

# Lep-KO

系統名	B6;129S-Lep <sup>tm1Smoc</sup>
SMOC番号	NM-KO-00034
維持形態	Repository Live

## 遺伝子の概要

<b>Gene Symbol</b> <b>Lep</b>	<b>Synonyms</b>	ob; obese
	<b>NCBI ID</b>	<a href="#">16846</a>
	<b>MGI ID</b>	<a href="#">104663</a>
	<b>Ensembl ID</b>	<a href="#">ENSMUSG00000059201</a>
	<b>Human Ortholog</b>	LEP

## 説明

Exon 2 was replaced by Neo cassette.

**応用分野:** Insulin resistance, obesity, and type 2 diabetes etc.

\*Literature published using this strain should indicate: Lep-KO mice (Cat. NO. NM-KO-00034) were purchased from Shanghai Model Organisms Center, Inc..

## 病気の予測

<b>Abdominal Obesity-Metabolic Syndrome</b>	<b>表現型</b>	<a href="#">MGI:4429407</a>
	<b>参考文献</b>	Xu A, Liu J, Liu P, Jia M, Wang H, Tao L, Mitochondrial translocation of Nur77 induced by ROS contributed to cardiomyocyte apoptosis in metabolic syndrome. Biochem Biophys Res Commun. 2014 Apr 18;446(4):1184-9
<b>Abdominal Obesity-Metabolic Syndrome 1</b>	<b>表現型</b>	<a href="#">MGI:2654709</a>
	<b>参考文献</b>	Ikels K, Kuschel S, Fischer J, Kaisers W, Eberhard D, Ruther U, FTO is a relevant factor for the development of the metabolic syndrome in mice. PLoS One. 2014;9(8):e105349

<b>Non-Alcoholic Fatty Liver Disease</b>	<b>表現型</b>	<a href="#">MGI:5807153</a>
	<b>参考文献</b>	Trak-Smayra V, Paradis V, Massart J, Nasser S, Jebara V, Fromenty B, Pathology of the liver in obese and diabetic ob/ob and db/db mice fed a standard or high-calorie diet. <i>Int J Exp Pathol.</i> 2011 Dec;92(6):413-21
<b>Obesity</b>	<b>表現型</b>	<a href="#">MGI:3623749</a>
	<b>参考文献</b>	Barouch LA, Berkowitz DE, Harrison RW, O'Donnell CP, Hare JM, Disruption of leptin signaling contributes to cardiac hypertrophy independently of body weight in mice. <i>Circulation.</i> 2003 Aug 12;108(6):754-9
<b>Type 2 Diabetes Mellitus</b>	<b>表現型</b>	<a href="#">MGI:5428893</a>
	<b>参考文献</b>	Clee SM, Nadler ST, Attie AD, Genetic and genomic studies of the BTBR ob/ob mouse model of type 2 diabetes. <i>Am J Ther.</i> 2005 Nov-Dec;12(6):491-8

## 表現型データ

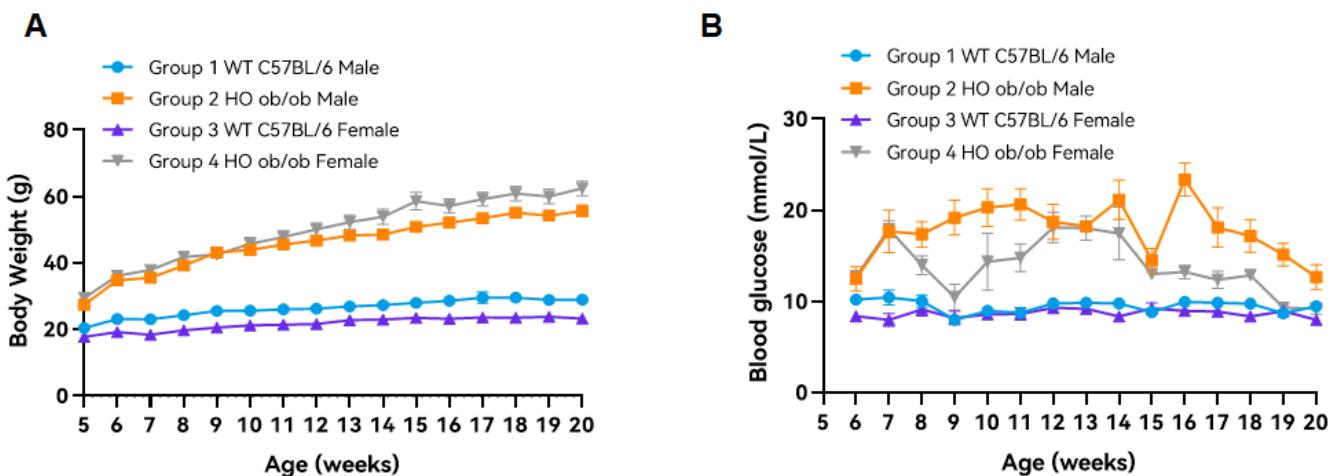


Fig1. Body weight and Blood glucose monitoring of ob/ob mice.

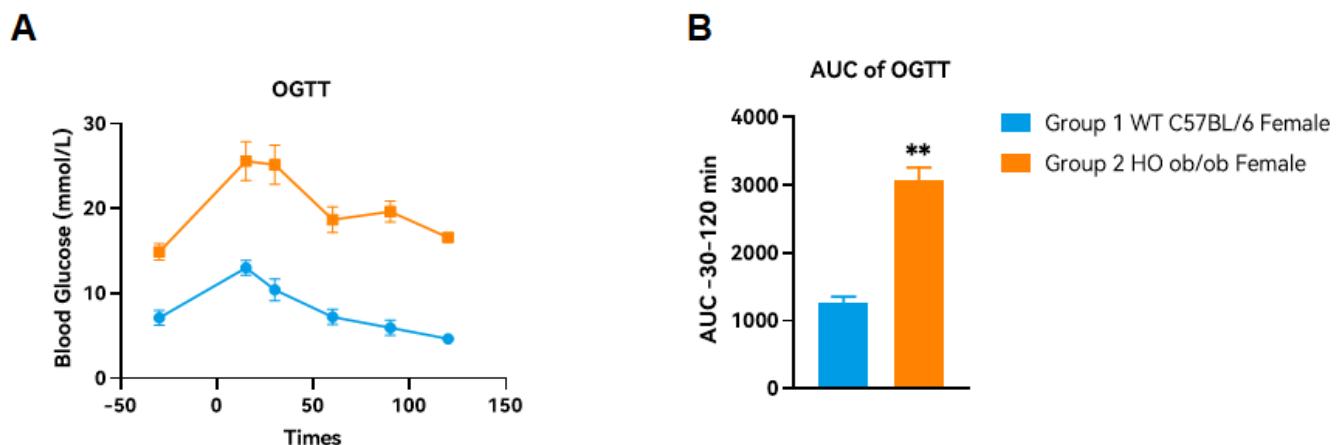


Fig2. Evaluation of Oral glucose tolerance test (OGTT) of female ob/ob mice.

Abbr. HO, homozygous; WT, wild type. \*\*, P ≤ 0.01.

Note. The tested ob/ob and C57BL/6 mice were 6 weeks old.

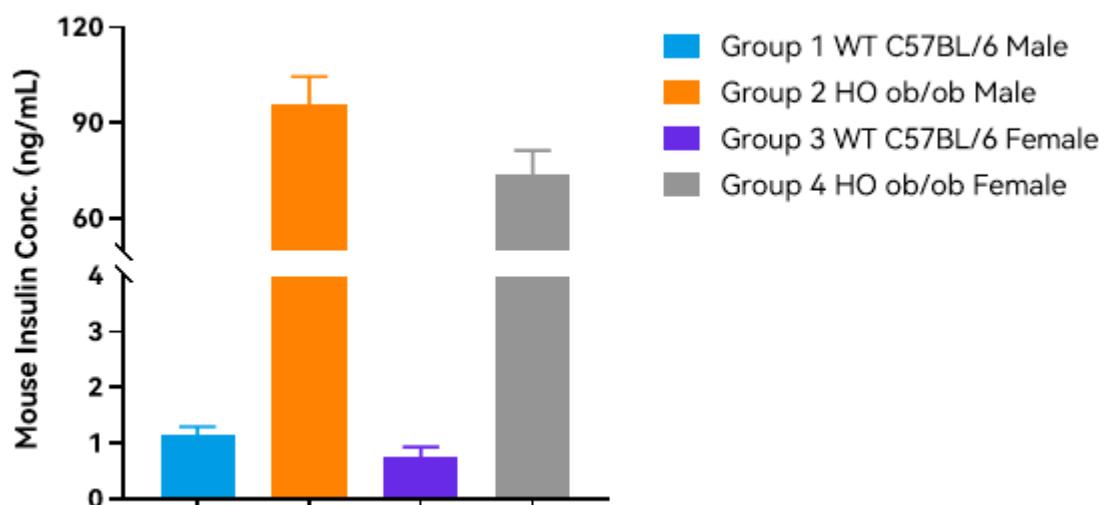


Fig3. Detection of insulin levels in serum by ELISA.

Abbr. Hom, homozygous; HE, heterozygous; WT, wild type.

Note. The tested ob/ob and C57BL/6 mice were 20 weeks old.

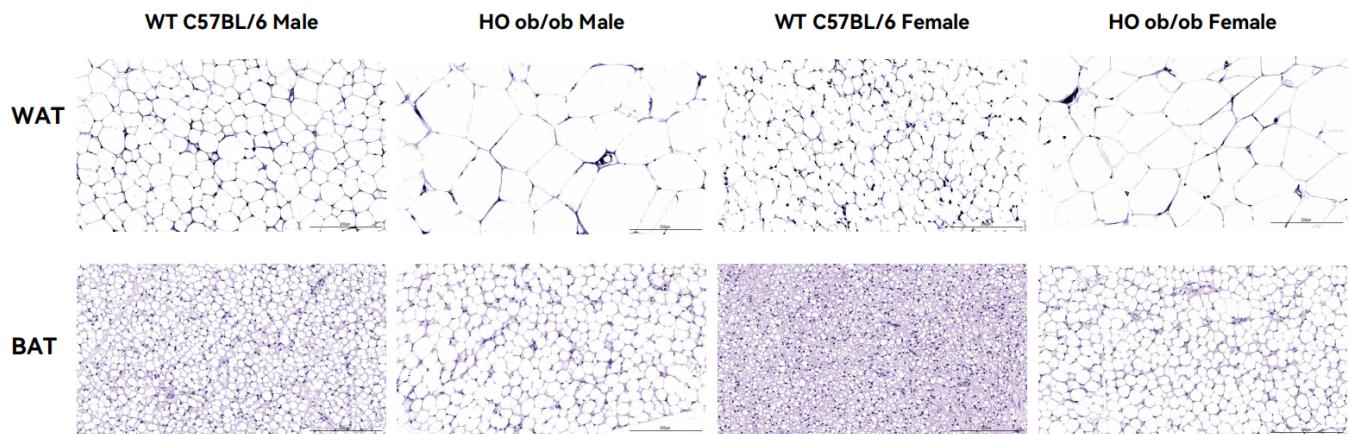


Fig4. H&E staining of the subcutaneous adipose tissue (SAT) in ob/ob mice.

Abbr. HO, homozygous; WT, wild type; WAT, white adipose tissue; BAT, brown adipose tissue.

Note. The tested ob/ob and C57BL/6 mice were 20 weeks old. Scale bar, 200  $\mu$ m; magnification,  $\times$  20.

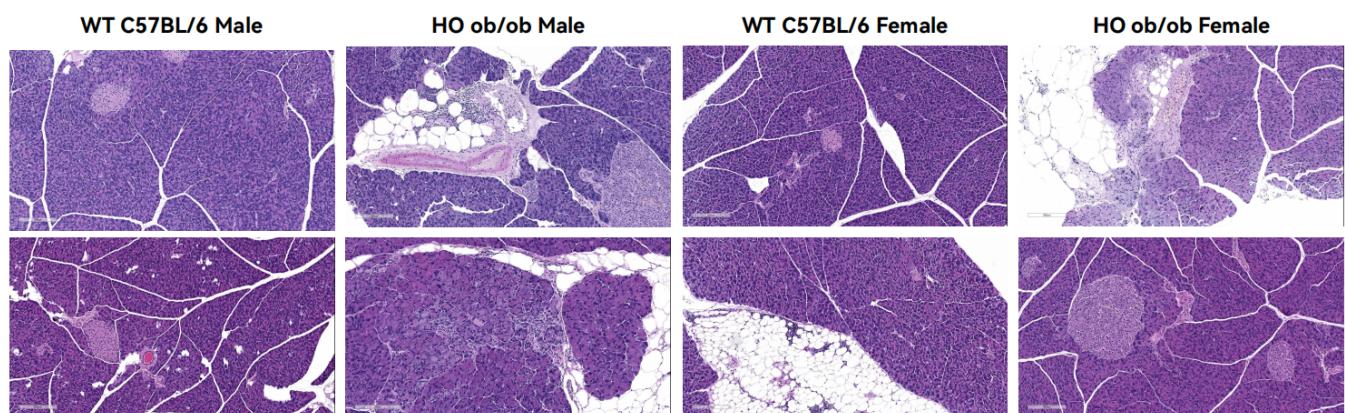


Fig5. Representative pictures of pancreas in ob/ob mice.

These results suggested the inflammatory cell infiltration of the pancreas in ob/ob mice.

Abbr. HO, homozygous; WT, wild type.

Note. The tested ob/ob and C57BL/6 mice were 20 weeks old. Scale bar, 200  $\mu$ m; magnification,  $\times$  10.