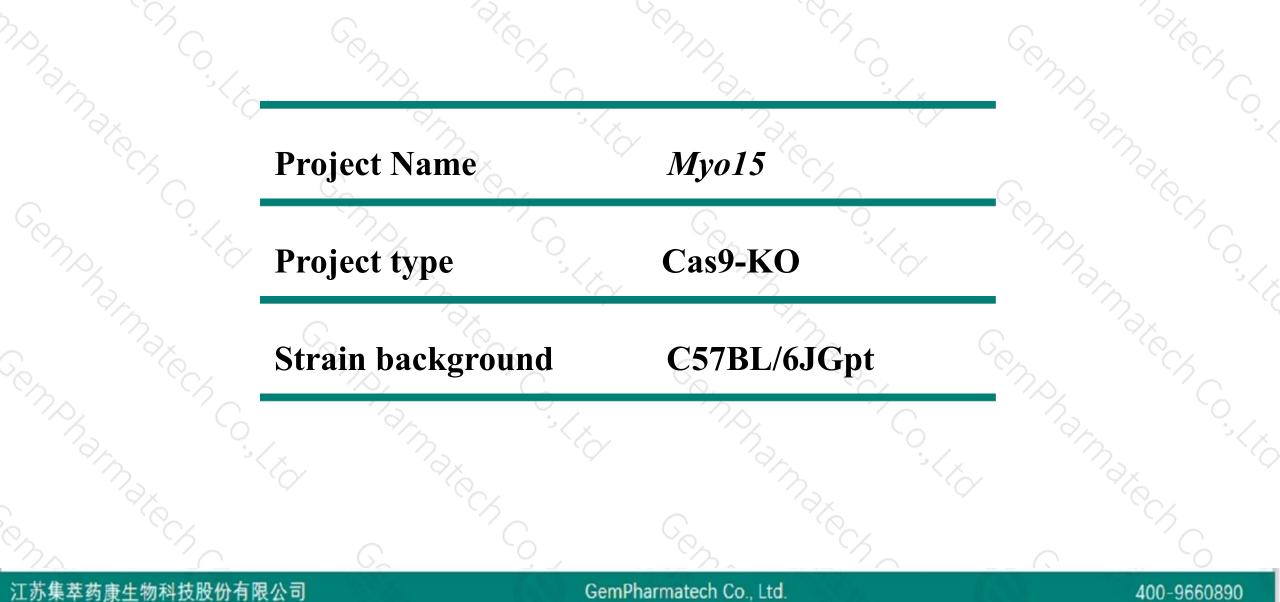


Myo15 Cas9-KO Strategy

Designer: Reviewer: Design Date: Huimin Su Ruirui Zhang 2020/2/10

Project Overview

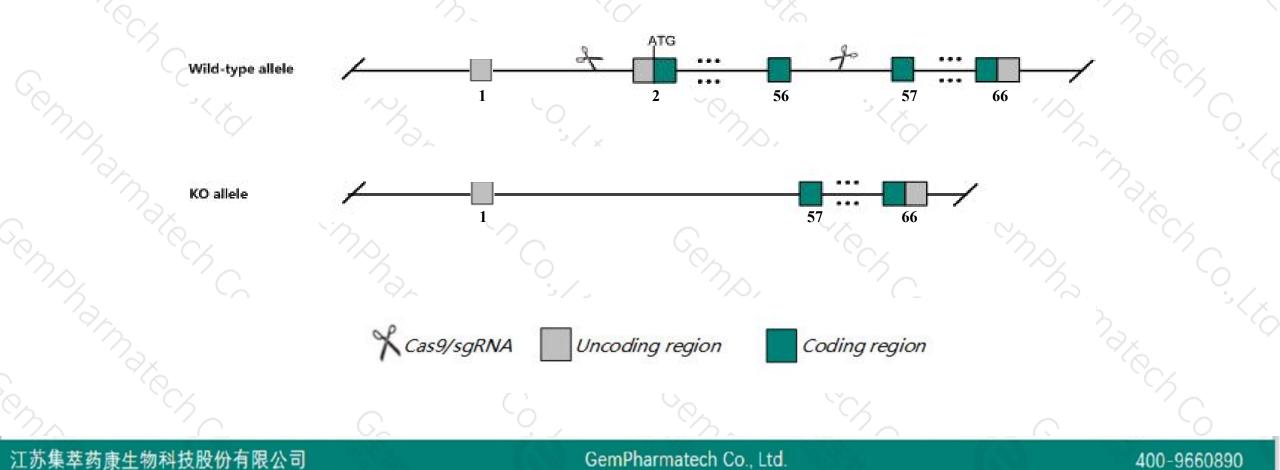




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Myo15 gene. The schematic diagram is as follows:





The Myo15 gene has 6 transcripts. According to the structure of Myo15 gene, exon2-exon56 of Myo15-201 (ENSMUST00000071880.8) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.

> In this project we use CRISPR/Cas9 technology to modify Myo15 gene. The brief process is as follows: CRISPR/Cas9 system



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- According to the existing MGI data, Mutations in this gene result in profound deafness and neurological behavior.
- The Myo15 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.

Notice

Gene information (NCBI)



Myo15 myosin XV [Mus musculus (house mouse)]

Gene ID: 17910, updated on 14-Aug-2019

- Summary

☆ ?

Official Symbol	Myo15 provided by MGI
Official Full Name	myosin XV provided by MGI
Primary source	MGI:MGI:1261811
See related	Ensembl:ENSMUSG0000042678
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	sh2; sh-2; Myo15a
Expression	Biased expression in testis adult (RPKM 2.4), cerebellum adult (RPKM 0.5) and 4 other tissues See more

Orthologs human all

- Genomic context

* ?

Location: 11 B2; 11 37.81 cM

See Myo15 in Genome Data Viewer

Exon count: 67

Ann	notation release	Status	Assembly	Chr	Location
108	1	current	GRCm38.p6 (GCF_000001635.26)	11	NC_000077.6 (6046933960528369)
Buil	ld 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	11	NC_000077.5 (6028284160341871)



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Transcript information (Ensembl)



Forward strand

The gene has 6 transcripts, all transcripts are shown below:

						· · · · · · · · · · · · · · · · · · ·		
Name 🖕	Transcript ID	bp 💧 Protein	Protein 💧	Biotype	CCDS	UniProt	Flags	
Myo15-201	ENSMUST0000071880.8	11769	<u>3511aa</u>	Protein coding	CCDS24792	<u>Q9QZZ4</u> @	TSL:5 GENCODE basic APPRIS P4	
Myo15-203	ENSMUST0000094135.8	11715	<u>3493aa</u>	Protein coding	CCDS48811@	Q9QZZ4&	TSL:5 GENCODE basic APPRIS ALT2	
Myo15-202	ENSMUST0000081823.11	7953	<u>2306aa</u>	Protein coding	<u>CCDS24793</u> &	<u>Q9QZZ4</u> &	TSL:5 GENCODE basic APPRIS ALT2	
Myo15-205	ENSMUST00000126522.3	5351	<u>1784aa</u>	Protein coding	-	F6VXK7团	CDS 5' and 3' incomplete TSL:1	
Myo15-206	ENSMUST00000238401.1	360	<u>77aa</u>	Protein coding	22	123	CDS 3' incomplete	
Myo15-204	ENSMUST00000122825.1	2285	No protein	Retained intron	5.	170	TSL:1	
		-						

The strategy is based on the design of Myo15-201 transcript, The transcription is shown below

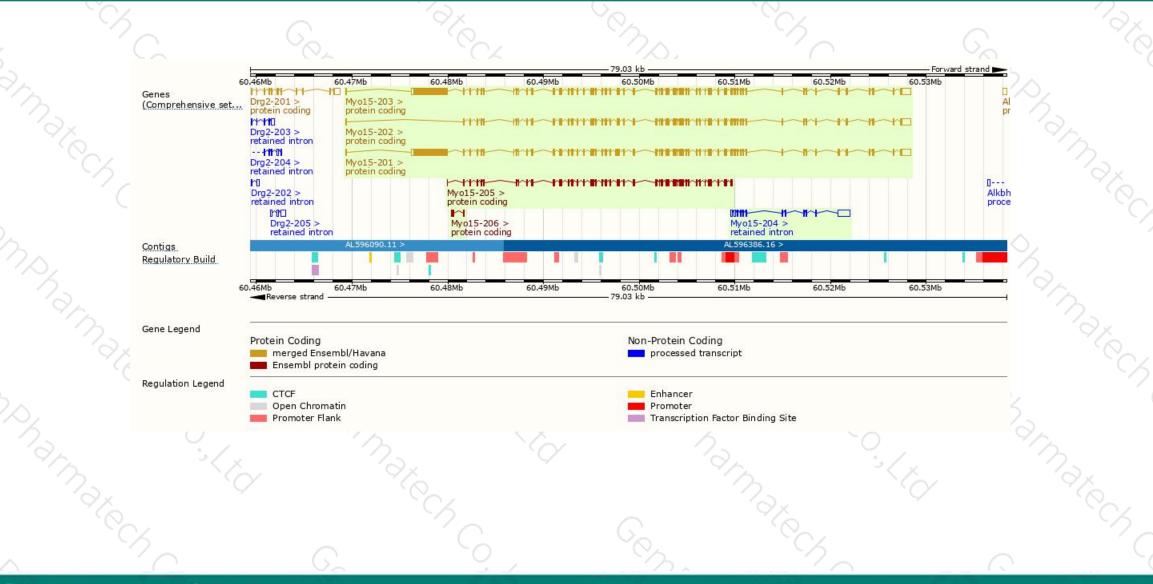
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Myo15-201 >

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Genomic location distribution



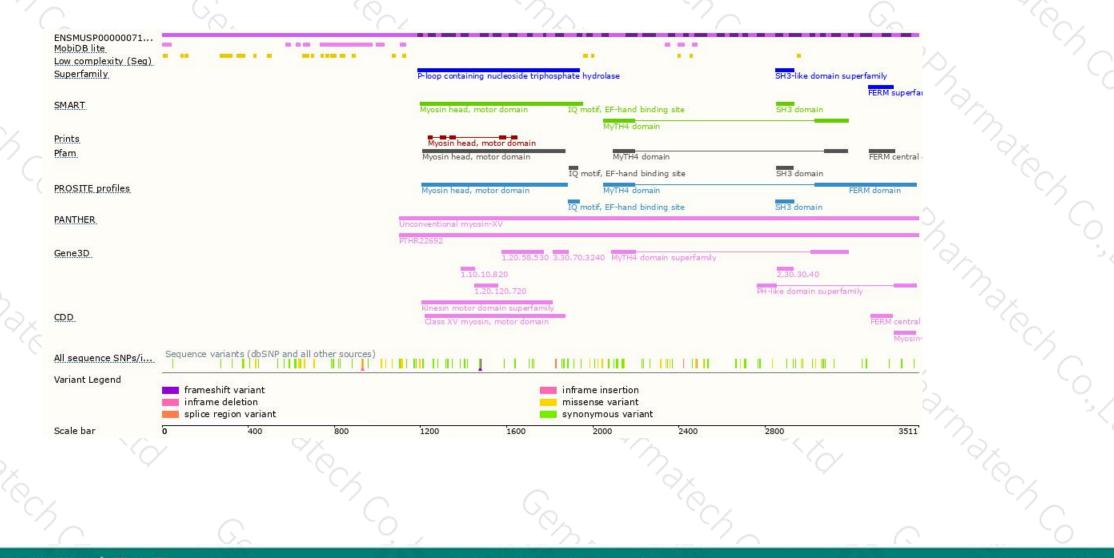


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Protein domain



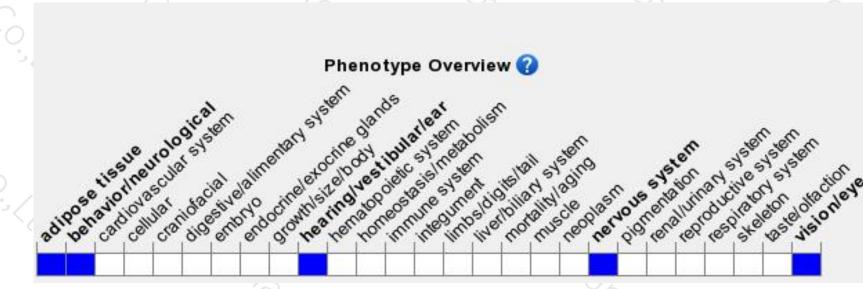


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Mouse phenotype description(MGI)





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, Mutations in this gene result in profound deafness and neurological behavior.



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



