

Datasheet for ABIN966473 anti-Lipase A antibody

Publications



Overview

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Quantity:	0.1 mg
Target:	Lipase A (LIPA)
Reactivity:	Human
Host:	Please inquire
Clonality:	Monoclonal
Conjugate:	This Lipase A antibody is un-conjugated
Application:	Western Blotting (WB), ELISA
Product Details	
Isotype:	lgG1
Specificity:	Ni-NTA purified truncated recombinant LAL expressed in E. Coli strain BL21 (DE3)
Purification:	Antibodies are purified by protein A affinity chromatography
Target Details	
Target:	Lipase A (LIPA)
Alternative Name:	LAL (LIPA Products)
Background:	Lysosomal acid lipase (LAL), with 378-amino acid protein(43-54 kDa), functions in the
	lysosome to catalyze the hydrolysis of cholesteryl esters and triglycerides which are taken up
	by receptor-mediated endocytosis. An inherited deficiency or low activity of human lysosomal
	acid lipase results in the intralysosomal storage of the respective lipid substrates. So it is also
	responsible for the rare conditions of Wolman disease and cholesteryl ester storage disease

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Application Details

Application Notes:	Western Blot: Dilution 1: 200- 1: 1,000
	ELISA: Propose dilution 1: 10,000.
	Determining optimal working dilutions by titration test.
Restrictions:	For Research Use only
Handling	
Storage:	-20 °C
Publications	
Product cited in:	Drebber, Andersen, Kasper, Lohse, Stolte, Dienes: "Severe chronic diarrhea and weight loss in
	cholesteryl ester storage disease: a case report." in: World journal of gastroenterology : WJG,
	Vol. 11, Issue 15, pp. 2364-6, (2005) (PubMed).
	Boldrini, Devito, Biselli, Filocamo, Bosman: "Wolman disease and cholesteryl ester storage
	disease diagnosed by histological and ultrastructural examination of intestinal and liver biopsy."
	in: Pathology, research and practice, Vol. 200, Issue 3, pp. 231-40, (2004) (PubMed).