

ASM rabbit pAb

Cat No.:ES11798

For research use only

Overview

Product Name	ASM rabbit pAb	
Host species	Rabbit	
Applications	WB;ELISA	
Species Cross-Reactivity	Human;Mouse	
Recommended dilutions	WB 1:500-2000 ELISA 1:5000-20000	
Immunogen	Synthesized peptide derived from part region of	
	human protein	
Specificity	ASM Polyclonal Antibody detects endogenous levels	
	of protein.	
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and	
	0.02% sodium azide.	
Storage	Store at -20 $^\circ\!\mathrm{C}$. Avoid repeated freeze-thaw cycles.	
Protein Name	Sphingomyelin phosphodiesterase (EC 3.1.4.12)	
	(Acid sphingomyelinase) (aSMase)	
Gene Name	SMPD1 ASM	
Cellular localization	Lysosome . Lipid droplet . Secreted . The secreted	
	form is induced in a time- and dose-dependent by	
	IL1B and TNF as well as stress and viral infection.	
	This increase of the secreted form seems to be due	
	to exocytosis of the lysosomal form and is	
	Ca(2+)-dependent (PubMed:20807762,	
	PubMed:22573858, PubMed:20530211). Secretion	
	is dependent of phosphorylation at Ser-510	
	(PubMed:17303575). Secretion is induced by	
	inflammatory mediators such as IL1B, IFNG or TNF as	
	well as infection with bacteria and viruses	
	(PubMed:12563314, PubMed:20807762)	
Purification	The antibody was affinity-purified from rabbit	
	antiserum by affinity-chromatography using	
	epitope-specific immunogen.	
Clonality	Polyclonal	
Concentration	1 mg/ml	
Observed band	69kD	



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Human Gene ID6609Human Swiss-Prot NumberP17405Alternative NamesThe pro

The protein encoded by this gene is a lysosomal acid sphingomyelinase that converts sphingomyelin to ceramide. The encoded protein also has phospholipase C activity. Defects in this gene are a cause of Niemann-Pick disease type A (NPA) and Niemann-Pick disease type B (NPB). Multiple transcript variants encoding different isoforms have been identified. [provided by RefSeq, Jul 2010],



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