

Tcirg1 Cas9-KO Strategy

Designer:

Project Overview

Project Name

Tcirg1

Project type

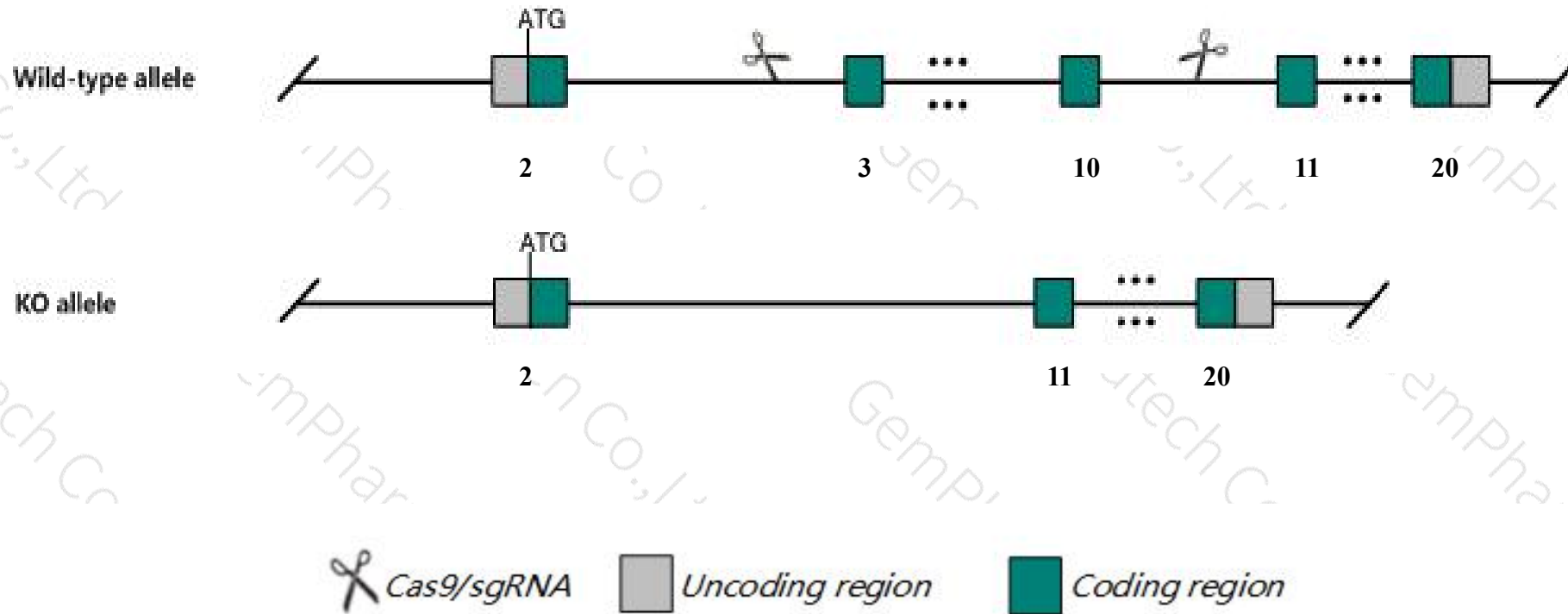
Cas9-KO

Strain background

C57BL/6J

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Tcirg1* gene. The schematic diagram is as follows:



- The *Tcirg1* gene has 14 transcripts. According to the structure of *Tcirg1* gene, exon3-exon10 of *Tcirg1*-204 (ENSMUST00000126070.8) transcript is recommended as the knockout region. The region contains 1051bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Tcirg1* gene. The brief process is as follows: sgRNA was transcribed in vitro. Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

- According to the existing MGI data, Homozygotes for mutant alleles exhibit severe osteopetrosis with increased bone density due to failure of secondary bone resorption. Mutants lack teeth and die around 30-40 days of age.
- The *Tcirg1* gene is located on the Chr19. 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)

Tcirg1 T cell, immune regulator 1, ATPase, H⁺ transporting, lysosomal V0 protein A3 [Mus musculus (house mouse)]

Gene ID: 27060, updated on 3-Feb-2019

Summary



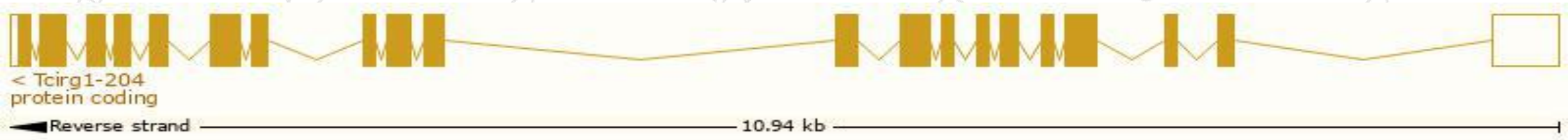
Official Symbol	Tcirg1 provided by MGI
Official Full Name	T cell, immune regulator 1, ATPase, H ⁺ transporting, lysosomal V0 protein A3 provided by MGI
Primary source	MGI:MGI:1350931
See related	Ensembl:ENSMUSG00000001750
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	ATP6N1C, ATP6a3, Atp6i, OC-116, OPTB1, Stv1, TIRC7, Vph1, oc
Expression	Ubiquitous expression in spleen adult (RPKM 61.8), adrenal adult (RPKM 46.6) and 27 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

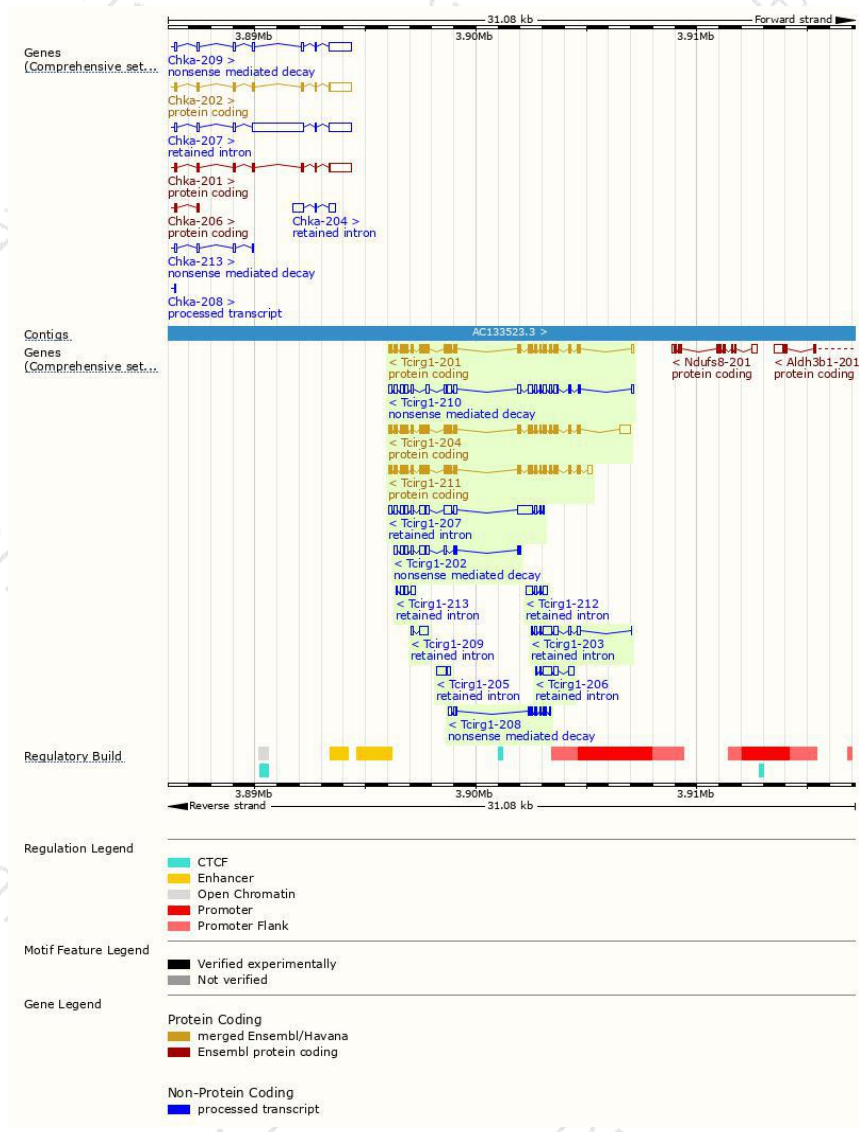
The gene has 14 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Tcirg1-204	ENSMUST00000126070.8	3041	834aa	Protein coding	CCDS29401	Q9JHF5	TSL:1 GENCODE basic APPRIS P1
Tcirg1-211	ENSMUST00000145791.7	2729	834aa	Protein coding	CCDS29401	Q9JHF5	TSL:1 GENCODE basic APPRIS P1
Tcirg1-201	ENSMUST00000001801.10	2688	834aa	Protein coding	CCDS29401	Q9JHF5	TSL:1 GENCODE basic APPRIS P1
Tcirg1-214	ENSMUST00000236590.1	333	65aa	Protein coding	-	-	CDS 3' incomplete
Tcirg1-210	ENSMUST00000135070.7	2424	72aa	Nonsense mediated decay	-	D6RFN1	TSL:5
Tcirg1-202	ENSMUST00000122885.7	1233	91aa	Nonsense mediated decay	-	F6ZFBB	CDS 5' incomplete TSL:5
Tcirg1-208	ENSMUST00000132164.1	813	191aa	Nonsense mediated decay	-	F6XRE6	CDS 5' incomplete TSL:5
Tcirg1-207	ENSMUST00000131327.7	2350	No protein	Retained intron	-	-	TSL:2
Tcirg1-203	ENSMUST00000125792.8	1066	No protein	Retained intron	-	-	TSL:5
Tcirg1-206	ENSMUST00000127308.1	923	No protein	Retained intron	-	-	TSL:5
Tcirg1-212	ENSMUST00000159824.7	692	No protein	Retained intron	-	-	TSL:2
Tcirg1-205	ENSMUST00000126643.2	568	No protein	Retained intron	-	-	TSL:2
Tcirg1-213	ENSMUST00000162688.1	467	No protein	Retained intron	-	-	TSL:1
Tcirg1-209	ENSMUST00000134698.1	463	No protein	Retained intron	-	-	TSL:3

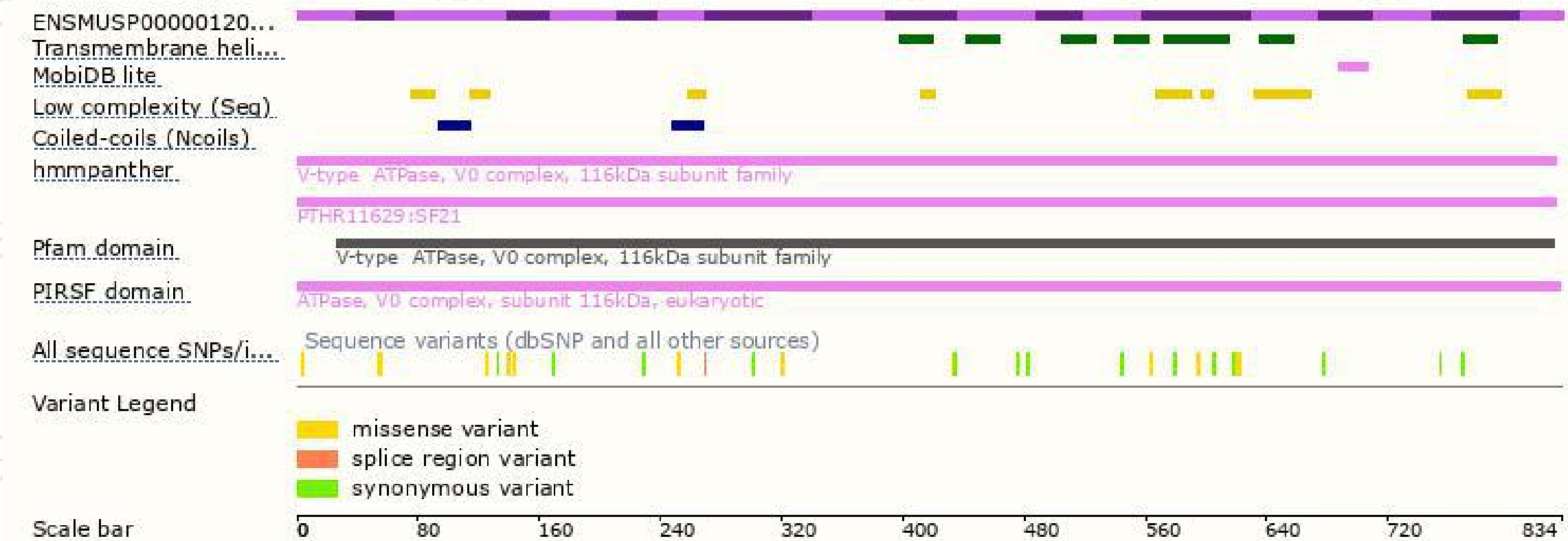
The strategy is based on the design of *Tcirg1-204* transcript,The transcription is shown below



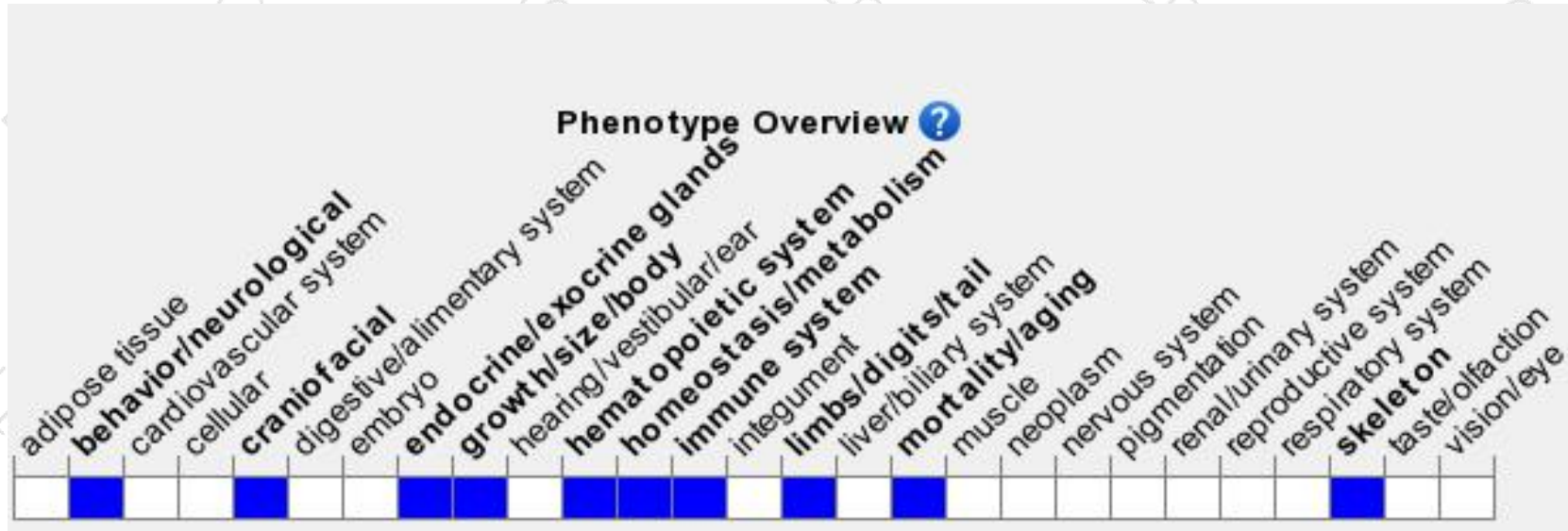
Genomic location distribution



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, Homozygotes for mutant alleles exhibit severe osteopetrosis with increased bone density due to failure of secondary bone resorption. Mutants lack teeth and die around 30-40 days of age.

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

Tel: 025-5864 1534

