

# Calcr Cas9-KO Strategy

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Design Date: 2019-8-5

# **Project Overview**



Project Name Calcr

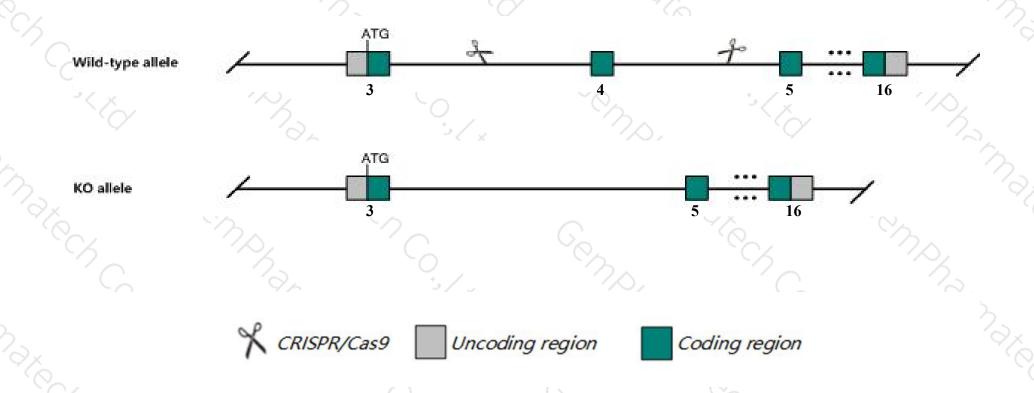
Project type Cas9-KO

Strain background C57BL/6JGpt

# **Knockout strategy**



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



### **Technical routes**



- ➤ The *Calcr* gene has 5 transcripts. According to the structure of *Calcr* gene, exon4 of *Calcr-201*(ENSMUST00000075644.12) transcript is recommended as the knockout region. The region contains 79bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Calcr* gene. The brief process is as follows: CRISPR/Cas9 system

### **Notice**



- ➤ According to the existing MGI data, Haploinsufficiency may result in increased bone density due to increased bone formation. Homozygous inactivation may result in embryonic lethality. Mice homozygous for another disruption allele at this locus show a normal phenotype.
- The knockout region is near to the N-terminal of *Mir653* and *Mir489* gene, this strategy may influence the regulatory function of the N-terminal of these gene.
- The *Calcr* gene is located on the Chr6. 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)



#### Calcr calcitonin receptor [Mus musculus (house mouse)]

Gene ID: 12311, updated on 19-Mar-2019

#### Summary

☆ ?

Official Symbol Calcr provided by MGI

Official Full Name calcitonin receptor provided by MGI

Primary source MGI:MGI:101950

See related Ensembl: ENSMUSG00000023964

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 Clr, Ct-r

Expression Biased expression in whole brain E14.5 (RPKM 1.5), CNS E18 (RPKM 1.2) and 5 other tissuesSee more

Orthologs human all

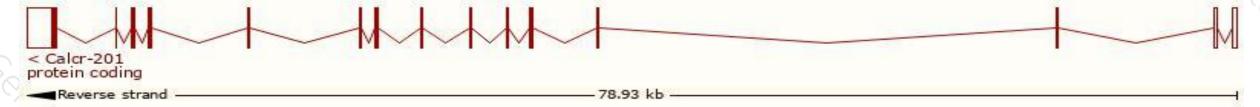
# Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

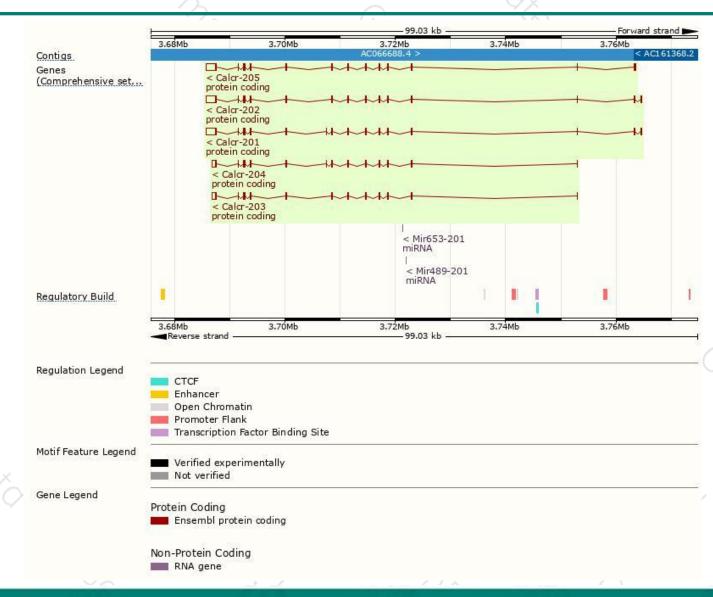
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Calcr-201	ENSMUST00000075644.12	3763	<u>533aa</u>	Protein coding	CCDS39417	Q60755	TSL:1 GENCODE basic APPRIS P4
Calcr-202	ENSMUST00000115622.7	3721	496aa	Protein coding	CCDS39416	Q60755	TSL:1 GENCODE basic APPRIS ALT2
Calcr-205	ENSMUST00000171613.7	3518	496aa	Protein coding	CCDS39416	Q60755	TSL:1 GENCODE basic APPRIS ALT2
Calcr-204	ENSMUST00000170266.2	2175	<u>533aa</u>	Protein coding	CCDS39417	Q60755	TSL:1 GENCODE basic APPRIS P4
Calcr-203	ENSMUST00000168592.8	2064	496aa	Protein coding	CCDS39416	Q60755	TSL:5 GENCODE basic APPRIS ALT2

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



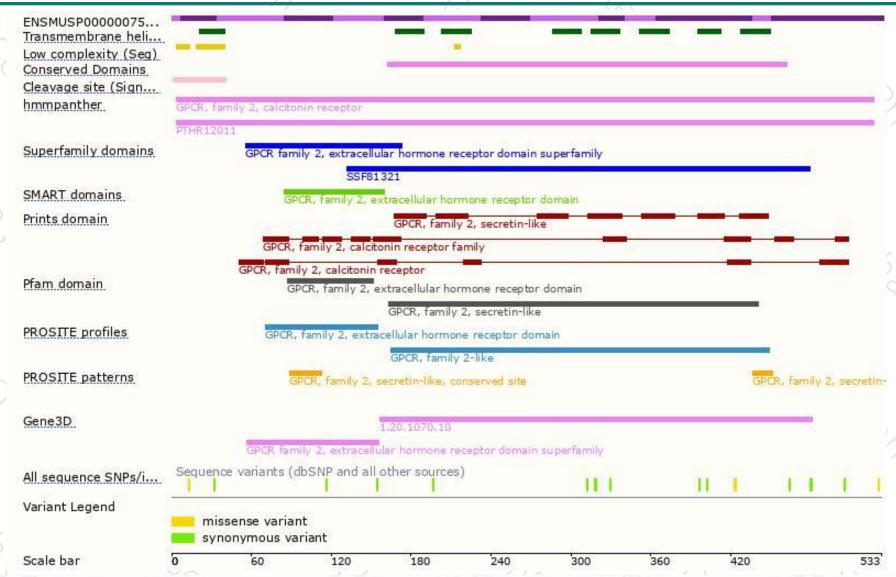
### Genomic location distribution





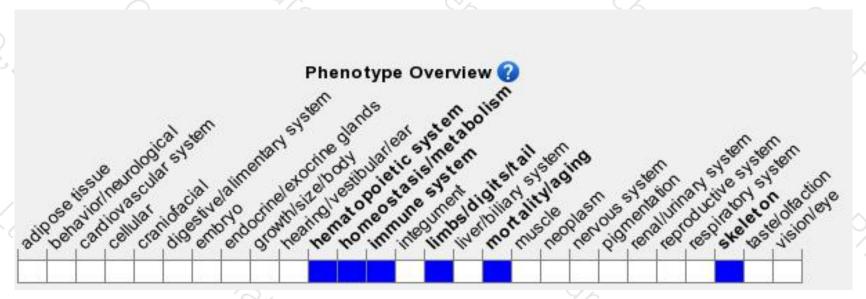
### Protein domain





# 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, Haploinsufficiency may result in increased bone density due to increased bone formation. Homozygous inactivation may result in embryonic lethality. Mice homozygous for another disruption allele at this locus show a normal phenotype.



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





