

Myh14 Cas9-CKO Strategy

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Reviewer:

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Project Overview

Project Name

Myh14

Project type

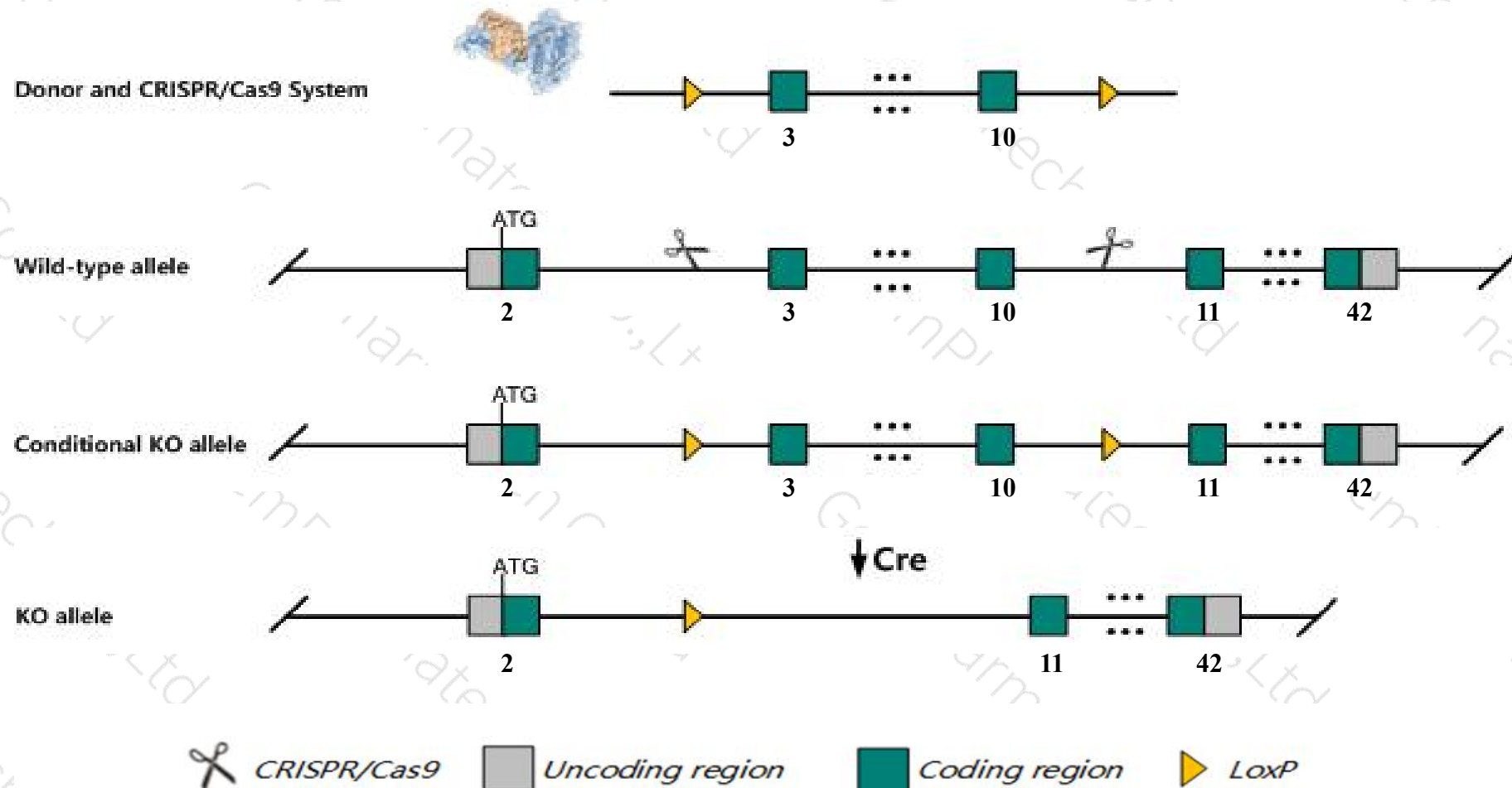
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Myh14* gene. The schematic diagram is as follows:



Technical routes

- The *Myh14* gene has 9 transcripts. According to the structure of *Myh14* gene, exon3-exon10 of *Myh14-201* (ENSMUST00000048102.14) transcript is recommended as the knockout region. The region contains 709bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Myh14* gene. The brief process is as follows: CRISPR/Cas9 system and Donor 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.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

Notice

- According to the existing MGI data, Mice homozygous for a knock-out allele are healthy and survive to adulthood with no apparent defects. About 30% of knock-in mice either heterozygous or homozygous for a single amino acid mutation exhibit increased lymphoma incidence.
- Transcript *Myh14-207* may not be affected.
- The *Myh14* gene is located on the Chr7. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)

Myh14 myosin, heavy polypeptide 14 [Mus musculus (house mouse)]

Gene ID: 71960, updated on 31-Jan-2019

Summary



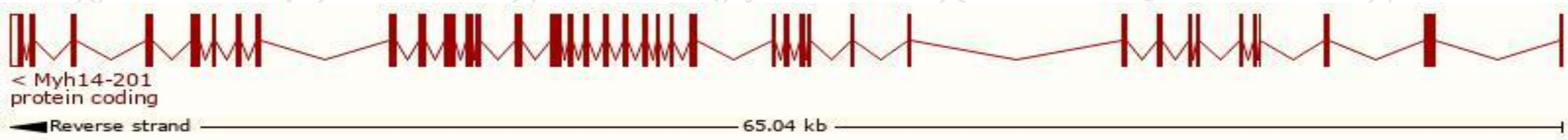
Official Symbol	Myh14 provided by MGI
Official Full Name	myosin, heavy polypeptide 14 provided by MGI
Primary source	MGI:MGI:1919210
See related	Ensembl:ENSMUSG00000030739
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	2400004E04Rik, II-C, NHMCII, NMHC II-C
Expression	Biased expression in small intestine adult (RPKM 46.6), colon adult (RPKM 41.7) and 8 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

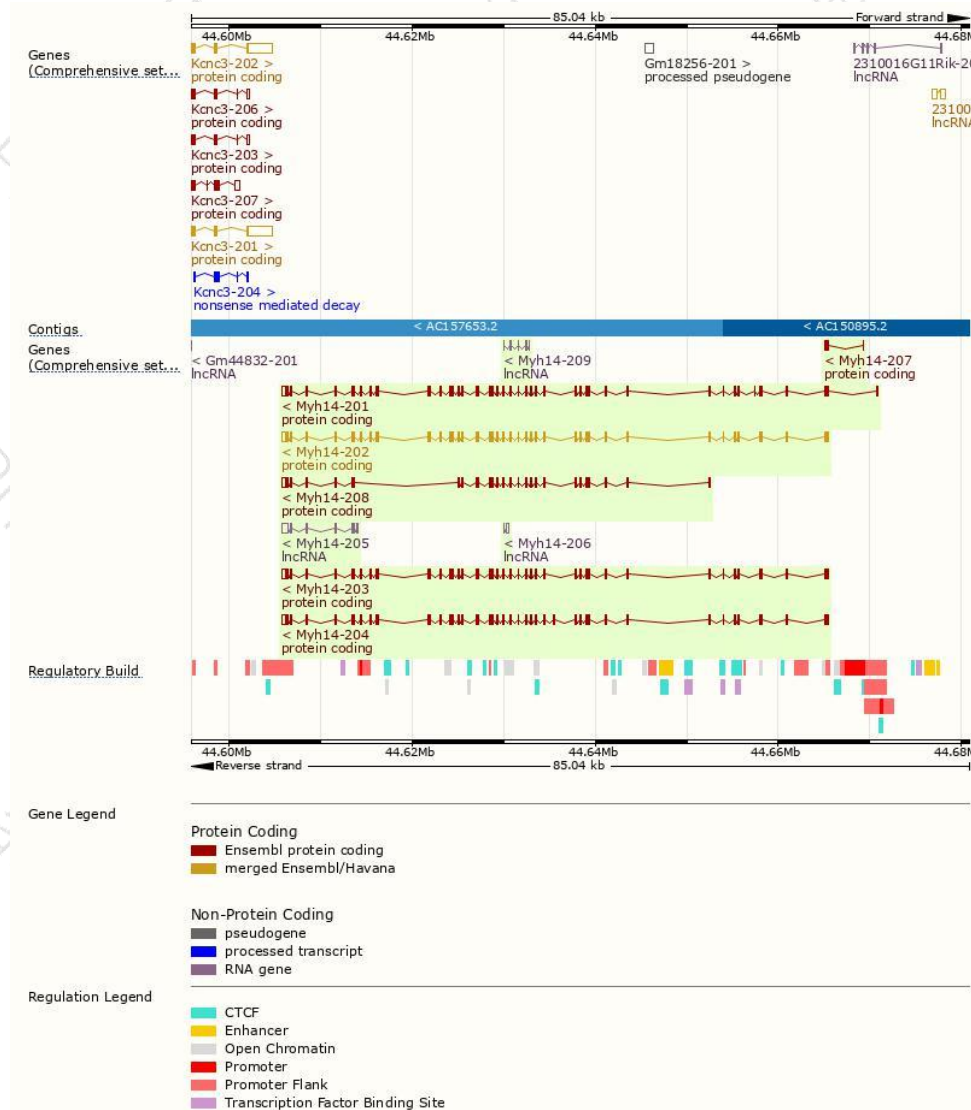
The gene has 9 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Myh14-201	ENSMUST00000048102.14	6457	2000aa	Protein coding	CCDS71951	Q6URW6	TSL:1 GENCODE basic APPRIS ALT2
Myh14-202	ENSMUST00000107899.9	6390	1992aa	Protein coding	CCDS39943	Q6URW6	TSL:1 GENCODE basic APPRIS P3
Myh14-204	ENSMUST00000207775.1	6499	2033aa	Protein coding	-	Q6URW6	TSL:5 GENCODE basic APPRIS ALT2
Myh14-203	ENSMUST00000107900.3	6400	2000aa	Protein coding	-	K3W4R2	TSL:5 GENCODE basic APPRIS ALT2
Myh14-208	ENSMUST00000208200.1	4143	1245aa	Protein coding	-	A0A140LI60	CDS 5' incomplete TSL:1
Myh14-207	ENSMUST00000208131.1	471	120aa	Protein coding	-	A0A140LHX1	CDS 3' incomplete TSL:2
Myh14-205	ENSMUST00000208044.1	1438	No protein	lncRNA	-	-	TSL:1
Myh14-209	ENSMUST00000209024.1	535	No protein	lncRNA	-	-	TSL:3
Myh14-206	ENSMUST00000208085.1	380	No protein	lncRNA	-	-	TSL:3

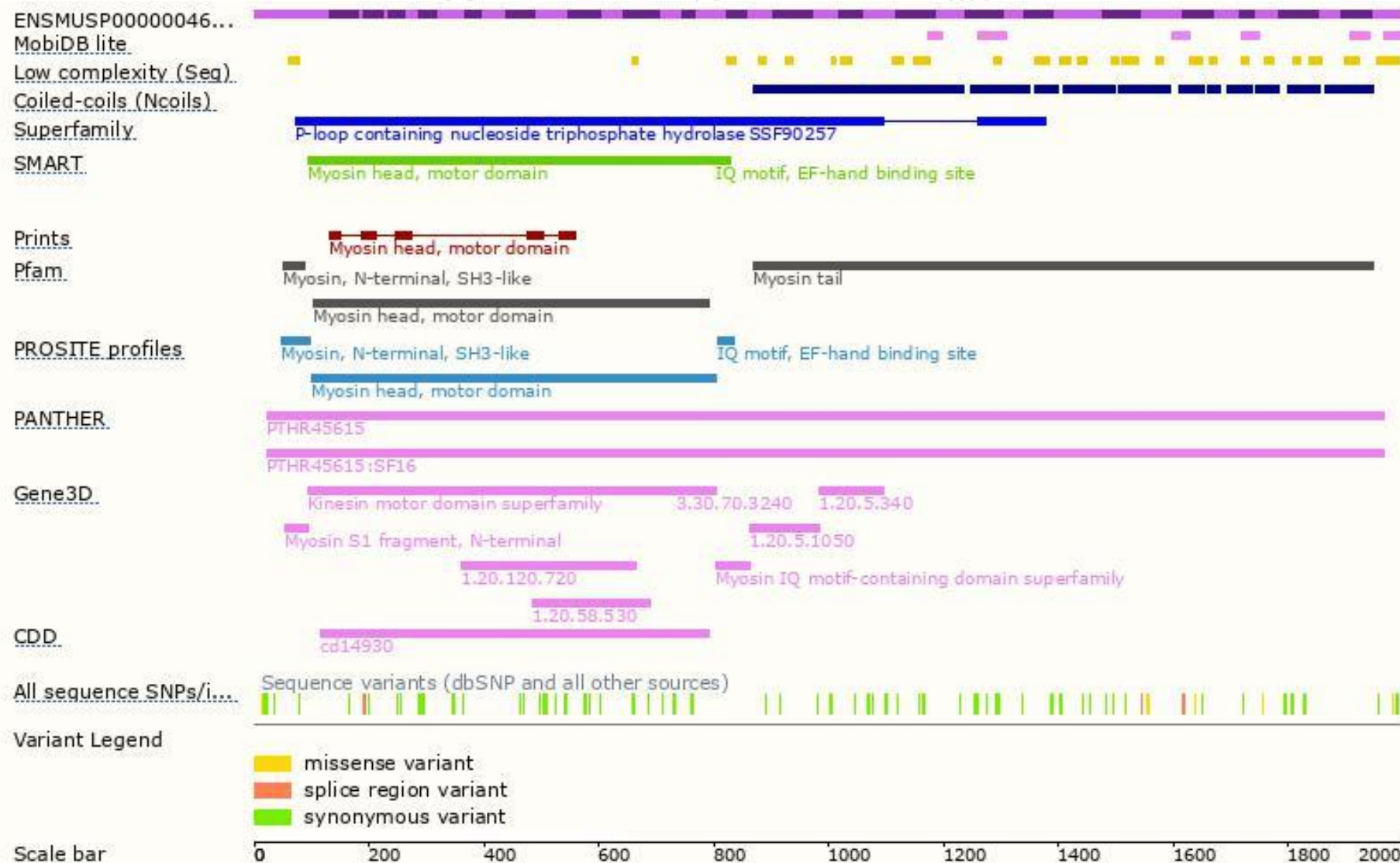
The strategy is based on the design of *Myh14-201* transcript,The transcription is shown below



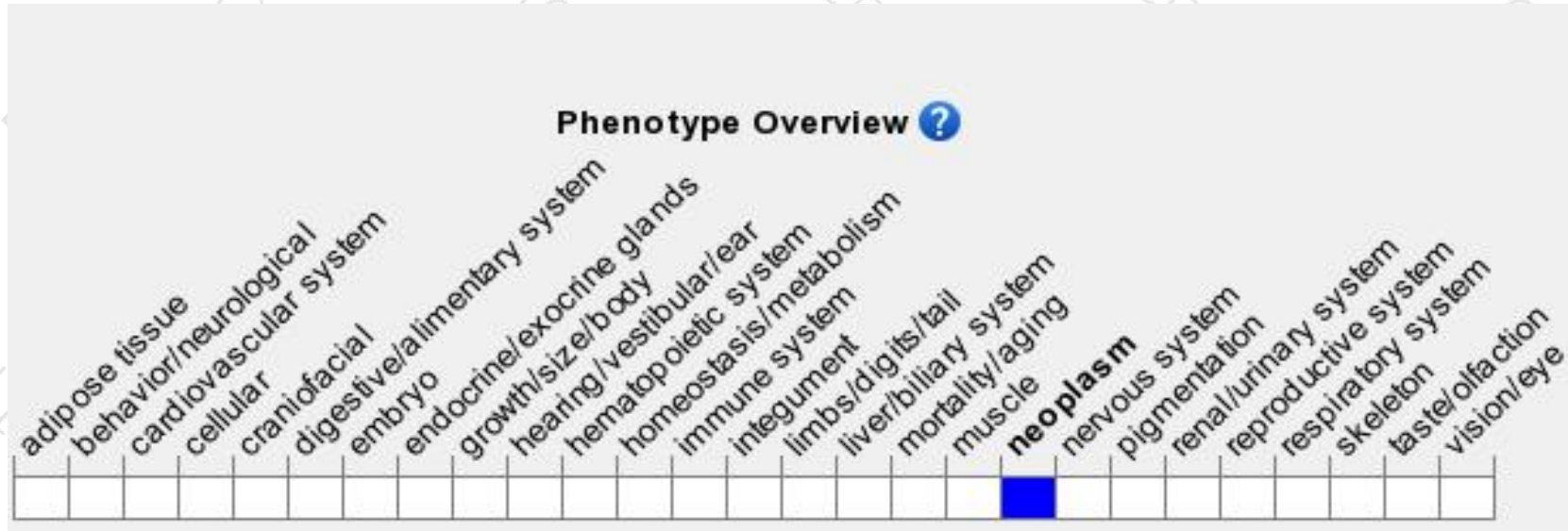
Genomic location distribution



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/>).

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

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