Store at -200

ASM Antibody



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Applications: Reactivity: Sensitivity: MW (kDa): Source: **UniProt ID:** Entrez-Gene Id: WR Н Endogenous 57, 70 Rabbit #P17405 6609

Product Usage Application Dilution Information Western Blotting 1:1000

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at -**Storage**

20°C. Do not aliquot the antibody.

Specificity / Sensitivity ASM Antibody detects endogenous levels of total human ASM protein.

Species predicted to react based on 100% sequence homology: Monkey

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues at the carboxyl terminus of human ASM. Antibody was purified by protein A and peptide affinity chromatography.

Background

Sphingomyelinases (SMases) catalyze the hydrolysis of sphingomyelin to produce ceramide and phosphocholine (1). Ceramide is an important bioactive lipid triggering signal transduction involved in cell proliferation, apoptosis and differentiation (1,2). A number of SMases have been described and categorized based on their optimum pH activity, cation dependence, tissue distribution, and subcellular localization (1). These include a lysosomal acid SMase, a Zn++-dependent secreted acid SMase, a membrane-bound Mg++-dependent neutral SMase, a Mg++-independent neutral SMase, and an alkaline SMase.

Acid sphingomyelinase (ASM or SMPD1) is a lysosomal enzyme responsible for the hydrolysis of sphingomyelin to ceramide and phosphocholine. The ASM gene encodes three proteins, ASM-1, ASM-2, and ASM-3, of which ASM-1 is the only catalytically active enzyme (3,4). ASM-1 can exist as a 70 kDa form as well as a 57 kDa proteoytic product (5). Expression of ASM is induced during monocytic cell differentiation (6). Defects in the ASM gene are associated with type A and type B Niemann-Pick disease

Background References

- 1. Marchesini, N. and Hannun, Y.A. (2004) Biochem Cell Biol 82, 27-44.
- 2. Ruvolo, P.P. (2001) Leukemia 15, 1153-60.
- 3. Quintern, L.E. et al. (1989) EMBO J 8, 2469-73.
- 4. Schuchman, E.H. et al. (1991) J Biol Chem 266, 8531-9.
- 5. Ferlinz, K. et al. (1994) Biochem J 301 (Pt 3), 855-62.
- 6. Langmann, T. et al. (1999) J Lipid Res 40, 870-80.
- 7. Levran, O. et al. (1991) Proc Natl Acad Sci USA 88, 3748-52.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key WB: Western Blotting

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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Limited Uses

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