Myh10 Cas9-KO Strategy

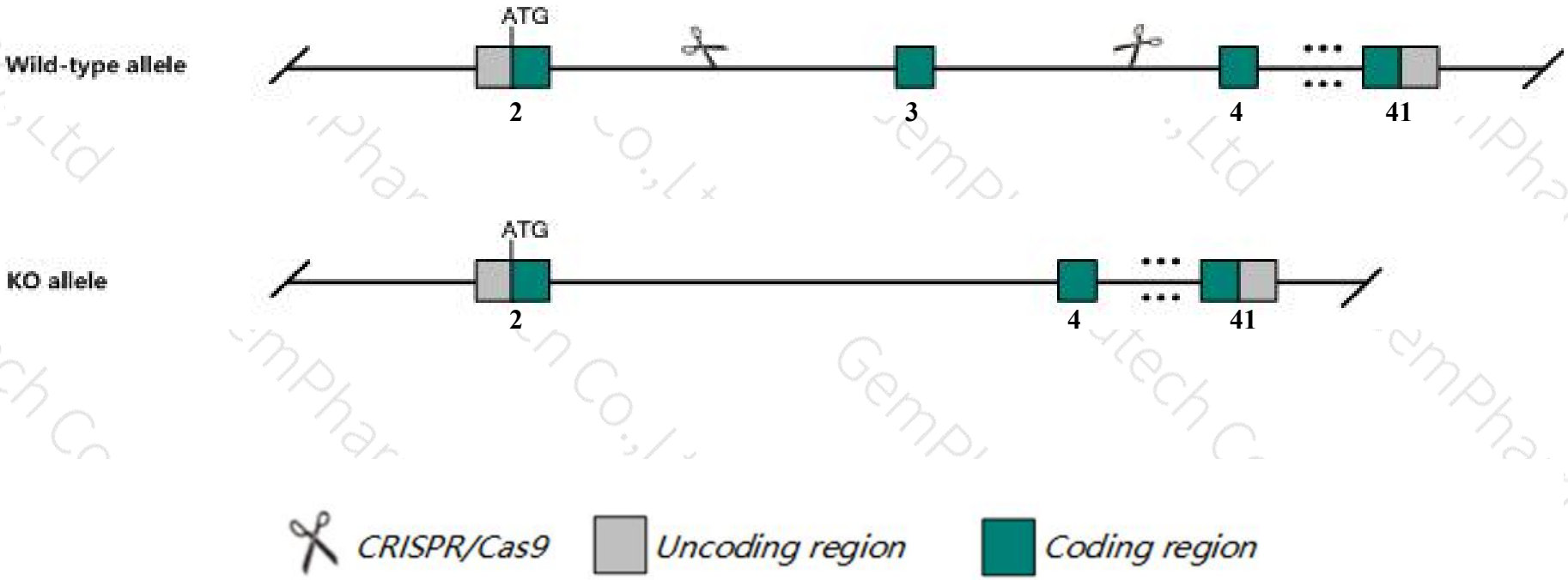
Designer:Yanhua Shen

Project Overview

Project Name	***Myh10***
Project type	**Cas9-KO**
Strain background	**C57BL/6JGpt**

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Myh10* gene. The schematic diagram is as follows:



Technical routes

➢ The *Myh10* gene has 10 transcripts. According to the structure of *Myh10* gene, exon3 of *Myh10-203* (ENSMUST00000102611.9) transcript is recommended as the knockout region.The region contains 157bp coding sequence. Knock out the region will result in disruption of protein function.

➢ In this project we use CRISPR/Cas9 technology to modify *Myh10* gene. The brief process is as follows: CRISPR/Cas9 systen

Notice

➢ According to the existing MGI data,Nullizygous mice show pre- and neonatal death, heart defects and hydrocephaly. Deletion of exon B1 disrupts migration of facial neurons, whereas deletion of exon B2 leads to Purkinje cell anomalies. Hypomorphs show hydrocephaly and defects in motor control, cerebellar foliation and neuron migration.

➢ The *Myh10* gene is located on the Chr11. 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）

集萃药康 GemPharmatech

Myh10 myosin, heavy polypeptide 10, non-muscle [Mus musculus (house mouse)]

Gene ID: 77579, updated on 7-Apr-2019

Summary

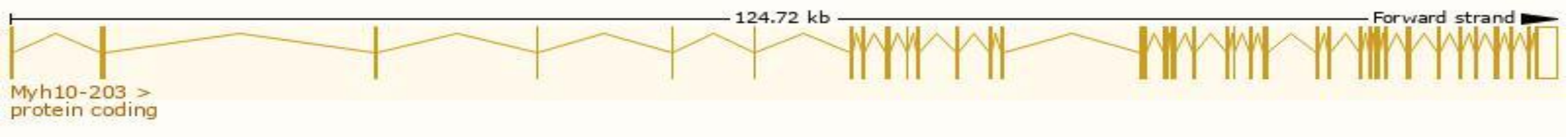
Official Symbol	Myh10 provided by MGI
Official Full Name	myosin, heavy polypeptide 10, non-muscle provided by MGI
Primary source	MGI:MGI:1930780
See related	Ensembl:ENSMUSG00000020900
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	5730504C04Rik, 9330167F11Rik, Fltn, Myhn-2, Myhn2, NMHC II-B, NMHC-B, NMHCII-B, NMMHC II-b, NMMHC-B, NMMHC-IIB, SMemb, mKIAA3005
Expression	Broad expression in CNS E11.5 (RPKM 43.2), CNS E14 (RPKM 26.8) and 19 other tissues See more
Orthologs	human all

Transcript information（Ensembl）

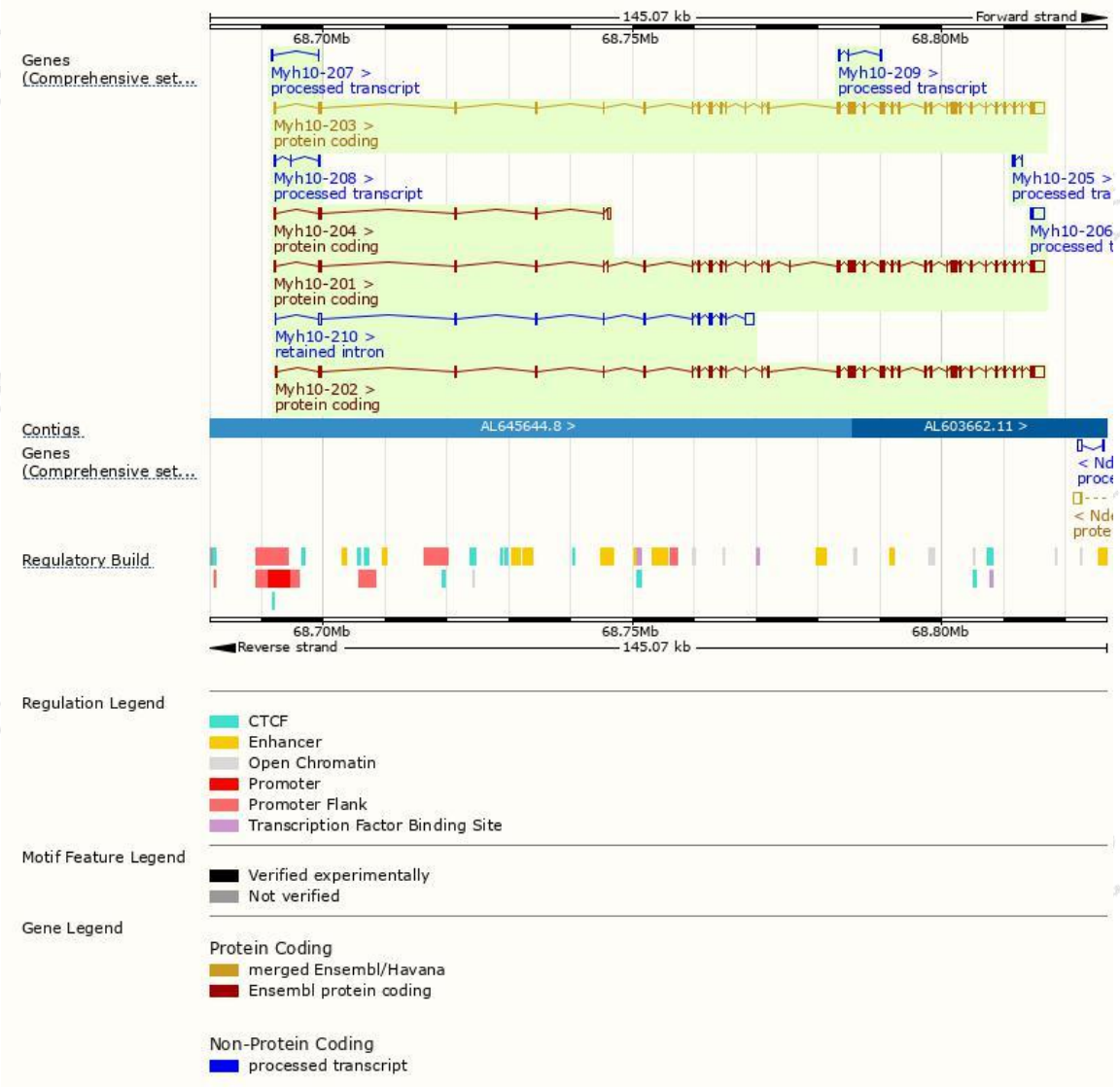
The gene has 10 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Myh10-203	ENSMUST00000102611.9	7791	1976aa	Protein coding	CCDS24869	Q61879	TSL:1 GENCODE basic APPRIS P2
Myh10-202	ENSMUST00000092984.5	7759	2013aa	Protein coding	-	Q3UH59	TSL:1 GENCODE basic
Myh10-201	ENSMUST00000018887.14	7667	2007aa	Protein coding	-	Q5SV64	TSL:5 GENCODE basic APPRIS ALT1
Myh10-204	ENSMUST00000108673.7	1315	231aa	Protein coding	-	Q8BXF2	TSL:1 GENCODE basic
Myh10-206	ENSMUST00000139059.1	1811	No protein	Processed transcript	-	-	TSL:1
Myh10-205	ENSMUST00000124006.1	558	No protein	Processed transcript	-	-	TSL:2
Myh10-208	ENSMUST00000143032.1	376	No protein	Processed transcript	-	-	TSL:2
Myh10-207	ENSMUST00000141558.1	334	No protein	Processed transcript	-	-	TSL:3
Myh10-209	ENSMUST00000145408.1	278	No protein	Processed transcript	-	-	TSL:3
Myh10-210	ENSMUST00000155765.1	2906	No protein	Retained intron	-	-	TSL:5

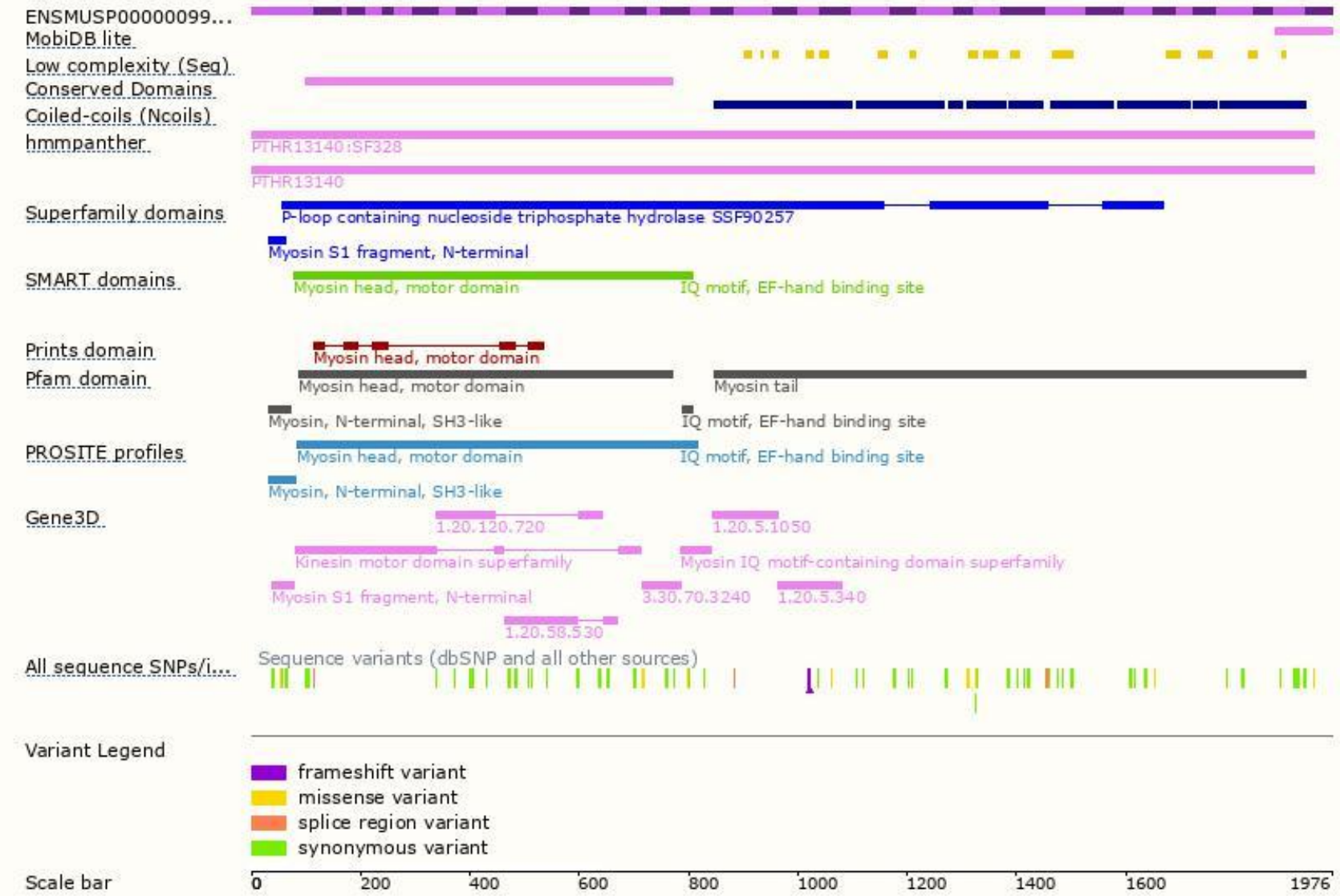
The strategy is based on the design of *Myh10-203* transcript,The transcription is shown below



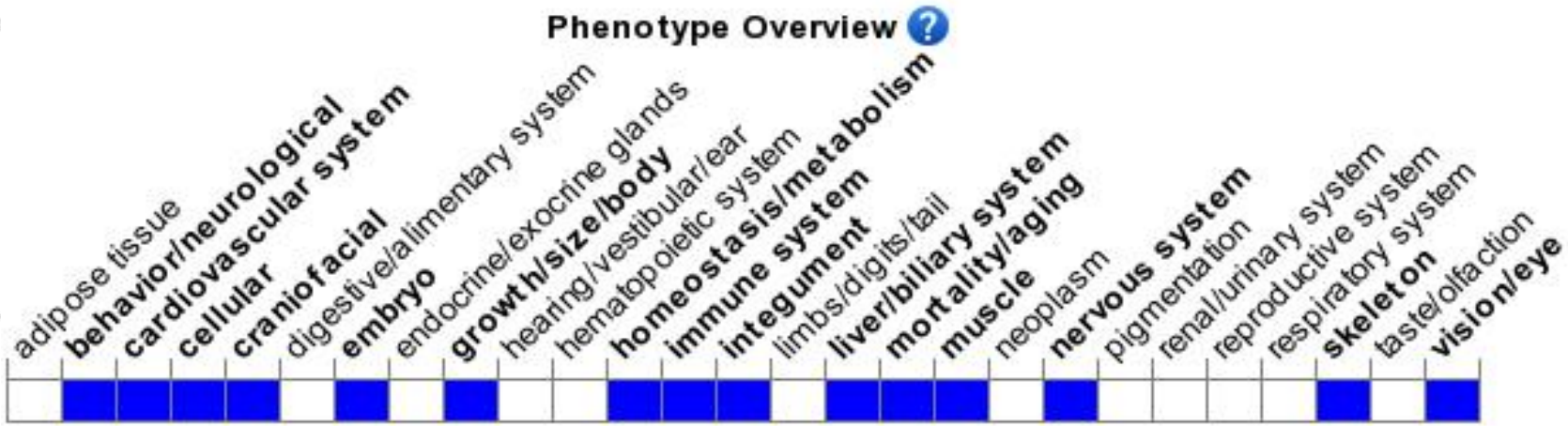
Genomic location distribution



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,Nullizygous mice show pre- and neonatal death, heart defects and hydrocephaly. Deletion of exon B1 disrupts migration of facial neurons, whereas deletion of exon B2 leads to Purkinje cell anomalies. Hypomorphs show hydrocephaly and defects in motor control, cerebellar foliation and neuron migration.

集萃药康
GemPharmatech

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

Tel: 400-9660890

集萃药康生物科技
GemPharmatech Co.,Ltd