

# Insc Cas9-CKO Strategy

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Reviewer: Ruirui Zhang

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## **Project Overview**



**Project Name** 

Insc

**Project type** 

Cas9-CKO

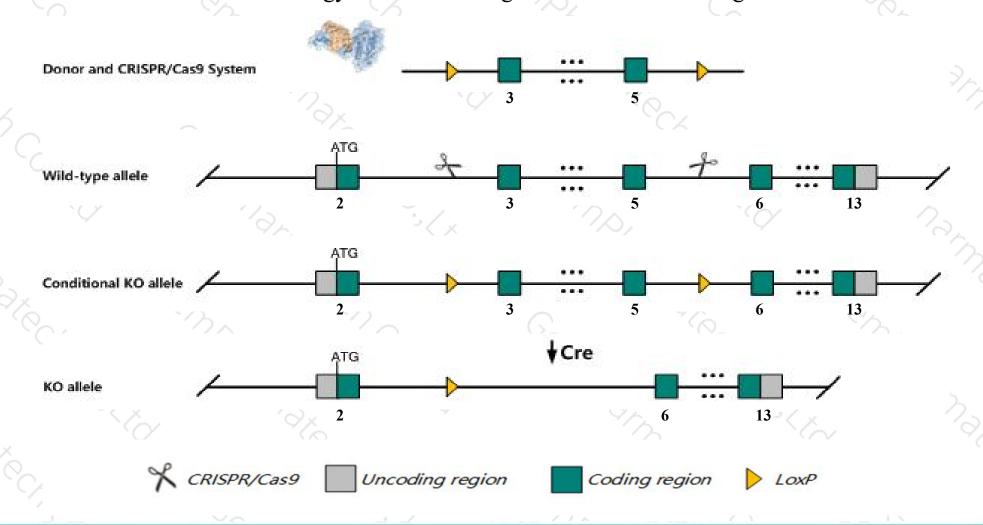
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



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



## Technical routes



- ➤ The *Insc* gene has 9 transcripts. According to the structure of *Insc* gene, exon3-exon5 of *Insc-208*(ENSMUST00000169913.7) transcript is recommended as the knockout region. The region contains 523bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Insc* gene. The brief process is as follows:CRISPR/Cas9 system and Donor 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 sequencing. A stable F1 generation mouse model was obtained by mating positive F0 generation mice with C57BL/6JGpt mice.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

### **Notice**



- ➤ According to the existing MGI data, Homozygous inactivation of this gene leads to abnormal cochlear hair cell morphology.
- The *Insc* gene is located on the Chr7. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

### Gene information (NCBI)



#### Insc INSC spindle orientation adaptor protein [ Mus musculus (house mouse) ]

Gene ID: 233752, updated on 17-Dec-2019

#### Summary

△ ?

Official Symbol Insc provided by MGI

Official Full Name INSC spindle orientation adaptor protein provided by MGI

Primary source MGI:MGI:1917942

See related Ensembl: ENSMUSG00000048782

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 C730020D23; 3830422K02Rik

Expression Broad expression in liver adult (RPKM 5.3), liver E18 (RPKM 3.5) and 21 other tissues See more

Orthologs human all



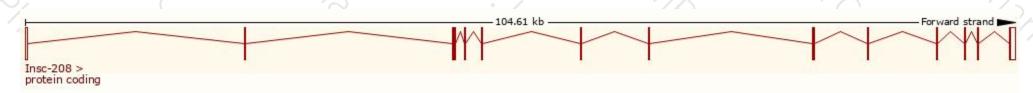
## Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

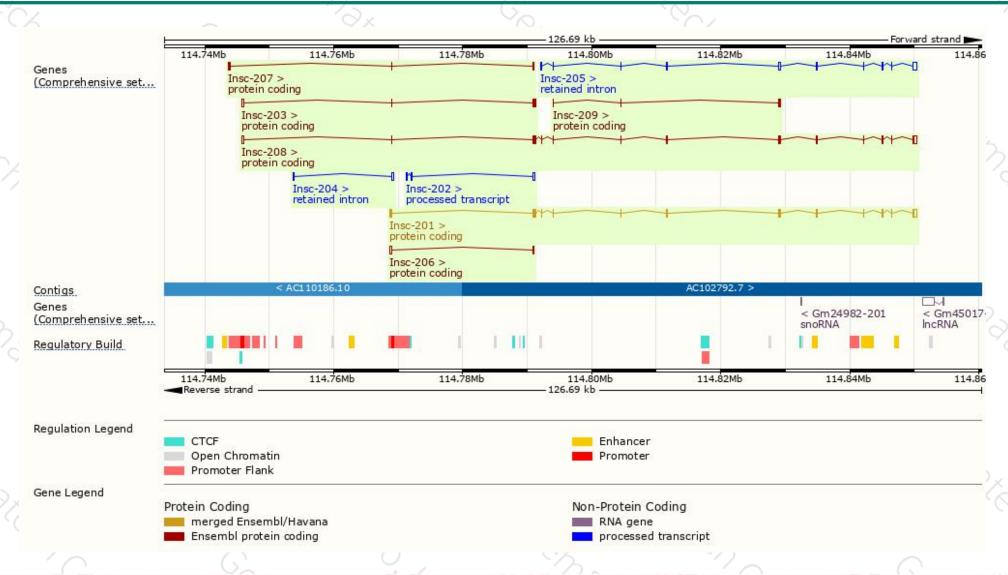
Name 🍦	Transcript ID	bp 🍦	Protein	Biotype	CCDS 🍦	UniProt 🍦	Flags
Insc-208	ENSMUST00000169913.7	2321	<u>532aa</u>	Protein coding	CCDS40095 ₽	<u>G3UW77</u> ₽	TSL:5 GENCODE basic APPRIS P1
Insc-201	ENSMUST00000117543.2	2170	<u>532aa</u>	Protein coding	CCDS40095₽	<u>G3UW77</u> ₽	TSL:5 GENCODE basic APPRIS P1
Insc-203	ENSMUST00000136645.7	713	<u>133aa</u>	Protein coding	2	D3Z267₽	CDS 3' incomplete TSL:2
Insc-207	ENSMUST00000161800.1	451	<u>88aa</u>	Protein coding	2	E0CXR9函	CDS 3' incomplete TSL:5
Insc-209	ENSMUST00000206274.1	405	<u>135aa</u>	Protein coding	<u> </u>	A0A0U1RNN4₽	CDS 5' and 3' incomplete TSL:3
Insc-206	ENSMUST00000151464.1	363	<u>53aa</u>	Protein coding		<u>D3Z5D7</u> ₽	CDS 3' incomplete TSL:1
Insc-202	ENSMUST00000136347.1	439	No protein	Processed transcript	2	2	TSL:1
Insc-205	ENSMUST00000150991.2	1826	No protein	Retained intron	2	2	TSL:1
Insc-204	ENSMUST00000139670.1	600	No protein	Retained intron	28	2	TSL:3

The strategy is based on the design of *Insc-208* transcript, The transcription is shown below



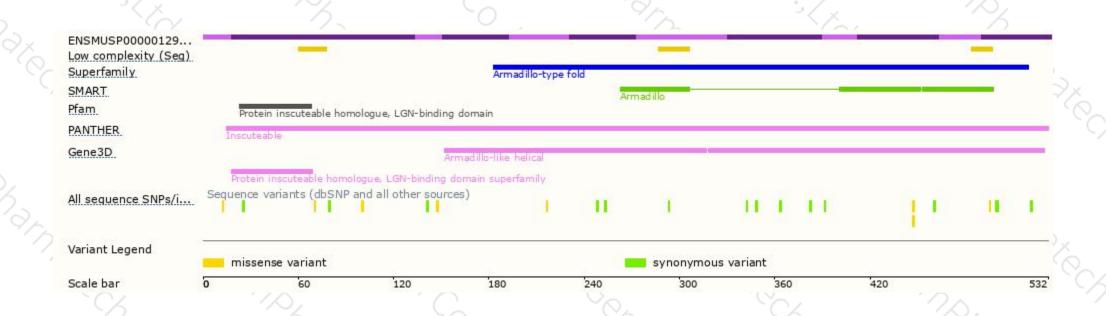
### Genomic location distribution





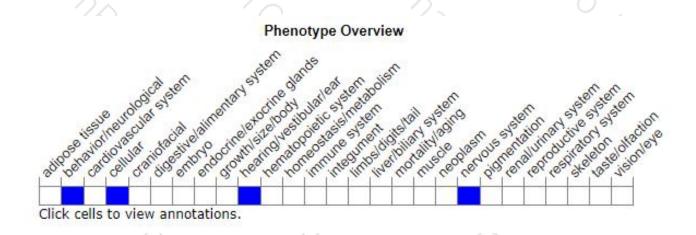
### Protein domain





## 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, Homozygous inactivation of this gene leads to abnormal cochlear hair cell morphology.



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





