

***Guca1b* Cas9-KO Strategy**

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Project Overview

Project Name

Guca1b

Project type

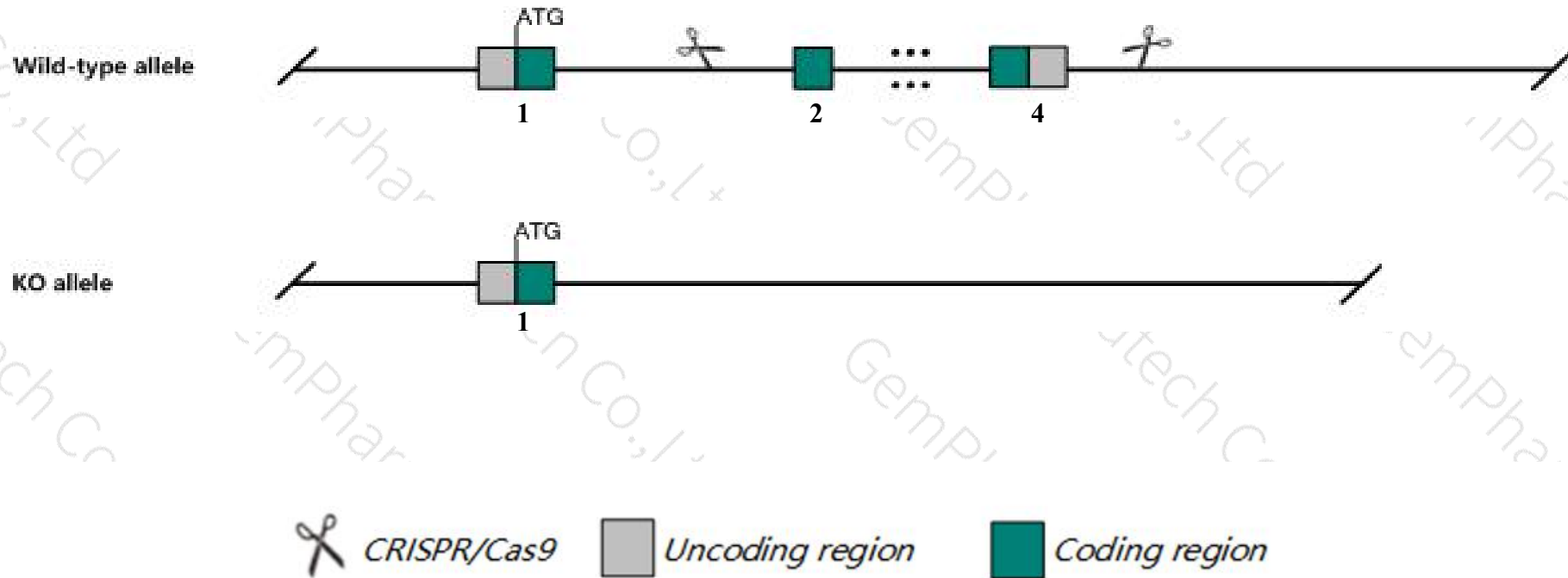
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Guca1b* gene. The schematic diagram is as follows:



- The *Guca1b* gene has 2 transcripts. According to the structure of *Guca1b* gene, exon2-exon4 of *Guca1b*-201 (ENSMUST00000024774.13) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Guca1b* gene. The brief process is as follows: gRNA was transcribed in vitro. Cas9 and gRNA 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.

- According to the existing MGI data, Mice homozygous for a null allele exhibit abnormal rod electrophysiology.
- The knockout region is near to the C-terminal of *AC112683.2* gene and *Guca1a* gene, this strategy may influence the regulatory function of the C-terminal of these genes.
- The *Guca1b* 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)

Guca1b guanylate cyclase activator 1B [*Mus musculus* (house mouse)]

Gene ID: 107477, updated on 19-Nov-2019

Summary

- Official Symbol
- Guca1b provided by [MGI](#)
- Official Full Name
- guanylate cyclase activator 1B provided by [MGI](#)
- Primary source
- [MGI:MGI:1194489](#)
- See related
- [Ensembl:ENSMUSG00000023979](#)
- 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
- GCAP2; BC018258
- Expression
- Ubiquitous expression in bladder adult (RPKM 1.3), adrenal adult (RPKM 0.7) and 28 other tissues [See more](#)
- Orthologs
- [human](#) [all](#)

Genomic context

Location: 17; 17 C

Exon count: 4

See Guca1b in [Genome Data Viewer](#)

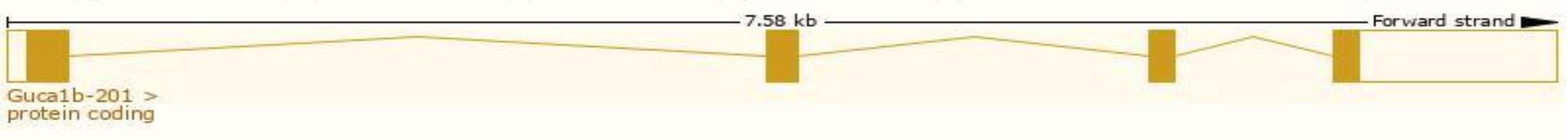
Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	17	NC_000083.6 (47384694..47392967)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	17	NC_000083.5 (47522342..47529916)

Transcript information (Ensembl)

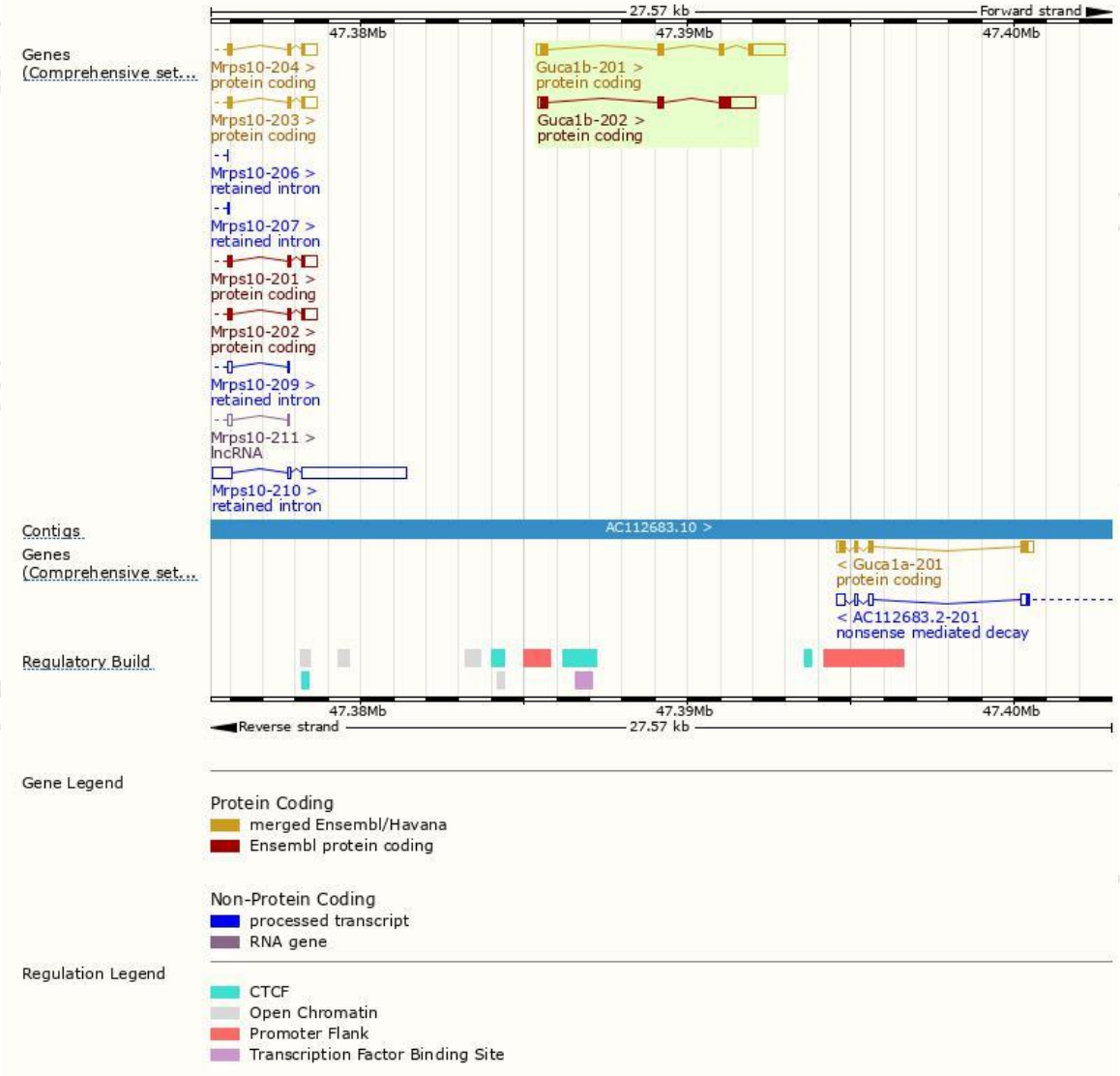
The gene has 2 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Guca1b-201	ENSMUST00000024774.13	1668	201aa	Protein coding	CCDS28847	Q8VBV8	TSL:1 GENCODE basic APPRIS P1
Guca1b-202	ENSMUST00000145462.1	1556	230aa	Protein coding	-	E9PWU1	TSL:2 GENCODE basic

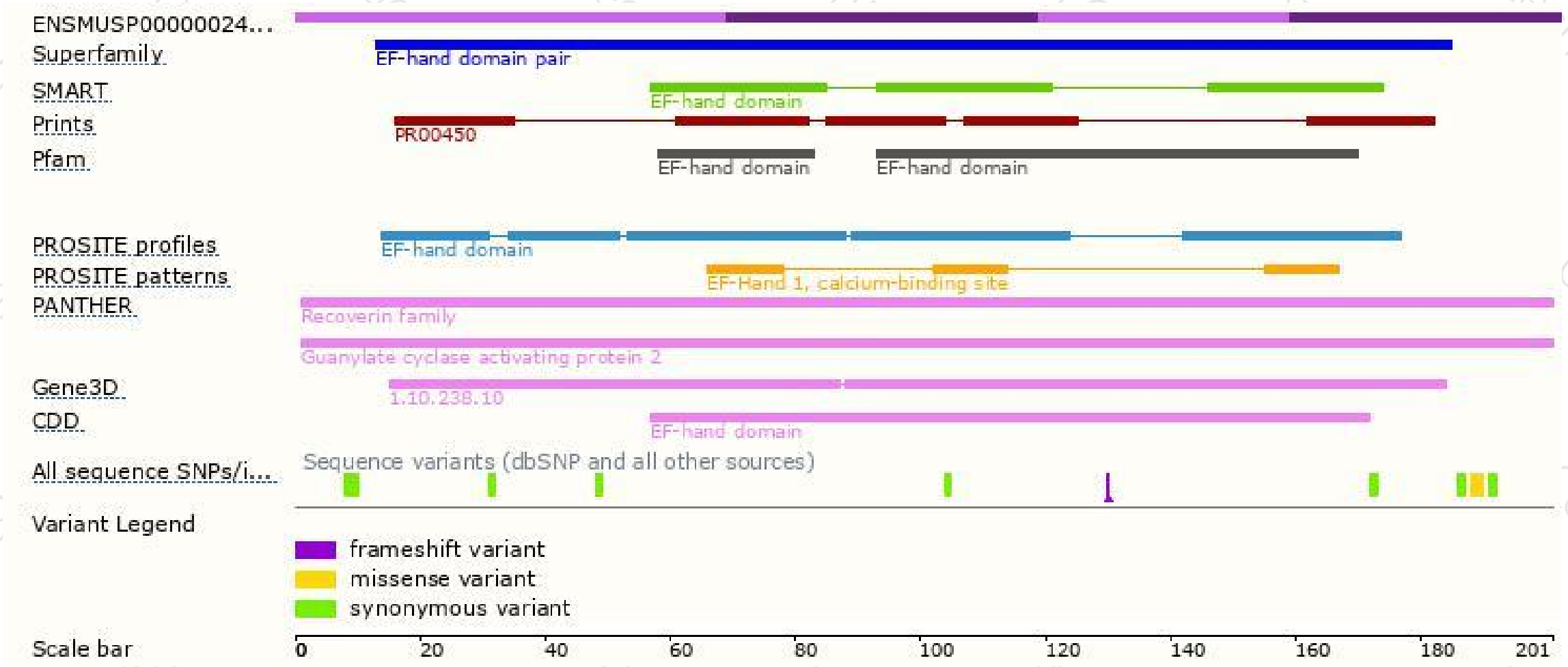
The strategy is based on the design of *Guca1b-201* transcript,The transcription is shown below



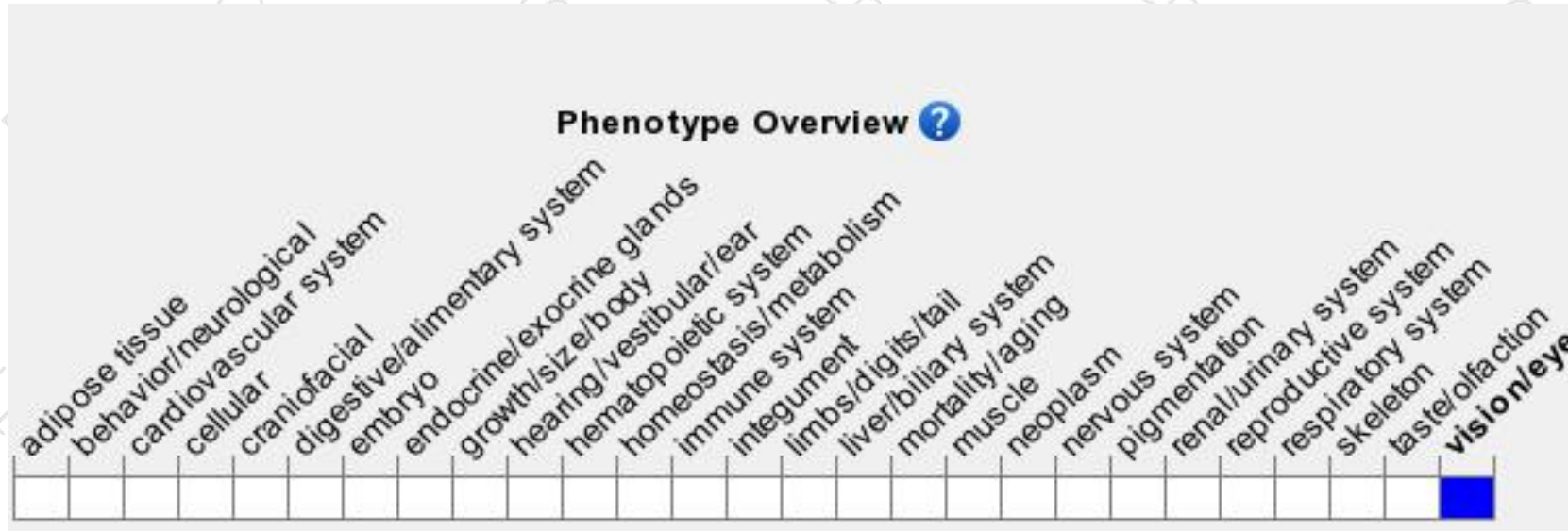
Genomic location distribution



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, Mice homozygous for a null allele exhibit abnormal rod electrophysiology.

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

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