

***Nhlh1* Cas9-CKO Strategy**

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Project Overview

Project Name

Nhlh1

Project type

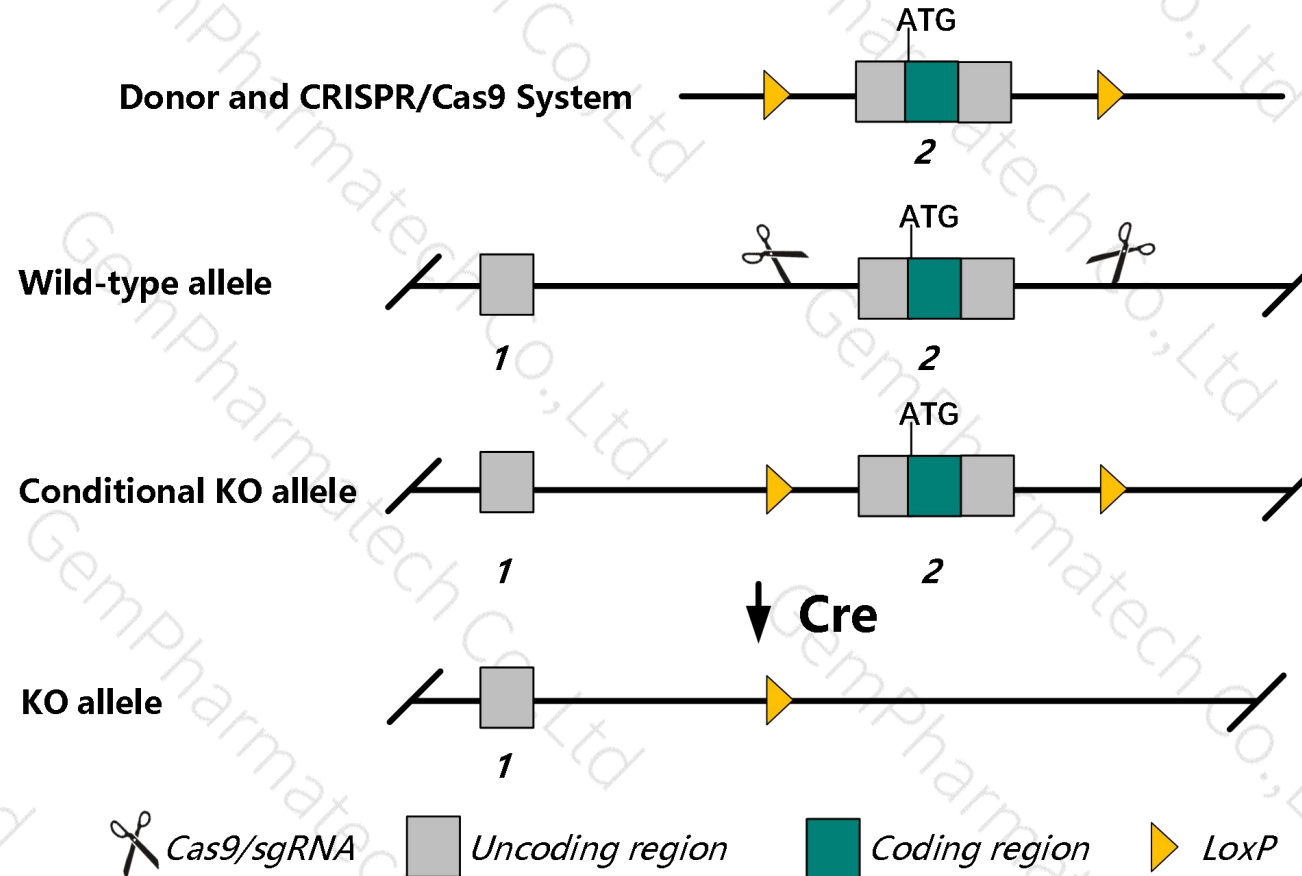
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Nhlh1* gene. The schematic diagram is as follows:



Technical routes

- The *Nhlh1* gene has 1 transcript. According to the structure of *Nhlh1* gene, exon2 of *Nhlh1*-201(ENSMUST00000059794.3) transcript is recommended as the knockout region. The region contains all 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 *Nhlh1* 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, homozygous targeted null mutants show cardiac arrhythmia and lack a heart rate response to water immersion. Adults are subject to sudden death, probably due to poor parasympathetic tone. Homozygotes for another targeted mutation had no abnormalities.
- The *Nhlh1* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)

Nhlh1 nescient helix loop helix 1 [Mus musculus (house mouse)]

Gene ID: 18071, updated on 13-Mar-2020

Summary

Official Symbol Nhlh1 provided by [MGI](#)

Official Full Name nescient helix loop helix 1 provided by [MGI](#)

Primary source [MGI:MGI:98481](#)

See related [Ensembl:ENSMUSG00000051251](#)

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 Hen1, Nscl, Tal2, bHLHa35

Expression Biased expression in CNS E11.5 (RPKM 11.7), CNS E14 (RPKM 7.5) and 2 other tissues [See more](#)

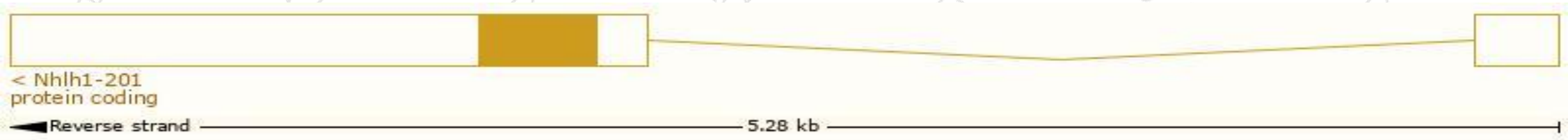
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

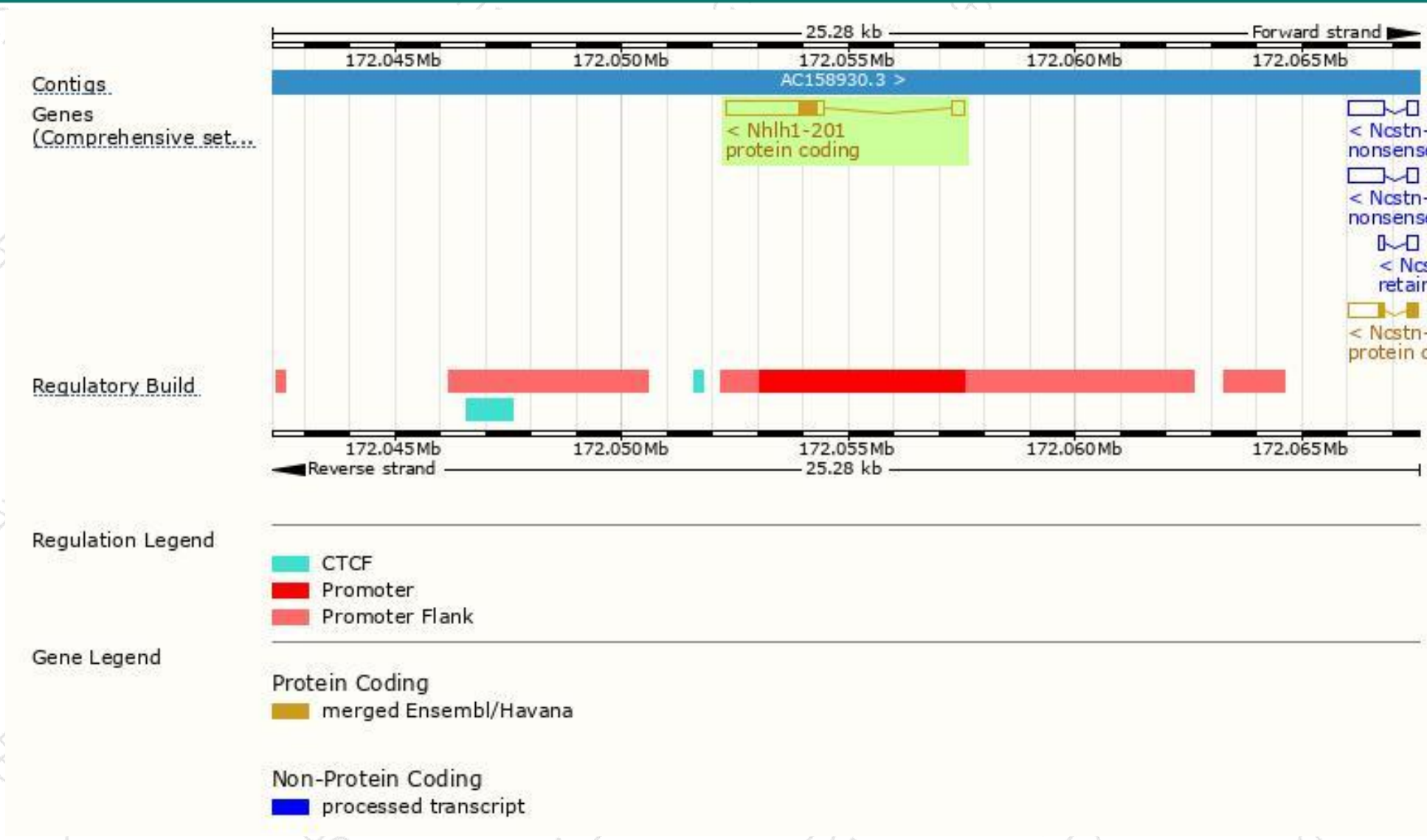
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Nhlh1-201	ENSMUST00000059794.3	2464	133aa	Protein coding	CCDS15506	Q02576	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1

The strategy is based on the design of *Nhlh1-201* transcript, the transcription is shown below:



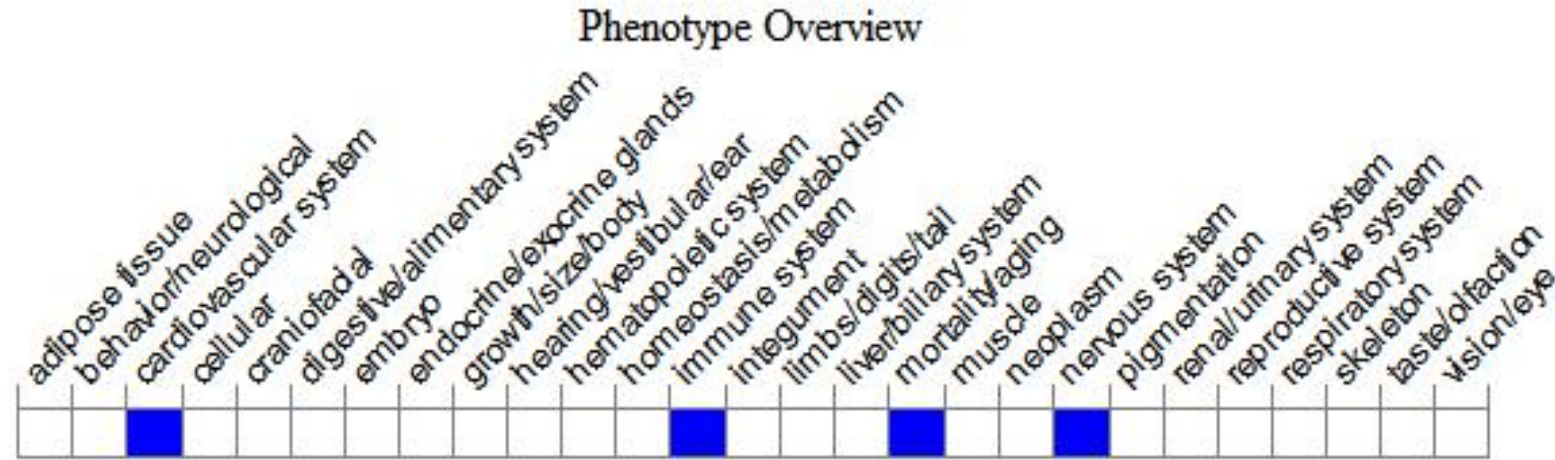
Genomic location distribution



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/>).

According to the existing MGI data, homozygous targeted null mutants show cardiac arrhythmia and lack a heart rate response to water immersion. Adults are subject to sudden death, probably due to poor parasympathetic tone.

Homozygotes for another targeted mutation had no abnormalities.

If you have any questions, you are welcome to inquire.

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