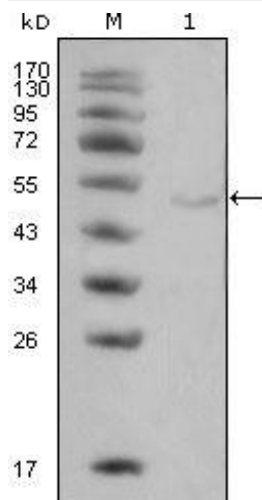
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis using LPL Monoclonal Antibody against HeLa cell lysate (1).