



LPL Monoclonal Antibody

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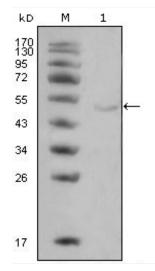


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	sometimes complicated with acute pancreatitis.,disease:Defects in LPL are the cause of lipoprotein lipase deficiency (LPL deficiency) [MIM:238600]. LPL deficiency leads to hypertriglyceridemia.,function:The primary function of this lipase is the hydrolysis 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).

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