

Lhx9 Cas9-KO Strategy

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Project Overview



Project Name

Lhx9

Project type

Cas9-KO

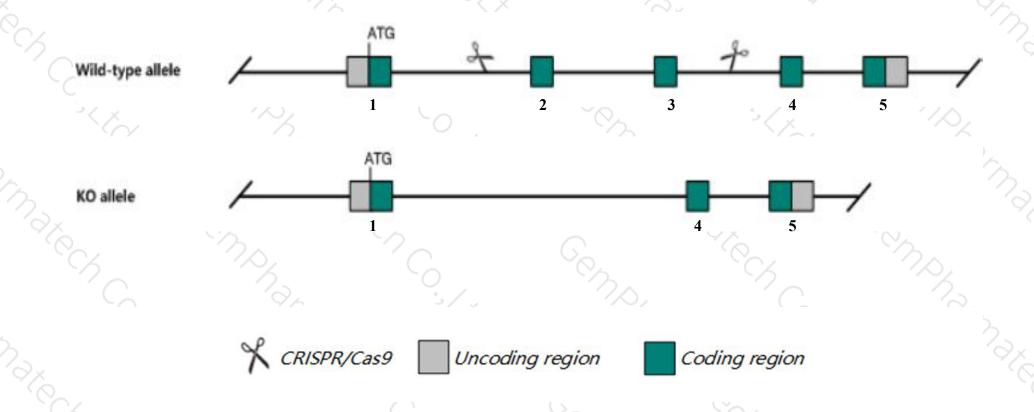
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Lhx9* gene. The schematic diagram is as follows:



Technical routes



- > The *Lhx9* gene has 7 transcripts. According to the structure of *Lhx9* gene, exon2-exon3 of *Lhx9*-204(ENSMUST00000112026.3) transcript is recommended as the knockout region. The region contains 559bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Lhx9* 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 failed proliferation of the somatic cells of the genital ridge resulting in lack of discrete gonad formation, infertility in both sexes, and female-like genitalia in genetically male mice.
- ightharpoonup Transcript *Lhx9*-206 may not be affected.
- \Rightarrow The *Lhx9* 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)



Lhx9 LIM homeobox protein 9 [Mus musculus (house mouse)]

Gene ID: 16876, updated on 13-Mar-2020





Official Symbol Lhx9 provided by MGI

Official Full Name LIM homeobox protein 9 provided byMGI

Primary source MGI:MGI:1316721

See related Ensembl: ENSMUSG00000019230

Gene type protein coding

RefSeq status VALIDATED

Organism <u>Mus musculus</u>

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

Muroidea; Muridae; Murinae; Mus; Mus

Also known as 3110009007Rik, BB104635, LH2B

Expression Biased expression in whole brain E14.5 (RPKM 7.8), CNS E14 (RPKM 6.5) and 4 other tissuesSee more

Orthologs <u>human</u> all

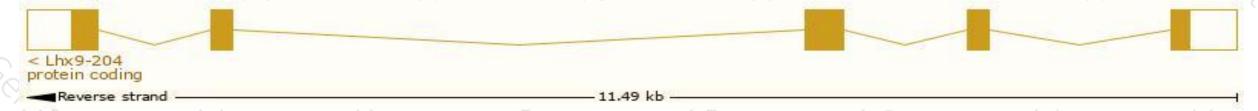
Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

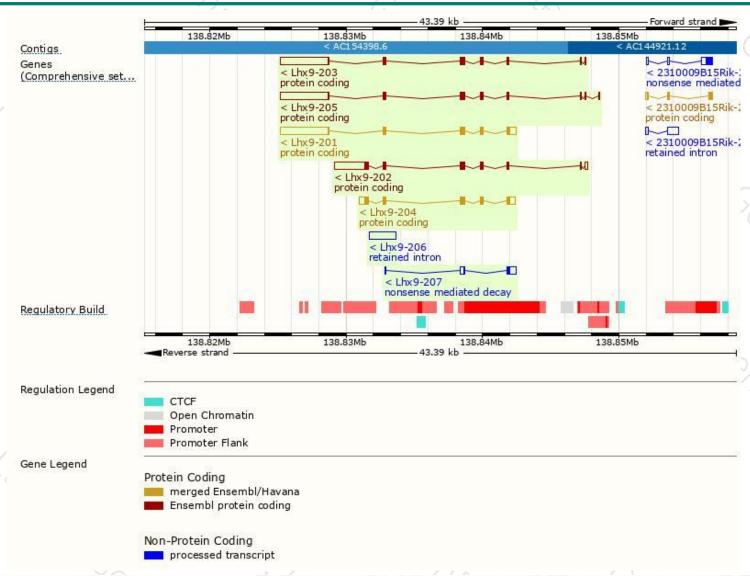
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Lhx9-201	ENSMUST00000019374.13	4937	330aa	Protein coding	CCDS15334	Q9WUH2	TSL:1 GENCODE basic APPRIS P3
Lhx9-205	ENSMUST00000112030.8	4633	<u>321aa</u>	Protein coding	CCDS15333	Q9WUH2	TSL:5 GENCODE basic
Lhx9-203	ENSMUST00000093486.9	4551	<u>321aa</u>	Protein coding	CCDS15333	Q9WUH2	TSL:1 GENCODE basic
Lhx9-204	ENSMUST00000112026.3	2065	<u>397aa</u>	Protein coding	CCDS35726	Q9WUH2	TSL:1 GENCODE basic APPRIS ALT1
Lhx9-202	ENSMUST00000046870.12	3645	388aa	Protein coding	-	Q9WUH2	TSL:5 GENCODE basic APPRIS ALT1
Lhx9-207	ENSMUST00000194557.1	1058	<u>75aa</u>	Nonsense mediated decay	170	A0A0A6YY42	TSL:5
Lhx9-206	ENSMUST00000192425.1	1946	No protein	Retained intron	-	-	TSL:NA

The strategy is based on the design of Lhx9-204 transcript, the transcription is shown below:



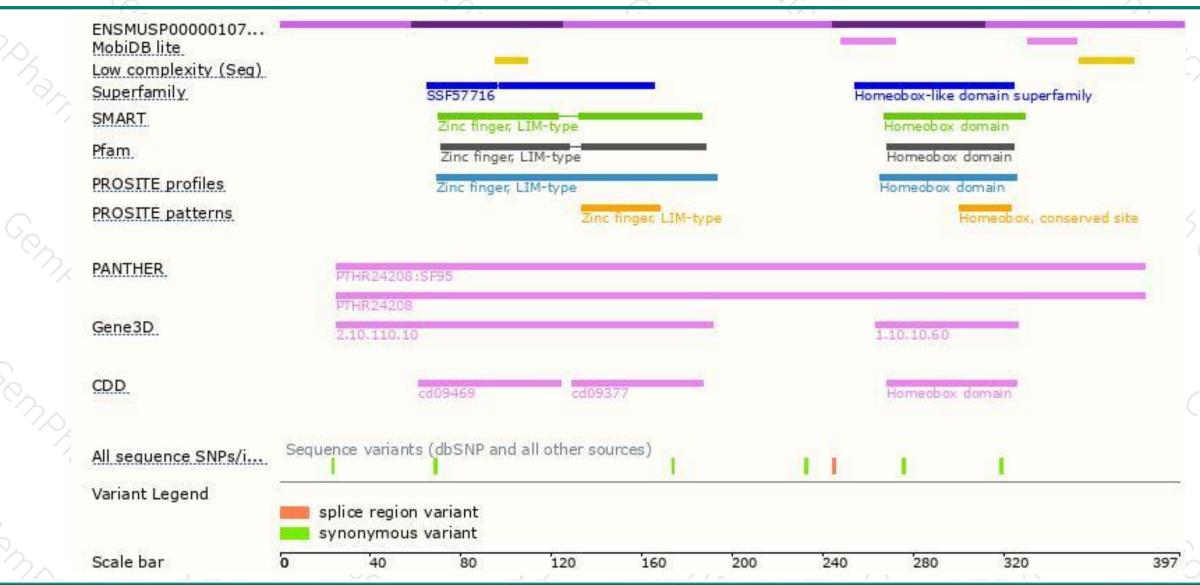
Genomic location distribution





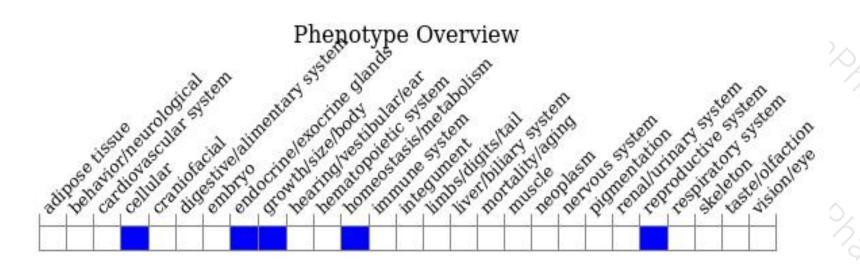
Protein domain





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, homozygotes for a targeted null mutation exhibit failed proliferation of the somatic cells of the genital ridge resulting in lack of discrete gonad formation, infertility in both sexes, and female-like genitalia in genetically male mice.



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





