

Star Cas9-CKO Strategy

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Project Overview



Project Name Star

Project type

Cas9-CKO

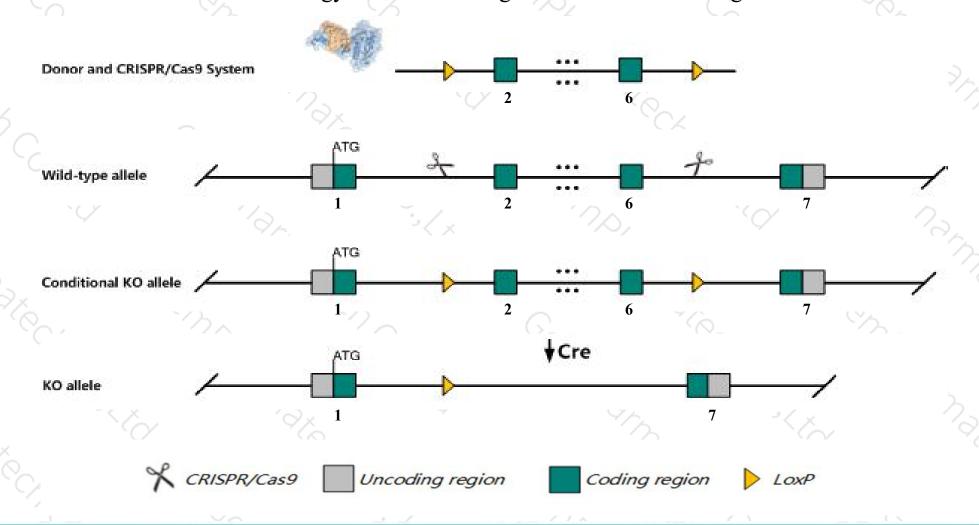
Strain background

C57BL/6JGpt

Conditional Knockout strategy



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



Technical routes



- ➤ The *Star* gene has 2 transcripts. According to the structure of *Star* gene, exon2-exon6 of *Star-201*(ENSMUST00000033979.5) transcript is recommended as the knockout region. The region contains 677bp coding sequence.

 Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Star* 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 null mice fail to thrive and die during the postnatal period due to adrenocortical insufficiency. Mice exhibit male pseudohermaprhoditism and show a progressive accumulation of lipids within steroidogenic cells of the adrenal glands and gonads.
- ➤ The effect on transcript *Star*-202 is unknown.
- > The *Star* gene is located on the Chr8. 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)



Star steroidogenic acute regulatory protein [Mus musculus (house mouse)]

Gene ID: 20845, updated on 12-Aug-2019

Summary

☆ ?

Official Symbol Star provided by MGI

Official Full Name steroidogenic acute regulatory protein provided by MGI

Primary source MGI:MGI:102760

See related Ensembl: ENSMUSG00000031574

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;

Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

Also known as stARD1; AV363654; D8Ertd419e

Expression Restricted expression toward adrenal adult (RPKM 849.8) See more

Orthologs human all

Genomic context



Location: 8 A2; 8 14.17 cM

See Star in Genome Data Viewer

Exon count: 7

Annotation release	Status	Assembly	Chr	Location	
108	current	GRCm38.p6 (GCF_000001635.26)		NC_000074.6 (2580847425815982)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	8	NC_000074.5 (2691898526926454)	

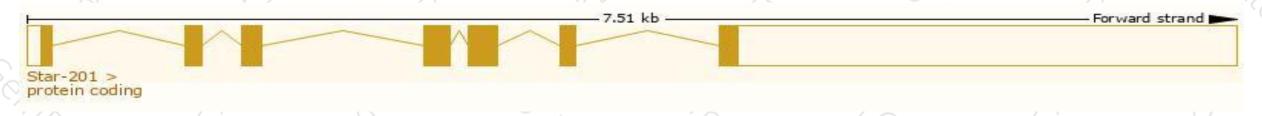
Transcript information (Ensembl)



The gene has 2 transcripts, all transcripts are shown below:

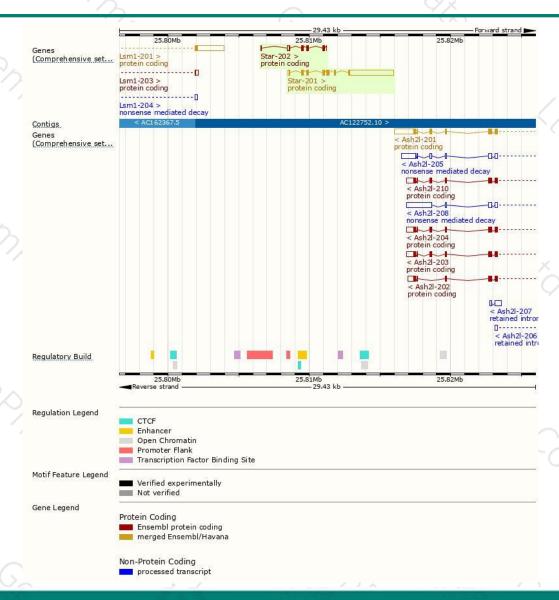
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Star-201	ENSMUST00000033979.5	4046	284aa	Protein coding	CCDS22203	P51557	TSL:1 GENCODE basic APPRIS P1
Star-202	ENSMUST00000210565.1	633	<u>157aa</u>	Protein coding	-	A0A1B0GSV6	CDS 3' incomplete TSL:5

The strategy is based on the design of Star-201 transcript, The transcription is shown below



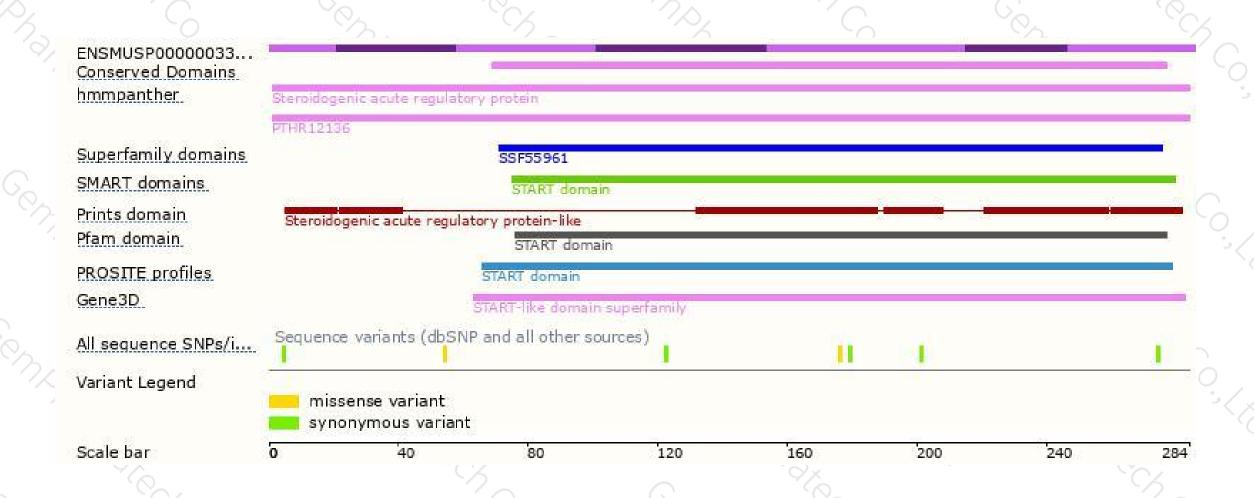
Genomic location distribution





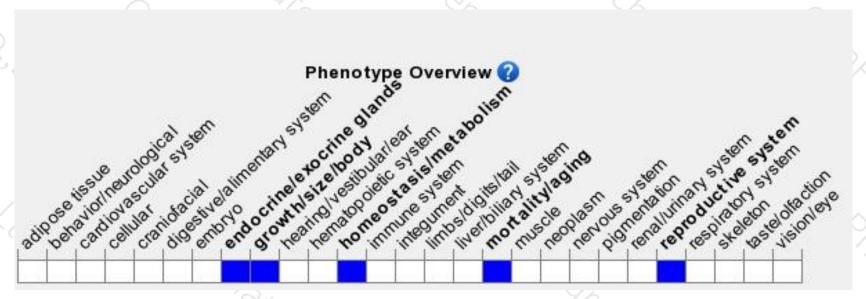
Protein domain





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 null mice fail to thrive and die during the postnatal period due to adrenocortical insufficiency. Mice exhibit male pseudohermaprhoditism and show a progressive accumulation of lipids within steroidogenic cells of the adrenal glands and gonads.



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





