

Srrm2 Cas9-KO Strategy

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Overview

Target Gene Name

- Srrm2

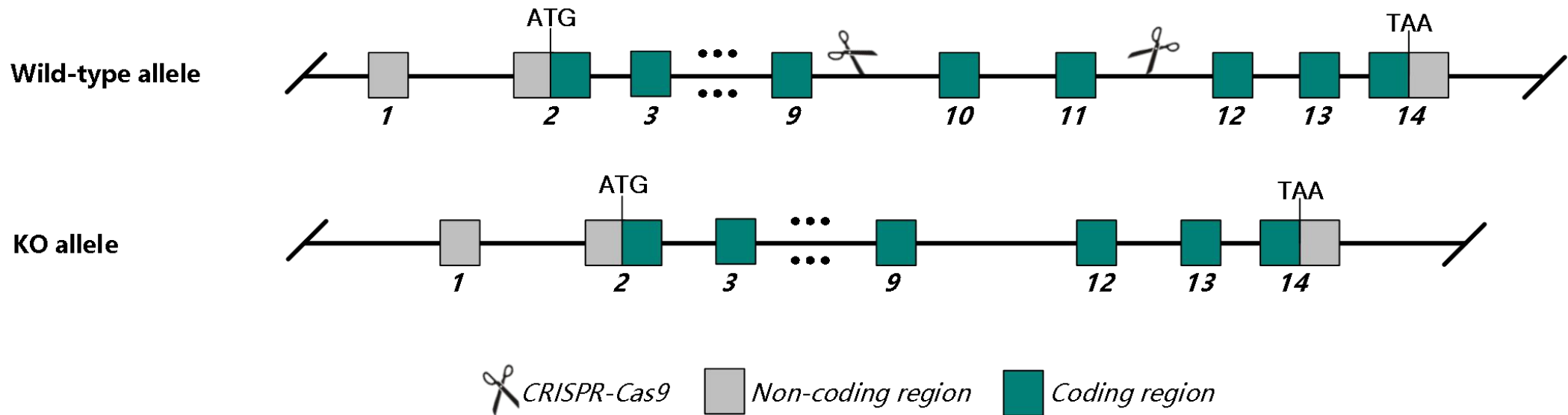
Project Type

- Cas9-KO

Genetic Background

- C57BL/6JGpt

Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the *Srrm2* gene.

Technical Information

- The *Srrm2* gene has 12 transcripts. According to the structure of *Srrm2* gene, exon 10-exon 11 of *Srrm2*-201 (ENSMUST00000088621.11) transcript is recommended as the knockout region. The region contains 6857 bp of coding sequences. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Srrm2* gene. The brief process is as follows: gRNAs were transcribed in vitro. Cas9 and gRNAs were microinjected into the fertilized eggs of C57BL/6JGpt mice. Fertilized eggs were transplanted to obtain positive F0 mice which were confirmed by PCR and on-target amplicon sequencing. A stable F1-generation mouse strain was obtained by mating positive F0-generation mice with C57BL/6JGpt mice and confirmation of the desired mutant allele was carried out by PCR and on-target amplicon sequencing.

Gene Information

Srrm2 serine/arginine repetitive matrix 2 [*Mus musculus* (house mouse)]

[Download Datasets](#)

Gene ID: 75956, updated on 30-Aug-2022

Summary

| | |
|---------------------------|---|
| Official Symbol | Srrm2 provided by MGI |
| Official Full Name | serine/arginine repetitive matrix 2 provided by MGI |
| Primary source | MGI:MGI:1923206 |
| See related | Ensembl:ENSMUSG00000039218 AllianceGenome:MGI:1923206 |
| Gene type | protein coding |
| RefSeq status | VALIDATED |
| Organism | Mus musculus |
| Lineage | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| Also known as | SRm300; mKIAA0324; 5033413A03Rik |
| Summary | Predicted to enable C2H2 zinc finger domain binding activity; mRNA binding activity; and protein N-terminus binding activity. Predicted to be involved in mRNA splicing, via spliceosome. Predicted to act upstream of or within RNA splicing and mRNA processing. Predicted to be located in Cajal body and nuclear speck. Predicted to be part of U2-type catalytic step 2 spliceosome and U2-type precatalytic spliceosome. Orthologous to human SRRM2 (serine/arginine repetitive matrix 2). [provided by Alliance of Genome Resources, Apr 2022] |
| Expression | Ubiquitous expression in adrenal adult (RPKM 122.5), ovary adult (RPKM 102.7) and 28 other tissues See more |
| Orthologs | human all |
| NEW | Try the new Gene table Try the new Transcript table |

Genomic context

Location: 17; 17 A3.3

See Srrm2 in [Genome Data Viewer](#)

Exon count: 20

Source: <https://www.ncbi.nlm.nih.gov/gene/75956>

Transcript Information

The gene has 12 transcripts, all transcripts are shown below:

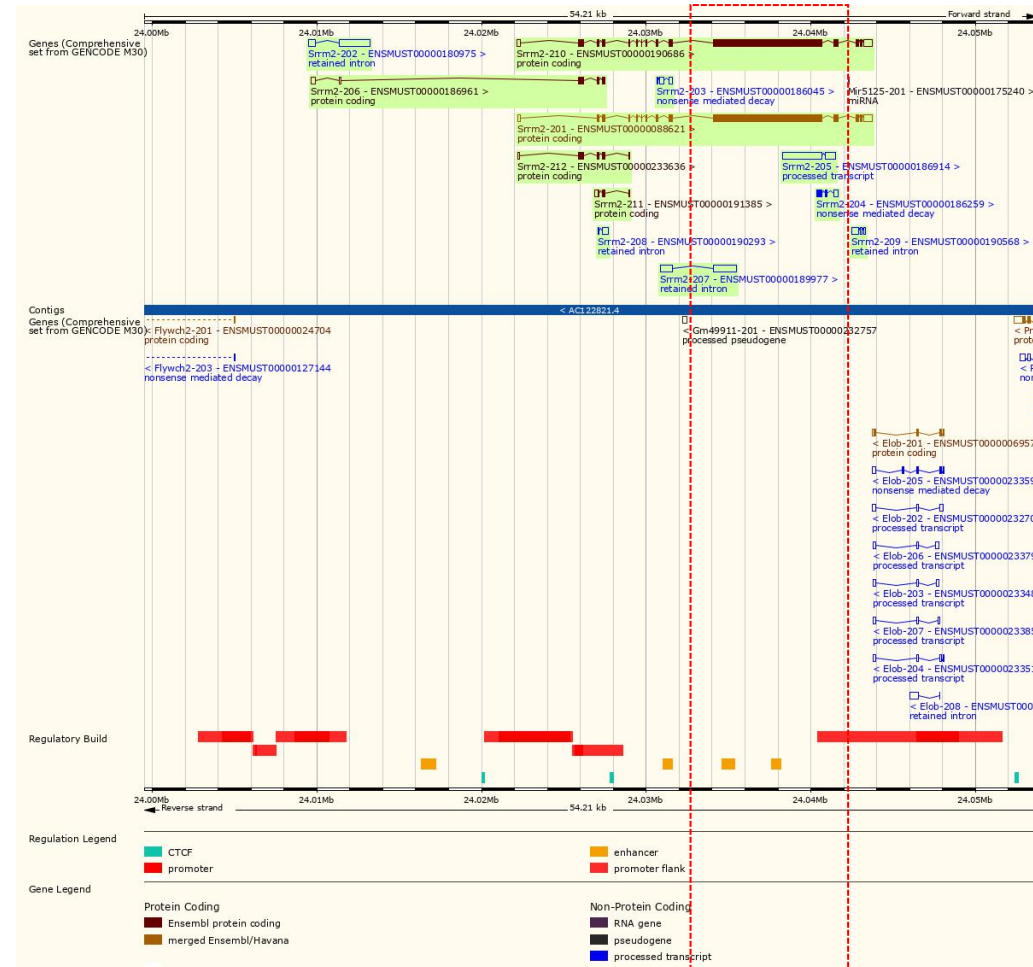
| Show/hide columns (1 hidden) | | | | | | | Filter |
|---------------------------------------|-----------|------|------------------------|-------------------------|---------------------------|----------------------------|---|
| Transcript ID | Name | bp | Protein | Biotype | CCDS | UniProt Match | Flags |
| ENSMUST00000233636.2 | Srrm2-212 | 678 | 191aa | Protein coding | | A0A3B2W3Z4 | CDS 3' incomplete |
| ENSMUST00000191385.3 | Srrm2-211 | 459 | 88aa | Protein coding | | A0A087WPS9 | TSL:2 CDS 3' incomplete |
| ENSMUST00000190686.7 | Srrm2-210 | 8864 | 2703aa | Protein coding | | Q8BTI8-1 | Ensembl Canonical Gencode basic APPRIS P1 TSL:5 |
| ENSMUST00000190568.2 | Srrm2-209 | 597 | No protein | Retained intron | | - | TSL:2 |
| ENSMUST00000190293.2 | Srrm2-208 | 431 | No protein | Retained intron | | - | TSL:3 |
| ENSMUST00000189977.2 | Srrm2-207 | 2092 | No protein | Retained intron | | - | TSL:1 |
| ENSMUST00000186961.7 | Srrm2-206 | 950 | 171aa | Protein coding | | A0A087WRX8 | TSL:5 CDS 3' incomplete |
| ENSMUST00000186914.2 | Srrm2-205 | 2991 | No protein | Processed transcript | | - | TSL:1 |
| ENSMUST00000186259.2 | Srrm2-204 | 647 | 105aa | Nonsense mediated decay | | A0A087WR98 | TSL:3 CDS 5' incomplete |
| ENSMUST00000186045.2 | Srrm2-203 | 494 | 34aa | Nonsense mediated decay | | A0A087WPU0 | TSL:3 CDS 5' incomplete |
| ENSMUST00000180975.3 | Srrm2-202 | 2266 | No protein | Retained intron | | - | TSL:1 |
| ENSMUST00000088621.11 | Srrm2-201 | 8564 | 2607aa | Protein coding | CCDS37475 | Q8BTI8-2 | Gencode basic TSL:1 |

The strategy is based on the design of *Srrm2-201* transcript, the transcription is shown below:



http://asia.ensembl.org/Mus_musculus/Transcript/Summary?db=core;g=ENSMUSG00000039218;r=17:24009506-24043715;t=ENSMUST00000088621

Genomic Information



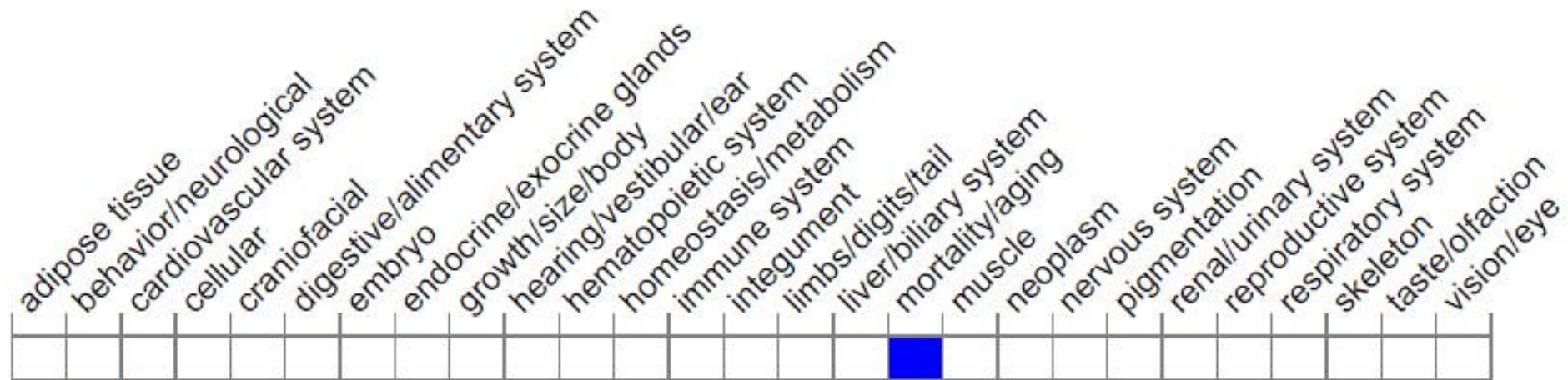
[http://asia.ensembl.org/Mus_musculus/Gene/Summary?db=core;g=E
NSMUSG00000039218;r=17:24009506-
24043715;t=ENSMUST000000088621](http://asia.ensembl.org/Mus_musculus/Gene/Summary?db=core;g=ENSMUSG00000039218;r=17:24009506-24043715;t=ENSMUST000000088621)

Protein Information



http://asia.ensembl.org/Mus_musculus/Transcript/ProteinSummary?db=core;g=ENSMUSG00000039218;r=17:24009506-24043715;t=ENSMUST000000088621

Mouse Phenotype Information (MGI)



Source: <http://www.informatics.jax.org/marker/MGI:1923206>

Important Information

- According to the available information (MGI), there is a certain lethality rate before weaning in mice homozygous for knock-out alleles.
<http://www.informatics.jax.org/diseasePortal/genoCluster/view/33639>
- The sequences encoding transcripts *Srrm2*-203, *Srrm2*-204, *Srrm2*-206, *Srrm2*-211, and *Srrm2*-212 are incomplete and the effect of knocking out the target region on them is unknown.
- The knockdown region designed in this strategy is about 1.6 kp away from the 3' end of the *Elob* gene, and knockdown of the target gene may affect the regulation of the 3' end of the *Elob* gene.
- The effect of this strategy on the uncoded transcript *Gm49911*-201、*Mir5125*-201 is unknown.
- *Srrm2* is located on Chr 17. If the knockout mice are crossed with other mouse strains to obtain double homozygous mutant offspring, please avoid the situation that the second gene is on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risks of the mutation on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.