

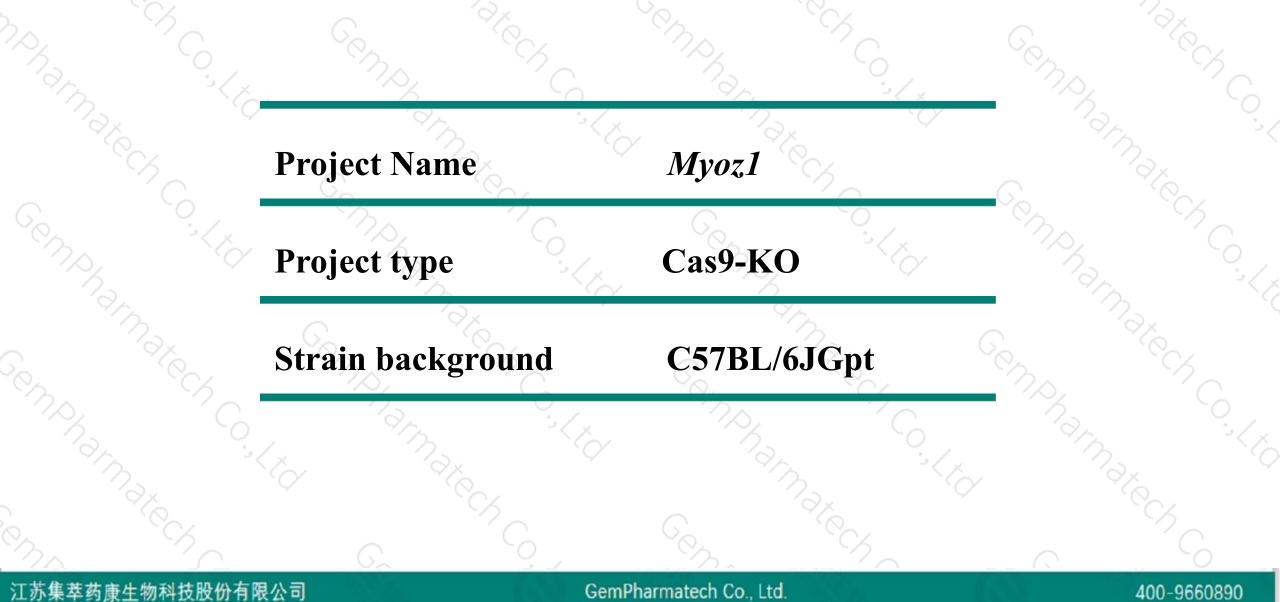
# Myoz1 Cas9-KO Strategy

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Designer: Xueting Zhang Reviewer:Yanhua Shen Date:2020-03-04

# **Project Overview**

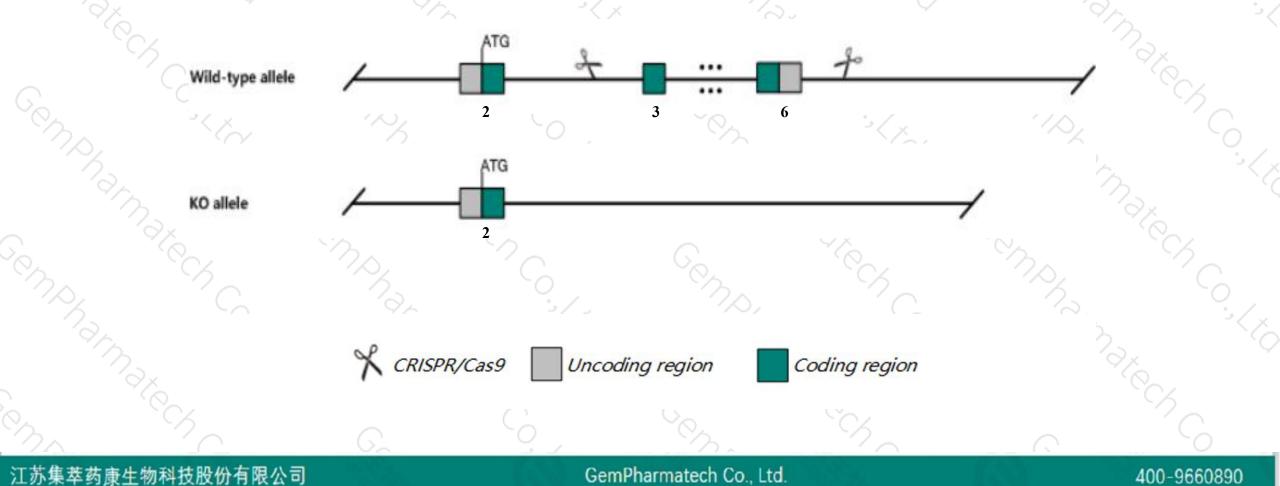




# **Knockout** strategy



This model will use CRISPR/Cas9 technology to edit the Myoz1 gene. The schematic diagram is as follows:





- The Myoz1 gene has 4 transcripts. According to the structure of Myoz1 gene, exon3-exon6 of Myoz1-201 (ENSMUST00000090469.7) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Myoz1 gene. The brief process is as follows: CRISPR/Cas9 system we



- According to the existing MGI data, Mice homozygous for a knock-out allele show reduced body weight and fast-twitch muscle mass, a fiber type shift toward more oxidative fibers, increased exercise capacity and calcineurin activity, and enhanced muscle regeneration after cardiotoxin injury.
- The Myoz1 gene is located on the Chr14. 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)



#### Myoz1 myozenin 1 [ Mus musculus (house mouse) ]

Gene ID: 59011, updated on 12-Aug-2019

Summary

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Official Symbol	Myoz1 provided by MGI
Official Full Name	myozenin 1 provided by MGI
Primary source	MGI:MGI:1929471
See related	Ensembl:ENSMUSG0000068697
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	FATZ; Myoz; AV090278; 2310001N11Rik
Expression	Biased expression in mammary gland adult (RPKM 65.7) and lung adult (RPKM 3.9) See more
Orthologs	human all

#### Genomic context

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See Myoz1 in Genome Data Viewer

Location: 14; 14 A3

Exon count: 6

 Annotation release
 Status
 Assembly
 Chr
 Location

 108
 current
 GRCm38.p6 (GCF\_000001635.26)
 14
 NC\_000080.6 (20649102..20656540, complement)

 Build 37.2
 previous assembly
 MGSCv37 (GCF\_000001635.18)
 14
 NC\_000080.5 (21468324..21475762, complement)

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# **Transcript information (Ensembl)**



# The gene has 4 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Myoz1-201	ENSMUST0000090469.7	1289	<u>296aa</u>	Protein coding	CCDS36819	<u>Q9JK37</u>	TSL:1 GENCODE basic APPRIS P1
Myoz1-202	ENSMUST00000224436.1	607	No protein	Retained intron	-	-8	
Myoz1-203	ENSMUST00000224472.1	405	No protein	Retained intron	-	28	
Myoz1-204	ENSMUST00000225231.1	346	No protein	IncRNA	22	20	

The strategy is based on the design of Myoz1-201 transcript, The transcription is shown below

#### < Myoz1-201 protein coding

Reverse strand

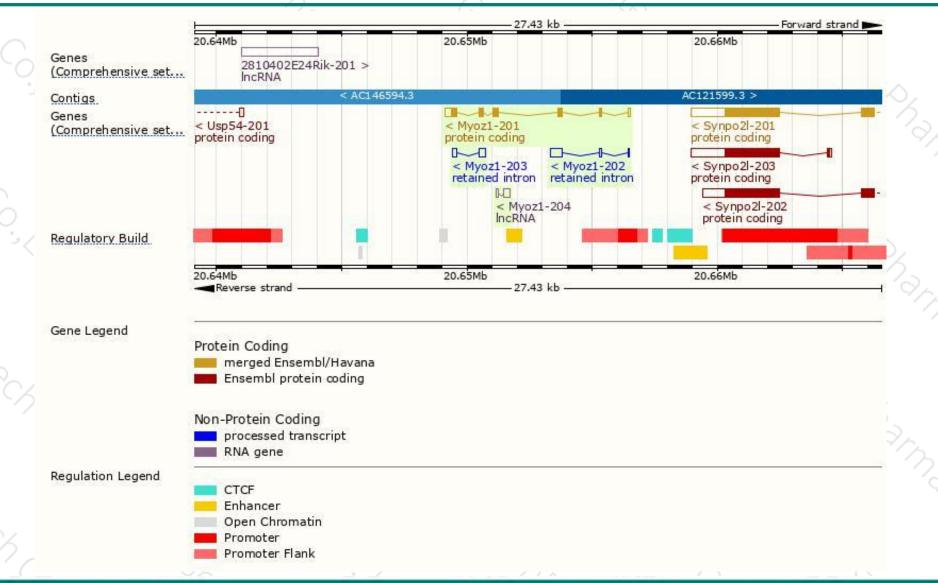
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7.43 kb

# **Genomic location distribution**





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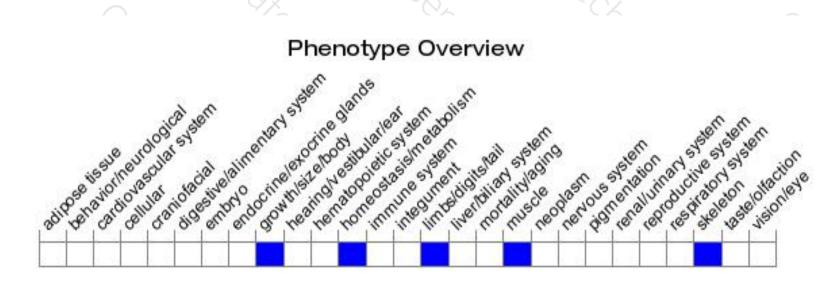
# **Protein domain**



130		S. S		26	10/3/	3		C M B H	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
60	ENSMUSP00000087 MobiDB lite Low complexity (Seg) Pfam PANTHER	Myozenin							
6	All sequence SNPs/i Variant Legend	PTHR15941:SF11 MyozenIn Sequence varia missense	ints (dbSNP an	d all other sourc	es)	Ť	111	1	
SNX	Scale bar		us variant 40	80	120	160	200	240	296
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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, Mice homozygous for a knock-out allele show reduced body weight and fast-twitch muscle mass, a fiber type shift toward more oxidative fibers, increased exercise capacity and calcineurin activity, and enhanced muscle regeneration after cardiotoxin injury.

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



