

# Slc7a5-Flox

系統名	C57BL/6Smoc- <i>Slc7a5</i> <sup>em1(flox)Smoc</sup>
SMOC番号	NM-CKO-232095
維持形態	Developing

## 遺伝子の概要

Gene Symbol <b>Slc7a5</b>	<b>Synonyms</b>	TA1; LAT1; 4F2LC; D0H16S474E
	<b>NCBI ID</b>	<a href="#">20539</a>
	<b>MGI ID</b>	<a href="#">1298205</a>
	<b>Ensembl ID</b>	<a href="#">ENSMUSG00000040010</a>
	<b>Human Ortholog</b>	SLC7A5

## 説明

These strains carry loxP sites flanking exon 1 of *Slc7a5* gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of *Slc7a5* gene.

\*Literature published using this strain should indicate: *Slc7a5*-Flox mice (Cat. NO. NM-CKO-232095) were purchased from Shanghai Model Organisms Center, Inc..

## 病気の予測

<b>Autism Spectrum Disorder</b>	<b>表現型</b>	<a href="#">MGI:5829465</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Tek-cre mice.
	<b>参考文献</b>	Tarlungeanu DC, Deliu E, Dotter CP, Kara M, Janiesch PC, Scalise M, Galluccio M, Tesulov M, Morelli E, Sonmez FM, Bilguvar K, Ohgaki R, Kanai Y, Johansen A, Esharif S, Ben-Omran T, Topcu M, Schlessinger A, Indiveri C, Duncan KE, Caglayan AO, Gunel M, Gleeson JG, Novarino G, Impaired Amino Acid Transport at the Blood Brain Barrier Is a Cause of Autism Spectrum Disorder. <i>Cell</i> . 2016 Dec 01;167(6):1481-1494.e18

## 表現型データ

No data

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