

# Srsf10 Cas9-KO Strategy

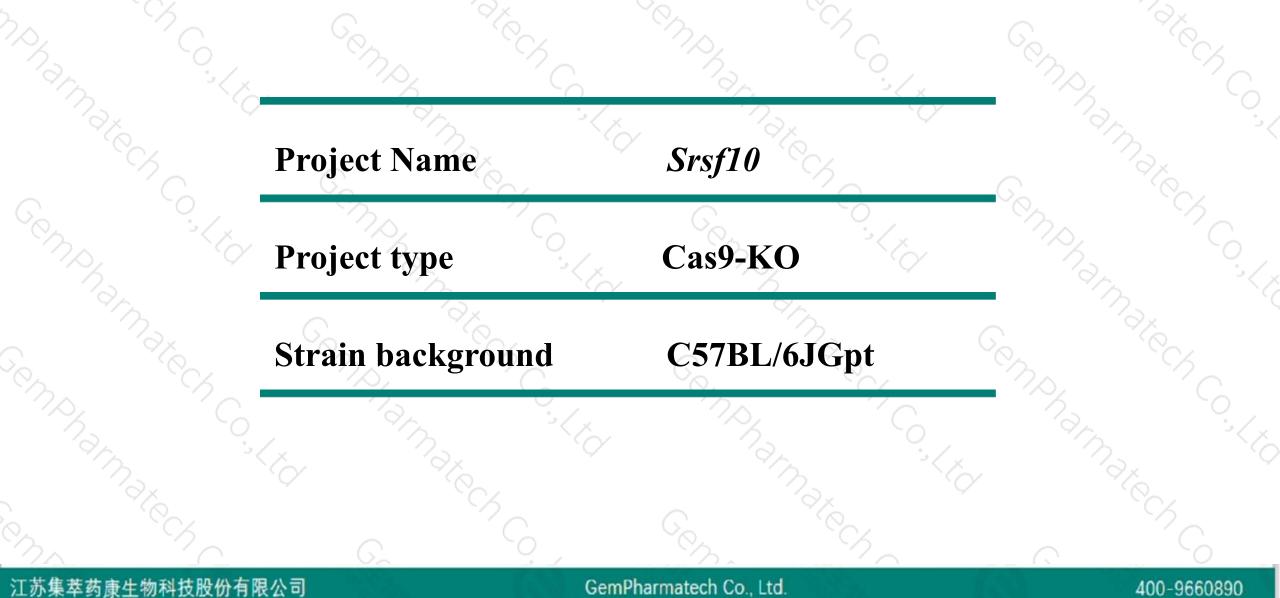
Designer: Design Date:

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Huan Wang 2019-8-8

### **Project Overview**

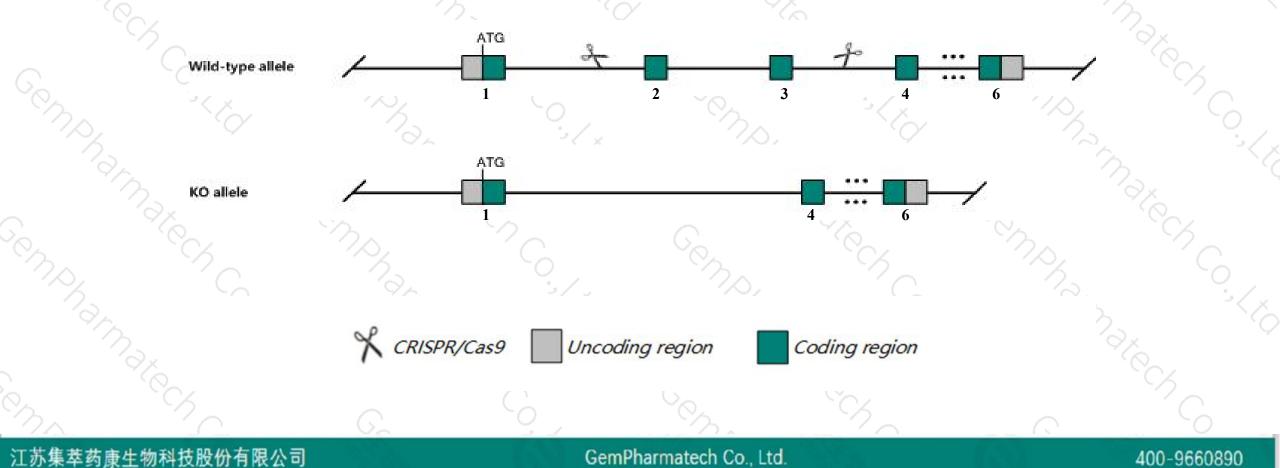




# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Srsf10 gene. The schematic diagram is as follows:





- The Srsf10 gene has 10 transcripts. According to the structure of Srsf10 gene, exon2-exon3 of Srsf10-205 (ENSMUST00000126641.1) transcript is recommended as the knockout region. The region contains 209bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Srsf10 gene. The brief process is as follows: CRISPR/Cas9 system

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- According to the existing MGI data, Mice homozygous for a null allele exhibit fetal and neonatal lethality associated with edema and cardiac defects.
- The Srsf10 gene is located on the Chr4. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Notice

# **Gene information (NCBI)**



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### Srsf10 serine/arginine-rich splicing factor 10 [Mus musculus (house mouse)]

Gene ID: 14105, updated on 3-Feb-2019

#### Summary

Official Symbol	Srsf10 provided by MGI
<b>Official Full Name</b>	serine/arginine-rich splicing factor 10 provided by MGI
Primary source	MGI:MGI:1333805
See related	Ensembl:ENSMUSG0000028676
Gene type	protein coding
<b>RefSeq status</b>	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	FUSIP2, Fusip1, NSSR1, NSSR2, Nssr, SRrp40, Sfrs13a, Srsf13a, TASR, TASR1, TASR2
Expression	Ubiquitous expression in CNS E11.5 (RPKM 24.5), limb E14.5 (RPKM 19.6) and 28 other tissues See more
Orthologs	human all

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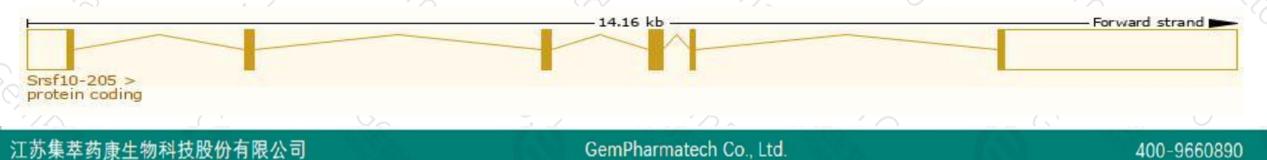
## **Transcript information (Ensembl)**



### The gene has 10 transcripts, all transcripts are shown below:

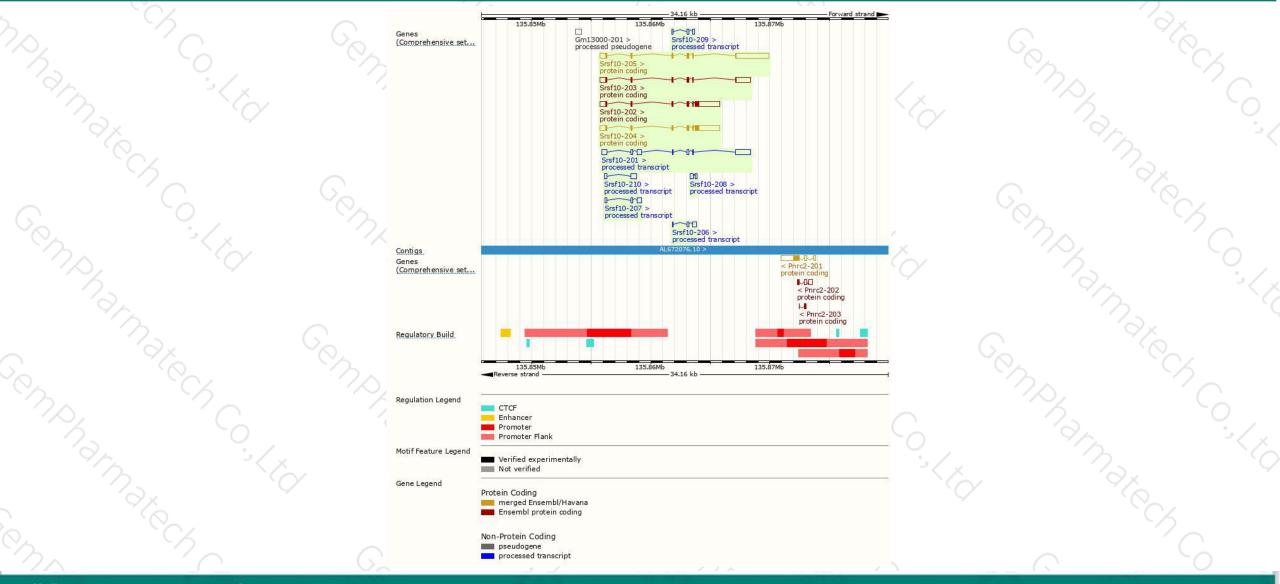
				Jun		> ^ _	the second s	at for a		
	Name 🍦	Transcript ID 🔹	bp 🖕	Protein 🖕	Biotype 🍦	CCDS 🖕	UniProt 🖕	Flags 🖕		
	Srsf10-210	ENSMUST00000154447.1	677	No protein	IncRNA	-	( <del>, ,</del> )	TSL:1		
	Srsf10-209	ENSMUST00000149878.7	394	No protein	IncRNA	-	(1 <b>-</b> )	TSL:3		
	Srsf10-208	ENSMUST00000142002.1	423	No protein	IncRNA	-	0.73	TSL:2		
	Srsf10-207	ENSMUST00000129718.1	717	No protein	IncRNA	3 <del>.</del>	(i <del>n</del> )	TSL:1		
	Srsf10-206	ENSMUST00000129198.1	522	No protein	IncRNA	-	070	TSL:3		
	Srsf10-205	ENSMUST00000126641.1	3754	<u>183aa</u>	Protein coding	<u>CCDS18791</u> &	<u>Q3UA07</u> @ <u>Q9R0U0</u> @	TSL:1 GENCODE basic APPRIS P3		
	Srsf10-204	ENSMUST00000105853.9	3058	<u>262aa</u>	Protein coding	CCDS38922	Q9R0U0团	TSL:1 GENCODE basic APPRIS ALT1		
	Srsf10-203	ENSMUST00000102544.8	2225	<u>182aa</u>	Protein coding	CCDS71494@	Q3TFP0 团	TSL:1 GENCODE basic APPRIS ALT1		
	Srsf10-202	ENSMUST0000097844.8	3055	<u>261aa</u>	Protein coding	<u>CCDS71493</u> &	Q9R0U0 团	TSL:2 GENCODE basic APPRIS ALT1		
	Srsf10-201	ENSMUST0000030438.14	2493	No protein	IncRNA	-	070	TSL:5		

The strategy is based on the design of Srsf10-205 transcript, The transcription is shown below



### **Genomic location distribution**





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### **Protein domain**



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	ENSMUSP00000114 MobiDB lite Low complexity (Seg) hmmpanther	PTHR23					-	. 1			
`` ?/;	Superfamily domains SMART domains	Serine/arginine-rich splicing factor 10. RNA-binding domain superfamily RNA recognition motif domain									
	Pfam domain	RNA	recognition	motif domain	1						
	PROSITE profiles	RNA r	ecognition m	otif domain							
	Gene3D	Nucleotide	binding alph	ia-beta plait d	lomain supe	rfamily					
52	All sequence SNPs/i	Sequence variants (dbSNP and all other sources)									6
	Variant Legend	splice region variant									5
	Scale bar	0	20	40	60	80	100	120	140	160	183
											X
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### Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

According to the existing MGI data, Mice homozygous for a null allele exhibit fetal and neonatal lethality associated with edema and cardiac defects.

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If you have any questions, you are welcome to inquire. Tel: 400-9660890



