

STOCK *Sst^{tm2.1(cre)Zjh} /J*

Stock No: **013044** | *Sst*-IRES-Cre (SOM-IRES-Cre)

 Targeted Mutation

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While *Sst*-IRES-Cre was designed as a 3' knock-in allele, additional characterization indicates it has significantly diminished endogenous *Sst* expression - see details below. As such, researchers should consider using heterozygous AVP-IRES2-Cre-D mice and wildtype littermate controls in all their studies.

Of note, the same *Sst*-IRES-Cre knock-in allele is also available as C57BL/6N-congenic (Stock No. [018973](#)) and C57BL/6J-congenic (Stock No. [028864](#)).

Donating Investigator

Z. Josh Huang, Cold Spring Harbor Laboratory

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GENETIC OVERVIEW

Genetic Background

Generation

[F?+F22](#)

(2019-12-31 00:00:00)

Sst^{tm2.1(cre)Zjh}

Allele Type

Targeted (Recombinase-expressing, Knockdown)

Gene Symbol

Sst

Gene Name

somatostatin

VIEW GENETICS

RESEARCH APPLICATIONS

Research Tools

Neurobiology Research

VIEW ALL RESEARCH APPLICATIONS

BASE PRICE

Starting at:

\$305.90 Domestic price for female 5-week

V I E W P R I C E L I S T

Details

Detailed Description

The Sst-IRES-Cre knock-in allele (or SOM-IRES-Cre) has an internal ribosome entry site and Cre recombinase in the 3' UTR of the somatostatin locus (*Sst*). As such, the endogenous *Sst* promoter/enhancer elements direct *cre* expression to somatostatin-expressing neurons. While Sst-IRES-Cre was designed as a 3' knock-in allele, additional characterization indicates it has significantly diminished endogenous *Sst* expression - see details below. As such, researchers should consider using heterozygous AVP-IRES2-Cre-D mice and wildtype littermate controls in all their studies.

When Sst-IRES-Cre mice are bred with mice containing *loxP*-flanked sequences, Cre-mediated recombination will result in deletion of the floxed sequences in the *Sst*-expressing cells in the offspring.

In 2010, the donating investigator reported Cre recombinase activity is specific and efficient; largely recapitulating the endogenous somatostatin expression pattern with efficient recombination. They reported Cre recombinase activity is observed in somatostatin positive neurons (including dendritic inhibitory interneurons such as Martinotti cells and Oriens-Lacunosum-Moleculare (O-LM) cells). The donating investigator did not examine *cre* expression in tissues other than brain. *Sst* expression from the Sst-IRES-Cre allele was not evaluated. They also reported that homozygous mice were viable, fertile and normal in size, with no gross physical abnormalities or behavioral abnormalities (see more recent information below).

Although Sst-IRES-Cre was designed as a 3' knock-in allele, additional characterization indicates it has significantly diminished endogenous *Sst* expression. Specifically, in 2016, unpublished research using Stock No. 013044 reported the Sst-IRES-Cre allele had diminished *Sst* RNA expression, and homozygous mice had abnormal locomotor activity (reduced in males during circadian cycle active phase, increased in females by the end of circadian cycle active phase). Heterozygous mice had partial recovery of *Sst* expression and normal behavioral responses. Furthermore, the findings of another group confirmed Sst-IRES-Cre imparts an allele dosage-dependent knock-down of endogenous *Sst* expression [Viollet *et al.* 2017 Front Endocrinol (Lausanne) 8:131 (PMID:28674519)]. Researchers should consider using heterozygous Sst-IRES-Cre mice and wildtype littermate controls in their studies.

For characterization information of the Sst-IRES-Cre knock-in allele, see images at the Allen Institute for Brain Science website ([Sst-IRES-Cre images](#)).

If the recombinase activity pattern of this allele is further characterized by the Genetic Resource Science group at The Jackson Laboratory, such findings will be reported on the Mouse Genome Informatics (MGI) Allele Detail entry ([Sst^{tm2.1\(cre\)Zjh}](#)). This same information would also be found searching the [MGI Recombinase Activity](#) database.

Development

Expression Data

Control Suggestions

Selected References

– Genetics

+ [Sst^{tm2.1\(cre\)}Zjh](#)

– Disease/Phenotype

+ [Disease Terms](#)

+ [Research Areas By Phenotype](#)

+ [Mammalian Phenotype Terms by Genotype](#)

+ [References](#)

– Technical Support

C O N T A C T T E C H N I C A L S U P P O R T

Genotyping Protocols

Separated PCR:[Sst Alternate1](#)

[Genotyping resources and troubleshooting](#)

Dietary Information

LabDiet® 5K52 formulation (6% fat)

Breeding Considerations

Homozygous mice are viable and fertile. When maintaining a live colony at The Jackson Laboratory Repository, homozygous mice may be bred together.

Researchers should consider using heterozygous Sst-IRES-Cre mice and wildtype littermate controls in their studies - please see strain description for more details.

[Additional Breeding and Husbandry Support](#)

Mating System

Homozygote x Homozygote

Citation

When using the Sst-IRES-Cre (SOM-IRES-Cre) mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #013044 in your Materials and Methods section.

Animal Health Reports

[Facility Barrier Level Descriptions](#)

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	Male	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
6 weeks	Female	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
	Male	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
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	Male	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
8 weeks	Female	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
	Male	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
9 weeks	Female	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90
	Male	Homozygous for Sst ^{tm2.1(cre)Zjh}	\$305.90

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