

Myl1 Cas9-KO Strategy

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Project Overview



Project Name

Myl1

Project type

Cas9-KO

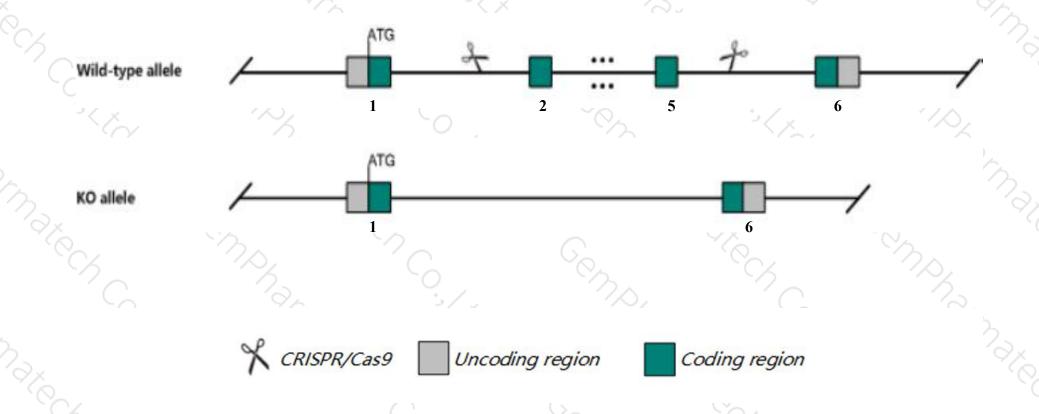
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Myl1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Myl1* gene has 8 transcripts. According to the structure of *Myl1* gene, exon2-exon5 of *Myl1-201*(ENSMUST00000027151.11) transcript is recommended as the knockout region. The region contains 424bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Myl1* gene. The brief process is as follows: CRISPR/Cas9 system 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.

Notice



- > According to the existing MGI data, homozygotes for a targeted null mutation exhibit developmental delay, fail to form mesoderm, and die by embryonic day 8.5.
- The *Myl1* 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)



Myl1 myosin, light polypeptide 1 [Mus musculus (house mouse)]

Gene ID: 17901, updated on 9-Jun-2020

Summary

☆ ?

Official Symbol Myl1 provided by MGI

Official Full Name myosin, light polypeptide 1 provided by MGI

Primary source MGI:MGI:97269

See related Ensembl: ENSMUSG00000061816

Gene type protein coding
RefSeq status REVIEWED
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae;

Mus; Mus

Also known as Mylf; MLC1f; MLC3f; Al325107

Summary Myosin is a hexameric ATPase cellular motor protein. It is composed of two heavy chains, two non-phosphorylatable alkali light chains, and two phosphorylatable

regulatory light chains. This gene encodes a myosin alkali light chain expressed in fast skeletal muscle. Multiple transcript variants encoding different isoforms have

been identified for this gene. [provided by RefSeq, Jul 2008]

Expression Biased expression in mammary gland adult (RPKM 118.5), limb E14.5 (RPKM 25.3) and 3 other tissues See more

Orthologs human all

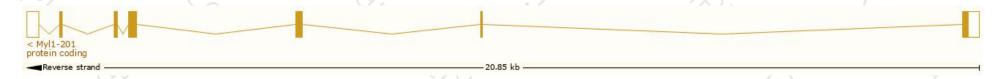
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

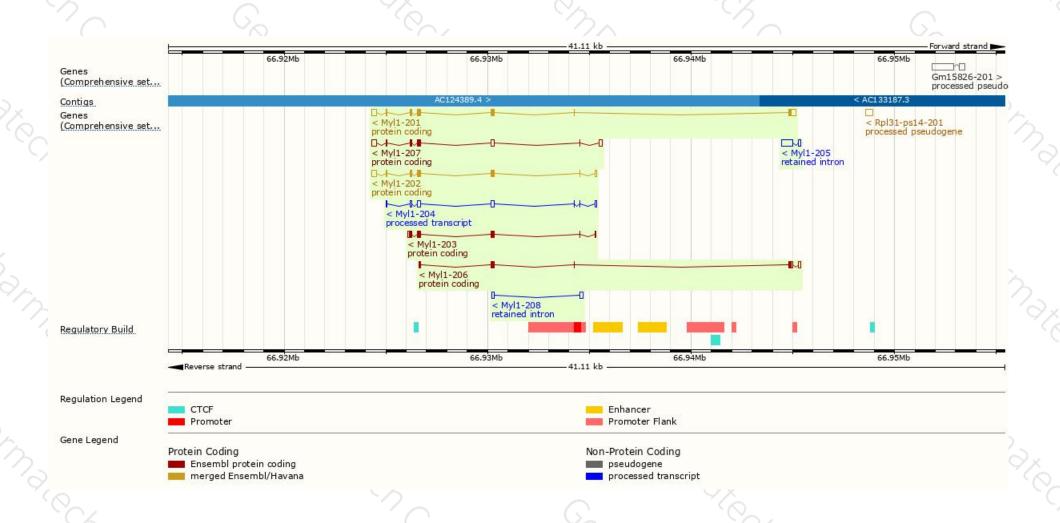
Name 🍦	Transcript ID	bp 🛊	Protein	Biotype	CCDS	UniProt	Flags
Myl1-201	ENSMUST00000027151.11	1082	<u>188aa</u>	Protein coding	CCDS15024 €	P05977@Q545T7@	TSL:1 GENCODE basic
MyI1-202	ENSMUST00000119429.7	803	<u>150aa</u>	Protein coding	CCDS48284 ₽	P05977@Q545G5@	TSL:1 GENCODE basic APPRIS P1
MyI1-207	ENSMUST00000186202.6	874	<u>92aa</u>	Protein coding	-	A0A087WRZ7₺	TSL:5 GENCODE basic
Myl1-203	ENSMUST00000120415.7	621	<u>161aa</u>	Protein coding	-	E9PWG4₽	TSL:1 GENCODE basic
MyI1-206	ENSMUST00000160100.1	571	<u>124aa</u>	Protein coding	2	E0CZ30@	CDS 3' incomplete TSL:3
MyI1-204	ENSMUST00000150542.1	567	No protein	Processed transcript	20	1.5	TSL:5
Myl1-205	ENSMUST00000151328.2	662	No protein	Retained intron			TSL:2
MyI1-208	ENSMUST00000186346.1	319	No protein	Retained intron	1.50	-	TSL:3

The strategy is based on the design of *Myl1-201* transcript, the transcription is shown below:



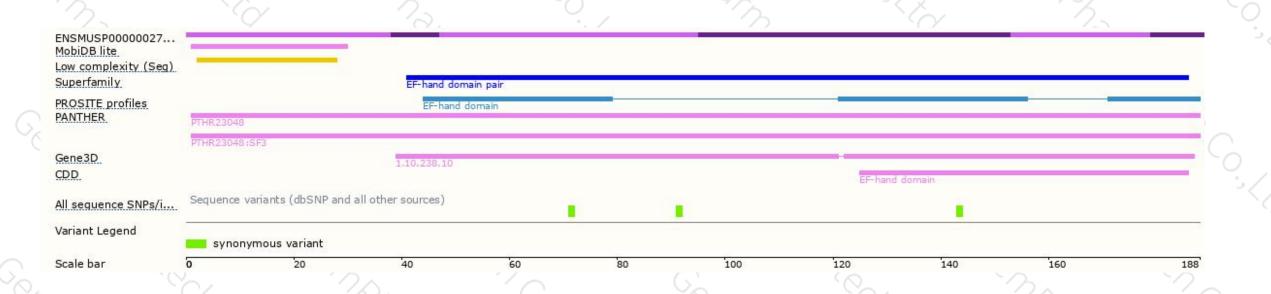
Genomic location distribution





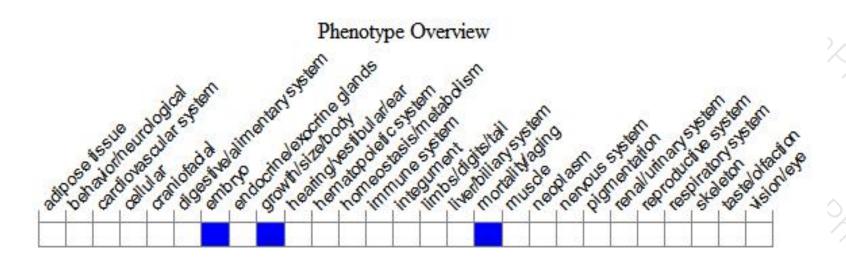
Protein domain





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. Tel: 400-9660890





