

Slc8a1 Cas9-KO Strategy

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Reviewer:

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Design Date:

2019-10-30

Project Overview

Project Name

Slc8a1

Project type

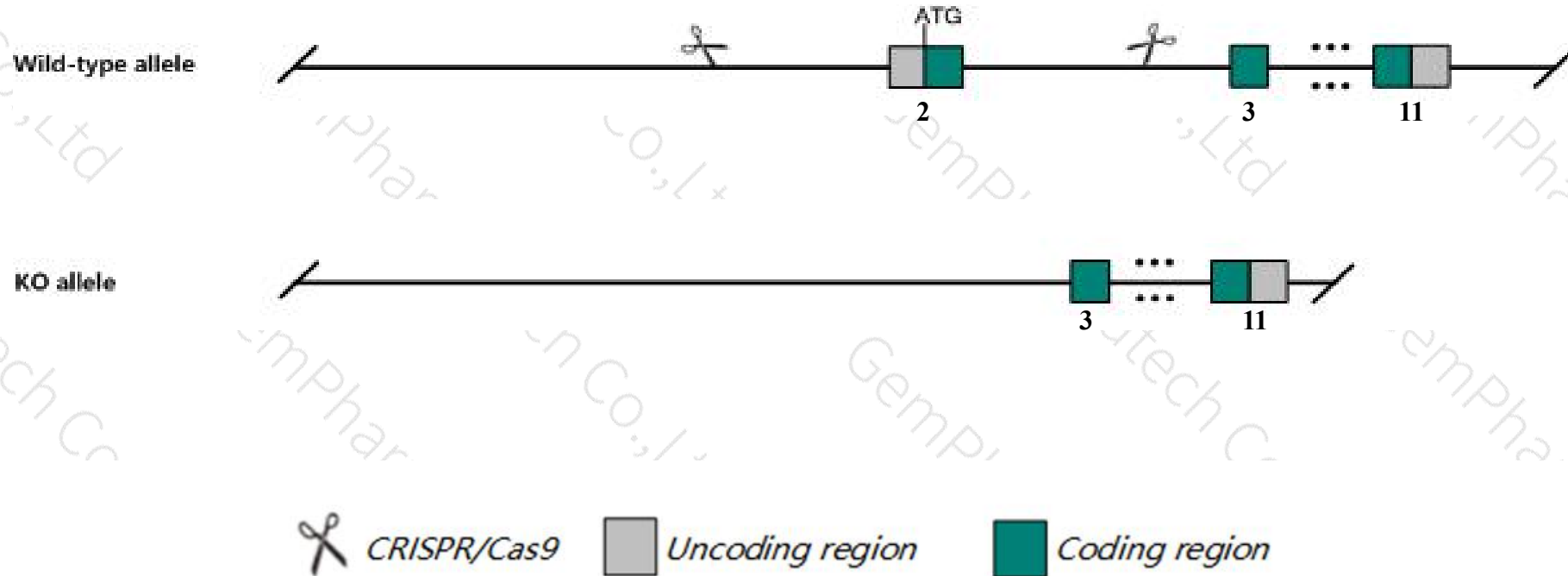
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slc8a1* gene. The schematic diagram is as follows:



- The *Slc8a1* gene has 6 transcripts. According to the structure of *Slc8a1* gene, exon2 of *Slc8a1-202* (ENSMUST00000163123.2) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc8a1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygotes for targeted null mutations have underdeveloped, nonbeating hearts with massive apoptosis of myocytes, a dilated pericardium and die around embryonic day 9.5. Heterozygotes exhibit altered responses to experimental cardiac pressure overload.
- The *Slc8a1* gene is located on the Chr17. 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.

Gene information (NCBI)

Slc8a1 solute carrier family 8 (sodium/calcium exchanger), member 1 [Mus musculus (house mouse)]

Gene ID: 20541, updated on 10-Feb-2019

Summary



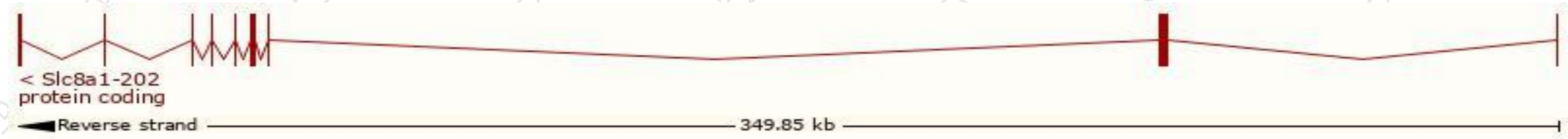
| | |
|---------------------------|---|
| Official Symbol | Slc8a1 provided by MGI |
| Official Full Name | solute carrier family 8 (sodium/calcium exchanger), member 1 provided by MGI |
| Primary source | MGI:MGI:107956 |
| See related | Ensembl:ENSMUSG00000054640 |
| Gene type | protein coding |
| 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 | AI852629, AV344025, D930008O12Rik, Ncx1 |
| Expression | Broad expression in heart adult (RPKM 10.8), frontal lobe adult (RPKM 7.0) and 16 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

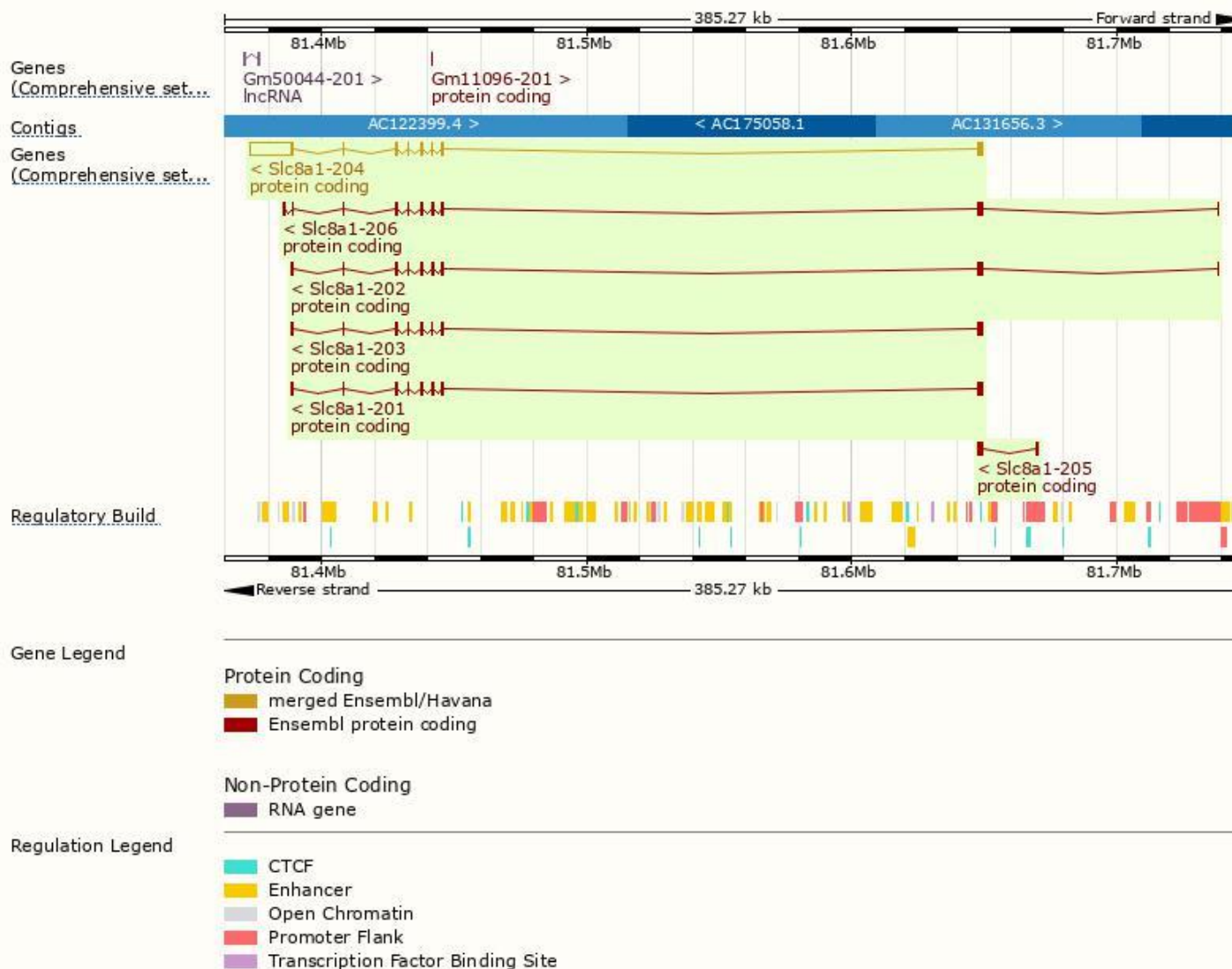
The gene has 6 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|------------|--------------------------------------|-------|-----------------------|----------------|---------------------------|------------------------|----------------------------------|
| Slc8a1-204 | ENSMUST00000234131.1 | 18464 | 958aa | Protein coding | CCDS50191 | - | GENCODE basic APPRIS ALT2 |
| Slc8a1-202 | ENSMUST00000163123.2 | 3180 | 970aa | Protein coding | CCDS37706 | Q68FL0 | TSL:1 GENCODE basic APPRIS P3 |
| Slc8a1-201 | ENSMUST00000086538.9 | 2913 | 970aa | Protein coding | CCDS37706 | G3X9J1 | TSL:5 GENCODE basic APPRIS P3 |
| Slc8a1-206 | ENSMUST00000235015.1 | 3339 | 940aa | Protein coding | - | - | GENCODE basic |
| Slc8a1-203 | ENSMUST00000163680.8 | 2913 | 970aa | Protein coding | - | G5E8Y0 | TSL:5 GENCODE basic APPRIS ALT 1 |
| Slc8a1-205 | ENSMUST00000234923.1 | 2017 | 599aa | Protein coding | - | - | CDS 3' incomplete |

The strategy is based on the design of *Slc8a1-202* transcript,The transcription is shown below



Genomic location distribution



Protein domain



集萃药康
GemPharmatech



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, Homozygotes for targeted null mutations have underdeveloped, nonbeating hearts with massive apoptosis of myocytes, a dilated pericardium and die around embryonic day 9.5. Heterozygotes exhibit altered responses to experimental cardiac pressure overload.

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

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