

SMPD1 Knockout HeLa Cell Lysate, Homozygous

Catalog No.: RM02357

Basic Information

Catalog No.

RM02357

Category

Cell Lysate

Parental Cell line

HeLa

Genotype

Knockout

Background

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]

Gene Information

Gene Symbol

SMPD1

Species

Human

Gene ID

6609

Swiss Prot

P17405

Synonyms

ASM; ASMASE; NPD

Contact

a	400-999-6126
\bowtie	cn.market@abclonal.com.cn
•	www.abclonal.com.cn

Product Information

Description

SMPD1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.

Allele-1:142bp deletion in exon1

Allele-2:142bp deletion in exon1

Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

Packaging

1 vial parental cell Lysate and 1 vial knockout cell Lysate

Shipping Conditions

Amount

4°C

50μL, 2μg/μL.

Storage

Lysate is stable for 12 months when stored at -20 $^{\circ}$ C. Minimizing freeze-thaw cycles.

Protocol

To be used as WB control. Lysate is supplied in $1\times$ SDS sample buffer (2% SDS, 60 mM Tris-HCl pH 6.8, 10% Glycerol, 0.02% Bromophenol blue, 60 mM beta-mercaptoethanol). Lysate should be boiled for 3 - 5 minutes before loading onto gel.

Sequencing data

WT GGACAAGACGGGAC************AGGTTACATCGCAT
Mut GGACAAGACGGGAC***Deletion***AGGTTACATCGCAT
Allele-1: 142bp deletion in exon1

WT GGACAAGACGGGAC*********AGGTTACATCGCAT
Mut GGACAAGACGGGAC***Deletion***AGGTTACATCGCAT

Allele-2: 142bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and SMPD1 knockout (KO) HeLa cells, using sanger sequencing.