

Actc1 Cas9-CKO Strategy

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Project Overview

Project Name

Actc1

Project type

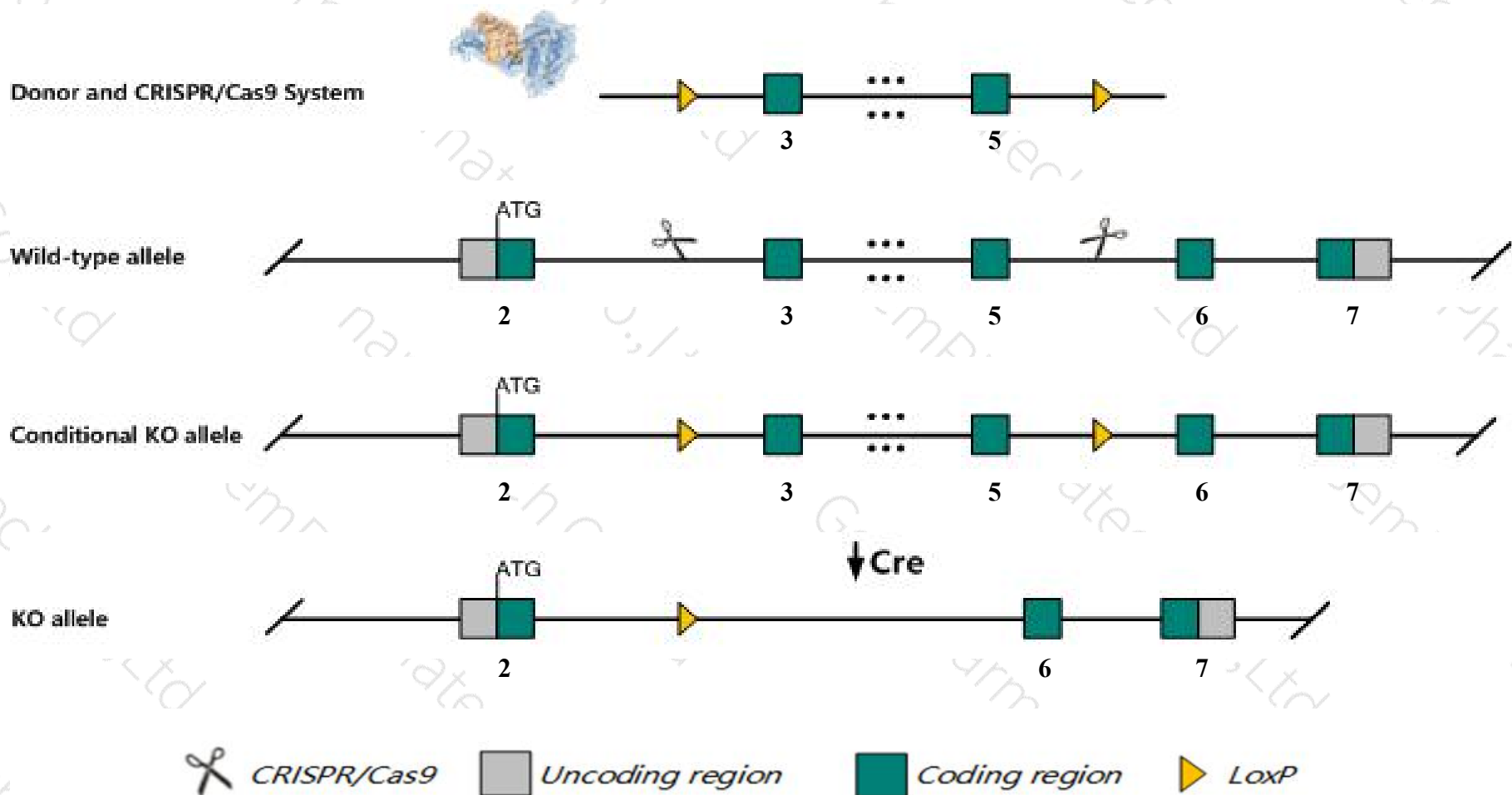
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Actc1* gene. The schematic diagram is as follows:



- The *Actc1* gene has 2 transcripts. According to the structure of *Actc1* gene, exon3-exon5 of *Actc1*-201 (ENSMUST00000090269.6) transcript is recommended as the knockout region. The region contains 679bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Actc1* 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.

- According to the existing MGI data, Homozygous null mutation of this gene results in embryonic and postnatal lethality. Animals that survive to birth die within the first 2 weeks and display reduced body size and heart muscle defects.
- The partial sequence of the intron of *A530058N18Rik* gene will be deleted.
- The floxed region is near to the N-terminal of *C130080G10Rik* gene, this strategy may influence the regulatory function of the N-terminal of *C130080G10Rik* gene.
- The effect on transcript *Actc1*-202 is unknown.
- The *Actc1* 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)

Actc1 actin, alpha, cardiac muscle 1 [*Mus musculus* (house mouse)]

Gene ID: 11464, updated on 9-Feb-2020

Summary

Official Symbol	Actc1 provided by MGI
Official Full Name	actin, alpha, cardiac muscle 1 provided by MGI
Primary source	MGI:MGI:87905
See related	Ensembl:ENSMUSG00000068614
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	Actc-1
Expression	Restricted expression toward heart adult (RPKM 9964.4) See more
Orthologs	human all

Genomic context

Location: 2 E4; 2 57.55 cM

See Actc1 in [Genome Data Viewer](#)

Exon count: 7

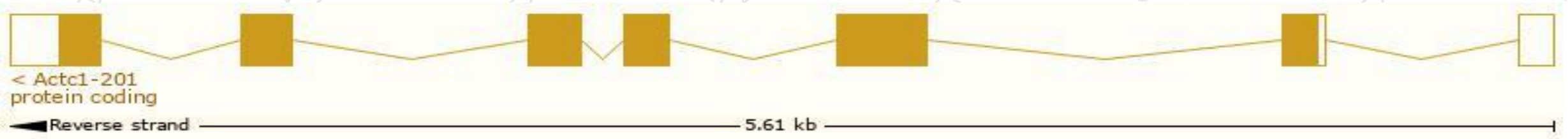
Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	2	NC_000068.7 (114047282..114052875, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	2	NC_000068.6 (113873025..113878547, complement)

Transcript information (Ensembl)

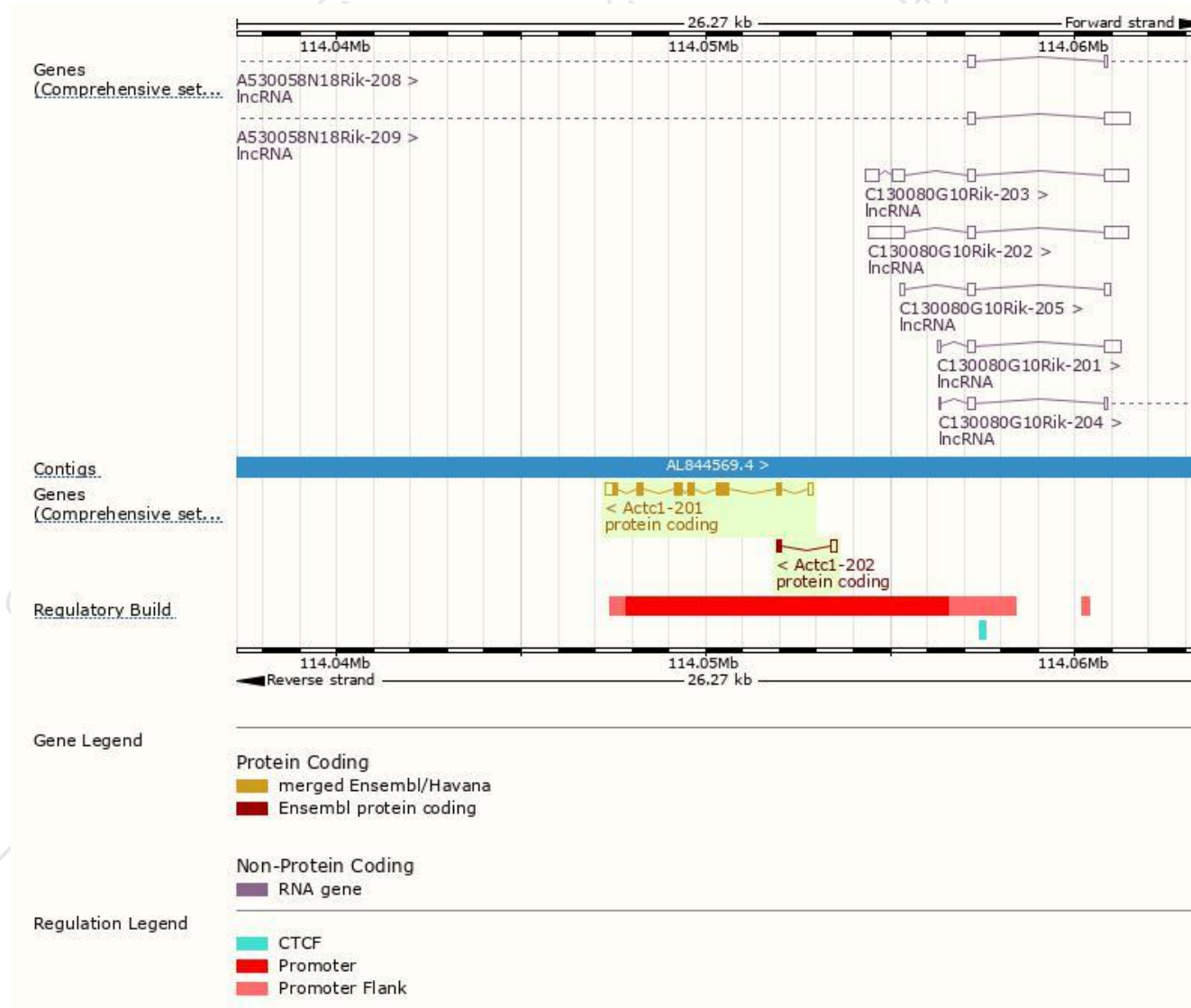
The gene has 2 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Actc1-201	ENSMUST00000090269.6	1473	377aa	Protein coding	CCDS16564	P68033 Q497E4	TSL:1 GENCODE basic APPRIS P1
Actc1-202	ENSMUST00000149125.1	303	43aa	Protein coding	-	F6WX90	CDS 3' incomplete TSL:3

The strategy is based on the design of *Actc1-201* transcript,The transcription is shown below



Genomic location distribution



Protein domain

ENSMUSP00000087...

Superfamily

SSF53067

SMART

Actin family

Prints

Actin family

Pfam

Actin family

PROSITE patterns

Actin, conserved site

Actin, c

Actin/actin-like conserved site

PANTHER

PTHR11937:SF416

Actin family

Gene3D

3.30.420.40

3.90.640.10

CDD

cd00012

All sequence SNPs/i...

Sequence variants (dbSNP and all other sources)

Variant Legend

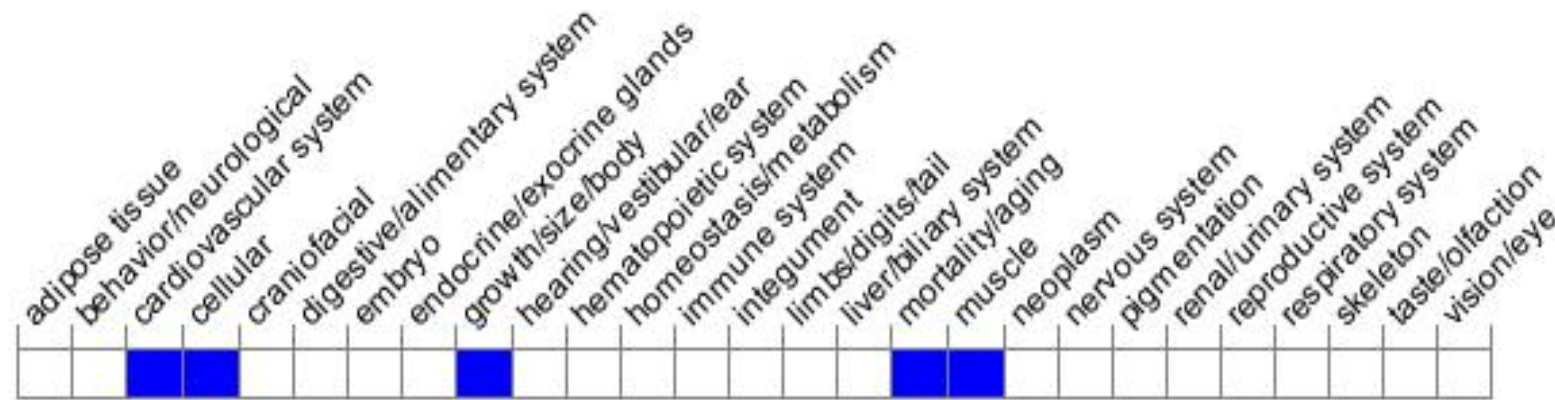
synonymous variant

Scale bar

0 40 80 120 160 200 240 280 320 377

Mouse phenotype description(MGI)

Phenotype Overview



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 mutation of this gene results in embryonic and postnatal lethality. Animals that survive to birth die within the first 2 weeks and display reduced body size and heart muscle defects.

If you have any questions, you are welcome to inquire.

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