

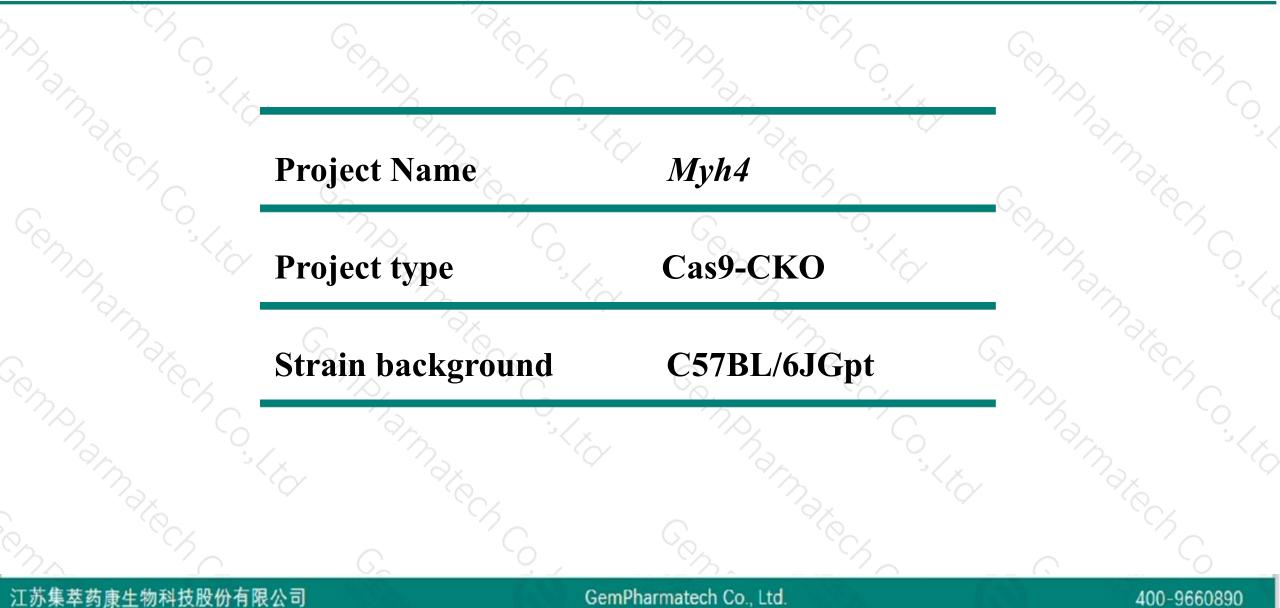
# Myh4 Cas9-CKO Strategy

Designer: Reviewer: Design Date: Yang Zeng Ruirui Zhang

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# **Project Overview**



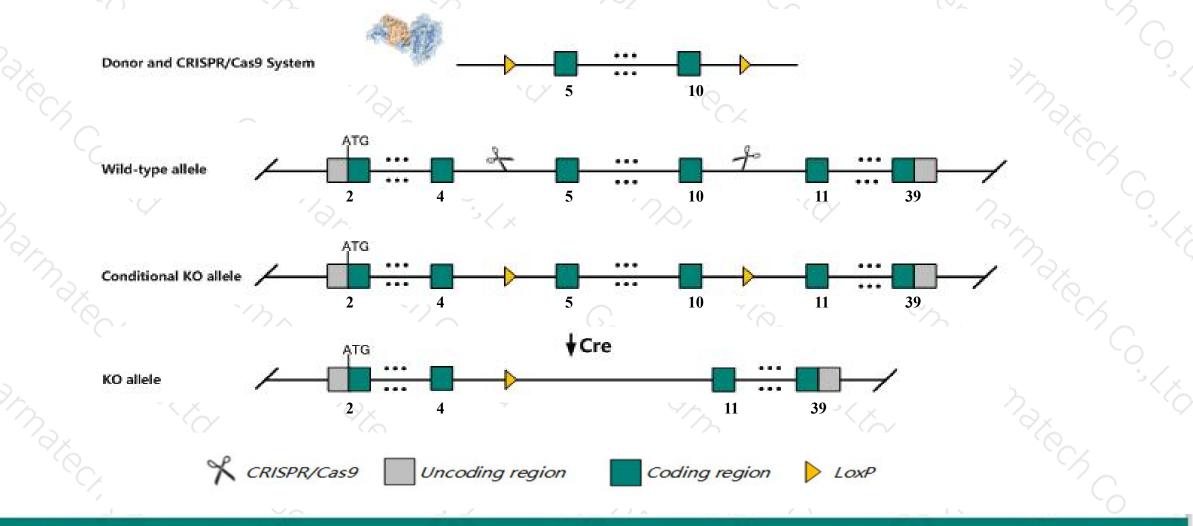


# **Conditional Knockout strategy**



400-9660890

This model will use CRISPR/Cas9 technology to edit the Myh4 gene. The schematic diagram is as follows:



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The Myh4 gene has 2 transcripts. According to the structure of Myh4 gene, exon5-exon10 of Myh4-201 (ENSMUST00000018632.10) transcript is recommended as the knockout region. The region contains 503bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Myh4* 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.



- According to the existing MGI data, Homozygotes for a targeted null mutation exhibit decreased growth and muscle defects including reduced muscle mass, muscle fiber loss, compensatory fiber hypertrophy, and impaired strength.
- The Myh4 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

# **Gene information (NCBI)**



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#### Myh4 myosin, heavy polypeptide 4, skeletal muscle [ Mus musculus (house mouse) ]

Gene ID: 17884, updated on 1-Oct-2019

#### Summary

Official SymbolWŋ44 provided by MGIOfficial Full Namemyosin, heavy polypeptide 4, skeletal muscle provided by MGIPrimary sourceMGI:MGI:1339713See relatedEnsembl:ENSMUSG0000057003Gene typeprotein codingGene typeprotein codingVALIDATEDMus musculusLineageEukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae;<br/>Murinae; Mus; MusAlso known asMM; MHC2B; Myhsf; MYH-2B; Minmus; Al506973; MyHC-IIb

Expression Restricted expression toward mammary gland adult (RPKM 118.7) See more

Orthologs human all

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# **Transcript information (Ensembl)**



The gene has 2 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Myh4-201	ENSMUST0000018632.10	6016	<u>1939aa</u>	Protein coding	CCDS24856	Q5SX39	TSL:5 GENCODE basic APPRIS P1
Myh4-202	ENSMUST00000170942.1	5912	<u>1939aa</u>	Protein coding	CCDS24856	Q5SX39	TSL:1 GENCODE basic APPRIS P1

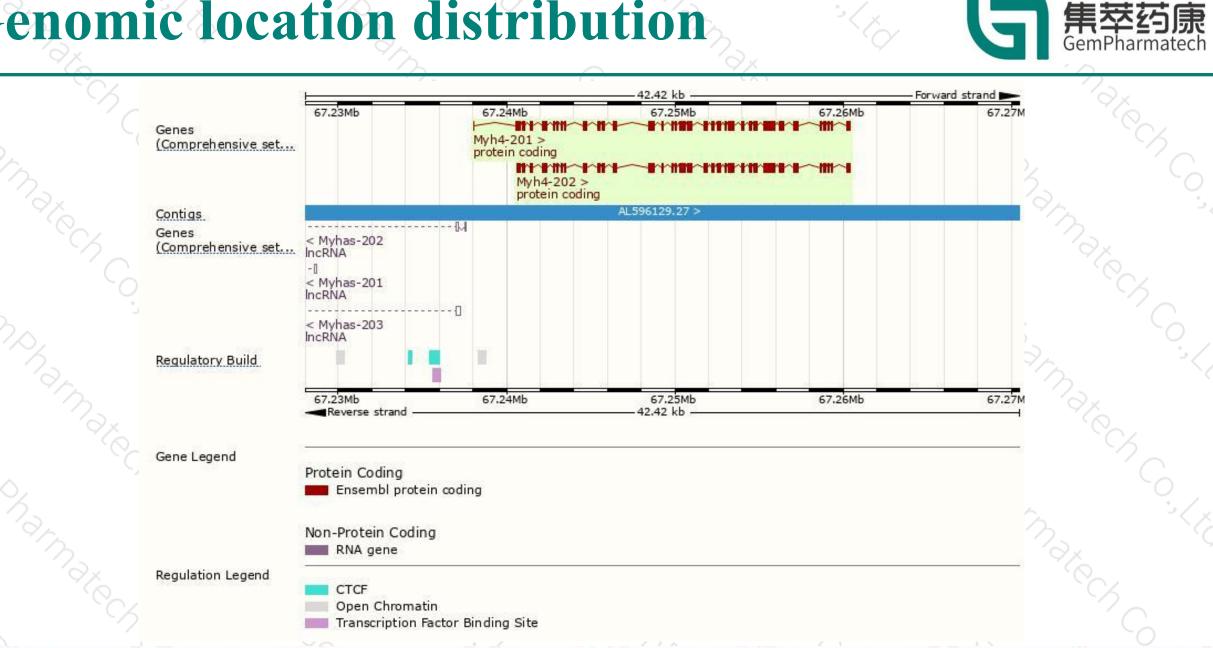
The strategy is based on the design of Myh4-201 transcript, The transcription is shown below

Myh4-201 > protein coding

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### **Genomic location distribution**



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# **Protein domain**



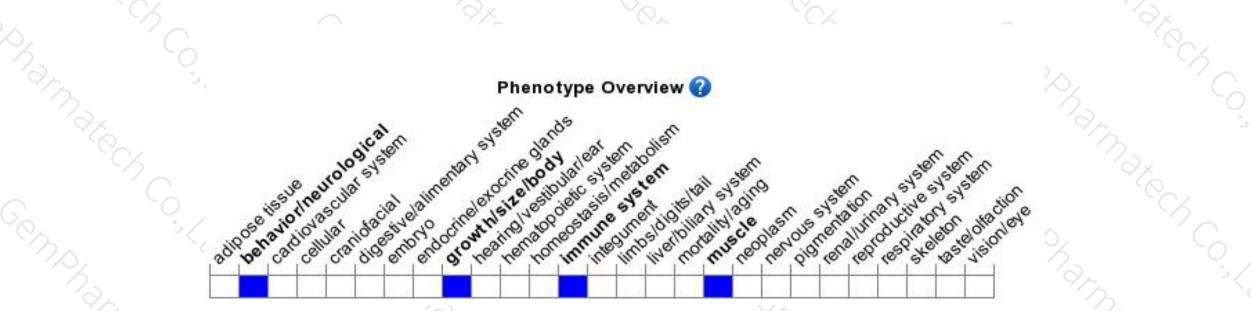
SK Co.	ENSMUSP00000018 MobiDB lite Low complexity (Seg) Coiled-coils (Ncoils) Superfamily	P-loop containing nucleoside triphosphate hydrolase SSF90257 SSF57997
	SMART	Myosin head, motor domain
50.	Prints	Myosin head, motor domain
	<u>Pfam</u>	Myosin head, motor domain Myosin head, motor domain Myosin head, motor domain Myosin, N-terminal, SH3-like
		Myosin, N-terminal, SH3-like
	PROSITE profiles	Myosin head, motor domain IQ motif, EF-hand binding site
	20.01200200	Myosin, N-terminal, SH3-like
	PANTHER	PTHR45615:SF3 7
	Gene3D	PTHR45615
	Geneso	1.20, 120, 720 3.30, 70, 3240 1.20, 5, 1050 1.20, 5, 370
3Kech		Myosin S1 fragment, N-terminal Myosin IQ motif-containing domain superfamily
		Kinesin motor domain superfamily
	CDD.	cd14915
	All sequence SNPs/i	Sequence variants (dbSNP and all other sources)
	Variant Legend	
	150	missense variant splice region variant
		synonymous variant
	Scale bar	0 200 400 600 800 1000 1200 1400 1600 1939
1		

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### 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 a targeted null mutation exhibit decreased growth and muscle defects including reduced muscle mass, muscle fiber loss, compensatory fiber hypertrophy, and impaired strength.



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



