

# Npc1-KO

系統名	C57BL/6Smoc- <i>Npc1</i> <sup>em1Smoc</sup>
SMOC番号	NM-KO-205053
維持形態	Sperm cryopreservation

## 遺伝子の概要

Gene Symbol	Synonyms	spm, lcsd, C85354, nmf164, D18ErtD139e, D18ErtD723e, A430089E03Rik
	NCBI ID	<a href="#">18145</a>
	MGI ID	<a href="#">1097712</a>
	Ensembl ID	<a href="#">ENSMUSG00000024413</a>
	Human Ortholog	NPC1

## 説明

Exon 9 of *Npc1* gene was deleted to generate *Npc1* knockout mice.

\*Literature published using this strain should indicate: *Npc1*-KO mice (Cat. NO. NM-KO-205053) were purchased from Shanghai Model Organisms Center, Inc..

## 病気の予測

Niemann-Pick disease	表現型	<a href="#">MGI:4359169</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with <i>Nr1H2</i> -KO(NM-KO-2110634) mice.
	参考文献	Repa JJ, Li H, Frank-Cannon TC, Valasek MA, Turley SD, Tansey MG, Dietschy JM, Liver X receptor activation enhances cholesterol loss from the brain, decreases neuroinflammation, and increases survival of the NPC1 mouse. <i>J Neurosci.</i> 2007 Dec 26;27(52):14470-80

<b>Niemann-Pick disease</b>	<b>表現型</b>	<a href="#">MGI:5305067</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with App-KO(NM-KO-190444) mice.
	<b>参考文献</b>	Nunes A, Pressey SN, Cooper JD, Soriano S, Loss of amyloid precursor protein in a mouse model of Niemann-Pick type C disease exacerbates its phenotype and disrupts tau homeostasis. <i>Neurobiol Dis.</i> 2011 Jun;42(3):349-59
<b>Niemann-Pick disease</b>	<b>表現型</b>	<a href="#">MGI:2386740</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Abcb1a-KO(NM-KO-200904) mice.
	<b>参考文献</b>	Erickson RP, Kiela M, Devine PJ, Hoyer PB, Heidenreich RA, mdr1a deficiency corrects sterility in Niemann-Pick C1 protein deficient female mice. <i>Mol Reprod Dev.</i> 2002 Jun;62(2):167-73
<b>Niemann-Pick disease</b>	<b>表現型</b>	<a href="#">MGI:4436744</a>
	<b>参考文献</b>	Elrick MJ, Pacheco CD, Yu T, Dadgar N, Shakkottai VG, Ware C, Paulson HL, Lieberman AP, Conditional Niemann-Pick C mice demonstrate cell autonomous Purkinje cell neurodegeneration. <i>Hum Mol Genet.</i> 2010 Mar 1;19(5):837-47
<b>Niemann-Pick Disease Type C1</b>	<b>表現型</b>	<a href="#">MGI:6359481</a>
	<b>参考文献</b>	Gomez-Grau M, Albaiges J, Casas J, Auladell C, Dierssen M, Vilageliu L, Grinberg D, New murine Niemann-Pick type C models bearing a pseudoexon-generating mutation recapitulate the main neurobehavioural and molecular features of the disease. <i>Sci Rep.</i> 2017 Feb 7;7:41931

## 表現型データ

No data