

Ddr2 Cas9-KO Strategy

Designer: Huimin Su

Reviewer: Ruirui Zhang

Date: 2019/9/25

Project Overview

Project Name

Ddr2

Project type

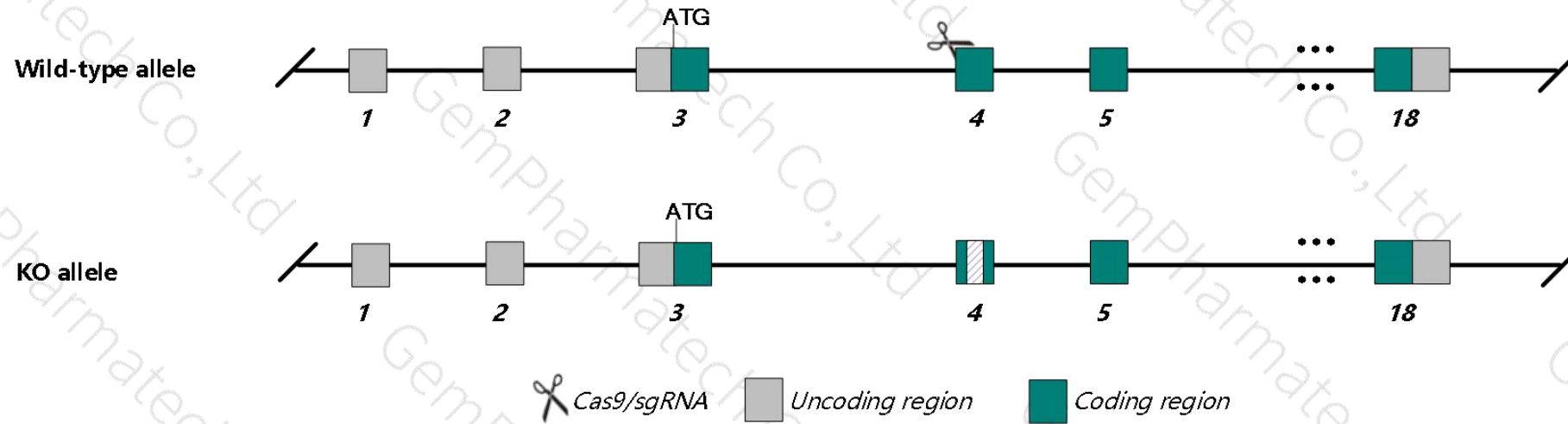
Cas9-KO

Strain background

C57BL/6N

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Ddr2* gene. The schematic diagram is as follows:



- In this project we use CRISPR/Cas9 technology to modify *Ddr2* gene. The brief process is as follows: sgRNA was transcribed in vitro. Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6N 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/6N mice.

- According to the MGI data, Homozygotes for a null allele show dwarfism, reduced chondrocyte proliferation, shortened long bones and snout, and skull anomalies. Homozygotes for another null allele show similar skeletal defects, small hearts, short cardiomyocytes, lower cardiac collagen density, and altered cardiac function.
- The *Ddr2* gene is located on the Chr1. 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)

Ddr2 discoidin domain receptor family, member 2 [*Mus musculus* (house mouse)]

Gene ID: 18214, updated on 3-Sep-2019

Summary

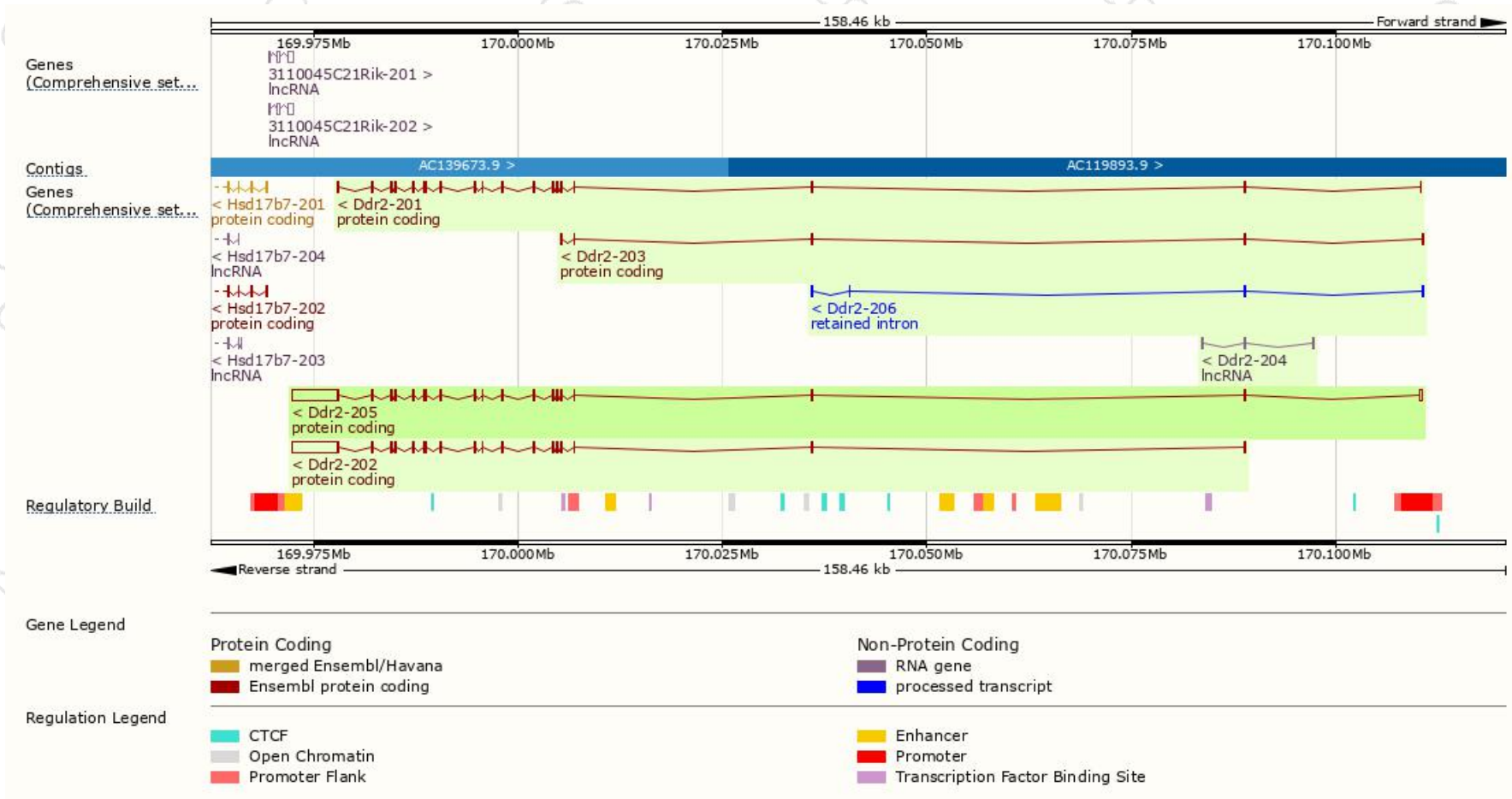
Official Symbol	Ddr2 provided by MGI
Official Full Name	discoidin domain receptor family, member 2 provided by MGI
Primary source	MGI:MGI:1345277
See related	Ensembl:ENSMUSG00000026674
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	Ntrkr3; tyro10; AW495251
Expression	Biased expression in bladder adult (RPKM 28.6), limb E14.5 (RPKM 18.7) and 13 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

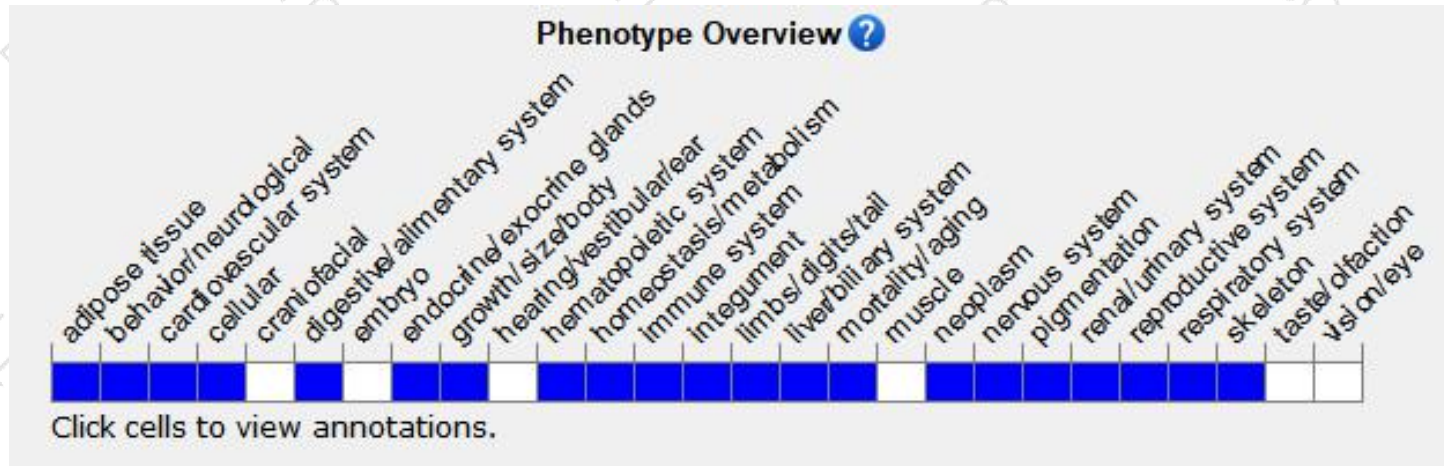
The gene has 6 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ddr2-205	ENSMUST00000194690.5	8590	854aa	Protein coding	CCDS48436	Q62371	TSL:1 GENCODE basic APPRIS P1
Ddr2-202	ENSMUST00000170800.7	8289	854aa	Protein coding	CCDS48436	Q62371	TSL:5 GENCODE basic APPRIS P1
Ddr2-201	ENSMUST00000027985.7	2928	854aa	Protein coding	CCDS48436	Q62371	TSL:1 GENCODE basic APPRIS P1
Ddr2-203	ENSMUST00000192312.5	672	129aa	Protein coding	-	A0A0A6YXY2	CDS 3' incomplete TSL:3
Ddr2-206	ENSMUST00000195867.5	641	No protein	Retained intron	-	-	TSL:3
Ddr2-204	ENSMUST00000194619.1	442	No protein	lncRNA	-	-	TSL:5

Genomic location distribution



Mouse phenotype description(MGI)



Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(<http://www.informatics.jax.org/>).

Homozygous null mice show Parkinson's disease like symptoms, including akinetic and bradykinetic behavior. Mice lacking only the long isoform are hypoactive and exhibit increased stereotypic behavior in response to dopamine agonists.

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

Tel: 025-5864 1534

