

Chst14 Cas9-CKO Strategy

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Project Overview



Project Name

Chst14

Project type

Cas9-CKO

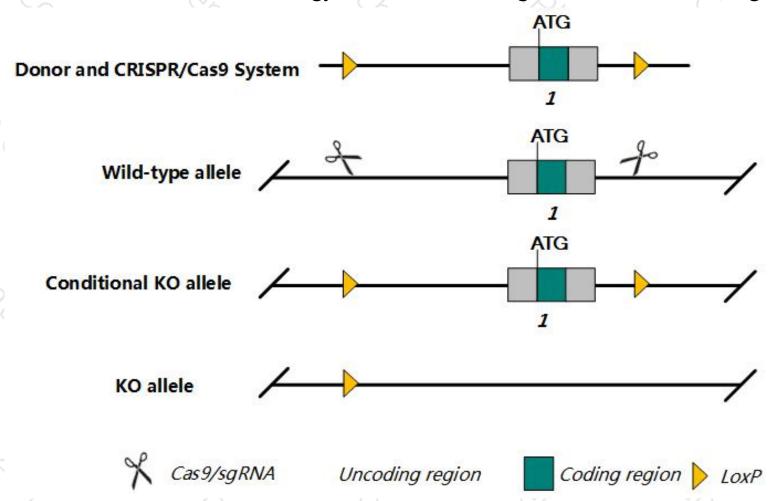
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Chst14 gene. The schematic diagram is as follows:



Technical routes



- The *Chst14* gene has 2 transcripts. According to the structure of *Chst14* gene, exon1 of *Chst14-201*(ENSMUST00000099546.5) transcript is recommended as the knockout region. The region contains all the coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Chst14* gene. The brief process is as follows:gRNA was transcribed in vitro, donor was constructed.Cas9, gRNA 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, Mice homozygous for a knock-out allele exhibit accelerated peripheral nerve regeneration, decreased body weight, premature death, fragile skin and background sensitive abnormal fertility, kinked tail and tooth growth.
- The floxed region is near to the C-terminal of *Bahd1* gene, this strategy may influence the regulatory function of the C-terminal of *Bahd1* gene.
- The *Chst14* gene is located on the Chr2. 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)



Chst14 carbohydrate sulfotransferase 14 [Mus musculus (house mouse)]

Gene ID: 72136, updated on 12-Aug-2019

Summary

☆ ?

Official Symbol Chst14 provided by MGI

Official Full Name carbohydrate sulfotransferase 14 provided by MGI

Primary source MGI:MGI:1919386

See related Ensembl: ENSMUSG00000074916

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 D4st1; D4ST-1; 2600016L03Rik

Orthologs human all

Genomic context



Location: 2; 2 E5

See Chst14 in Genome Data Viewer

Exon count: 1

Annotation release	Status	Assembly	Chr	Location
<u>108</u>	current	GRCm38.p6 (GCF_000001635.26)	2	NC_000068.7 (118926497118928585)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	2	NC_000068.6 (118752233118754321)

Transcript information (Ensembl)



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

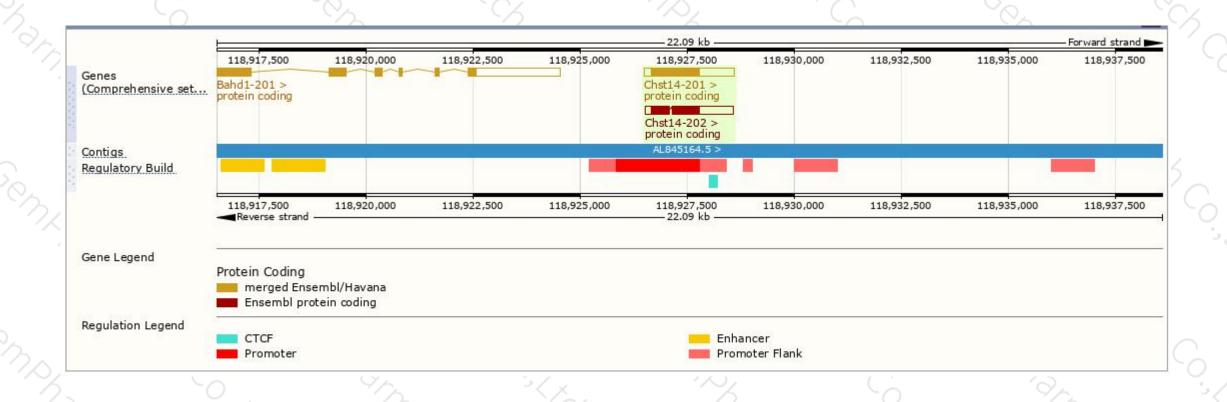
Name	Transcript ID A	bp 👙	Protein 🍦	Biotype	CCDS 🍦	UniProt 🌲	Flags		
Chst14-201	ENSMUST00000099546.5	2090	<u>376aa</u>	Protein coding	CCDS16587 ₽	Q80V53₽	TSL:NA GE	NCODE basic	APPRIS P1
Chst14-202	ENSMUST00000110837.1	1991	<u>351aa</u>	Protein coding		A2AQV2₽	TSL:1 GENCODE basic		basic

The strategy is based on the design of Chst14-201 transcript, The transcription is shown below



Genomic location distribution





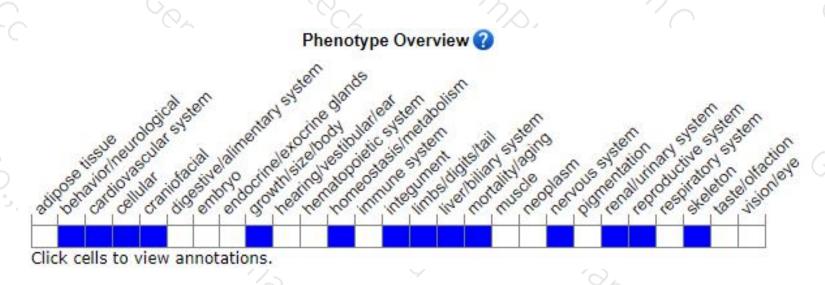
Protein domain





Mouse phenotype description(MGI)





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

Mice homozygous for a knock-out allele exhibit accelerated peripheral nerve regeneration, decreased body weight, premature death, fragile skin and background sensitive abnormal fertility, kinked tail and tooth growth.



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





