

Neurod2 Cas9-KO Strategy

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Project Overview



Project Name

Neurod2

Project type

Cas9-KO

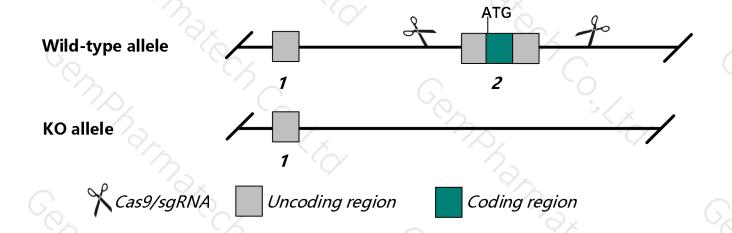
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Neurod2 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Neurod2* gene has 1 transcript. According to the structure of *Neurod2* gene, exon2 of *Neurod2*201(ENSMUST00000041685.6) 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 *Neurod2* 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 show elevated neuronal apoptosis resulting in ataxia, incoordination, elevated seizure susceptibility, and death by 35 days. Heterozygotes are moderately affected, and adults are subject to unexplained death.
- > The Neurod2 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.

Gene information (NCBI)



Neurod2 neurogenic differentiation 2 [Mus musculus (house mouse)]

Gene ID: 18013, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Neurod2 provided by MGI

Official Full Name neurogenic differentiation 2 provided by MGI

Primary source MGI:MGI:107755

See related Ensembl: ENSMUSG00000038255

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 Ndrf, bHLHa1

Expression Biased expression in CNS E14 (RPKM 26.6), whole brain E14.5 (RPKM 24.4) and 4 other tissues See more

Orthologs human all

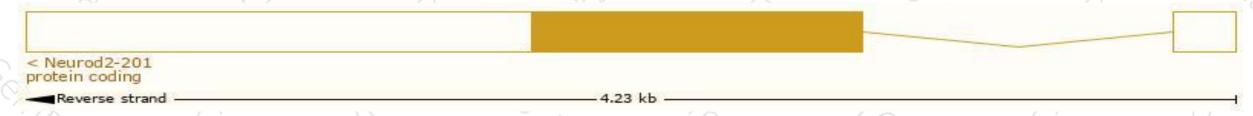
Transcript information (Ensembl)



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

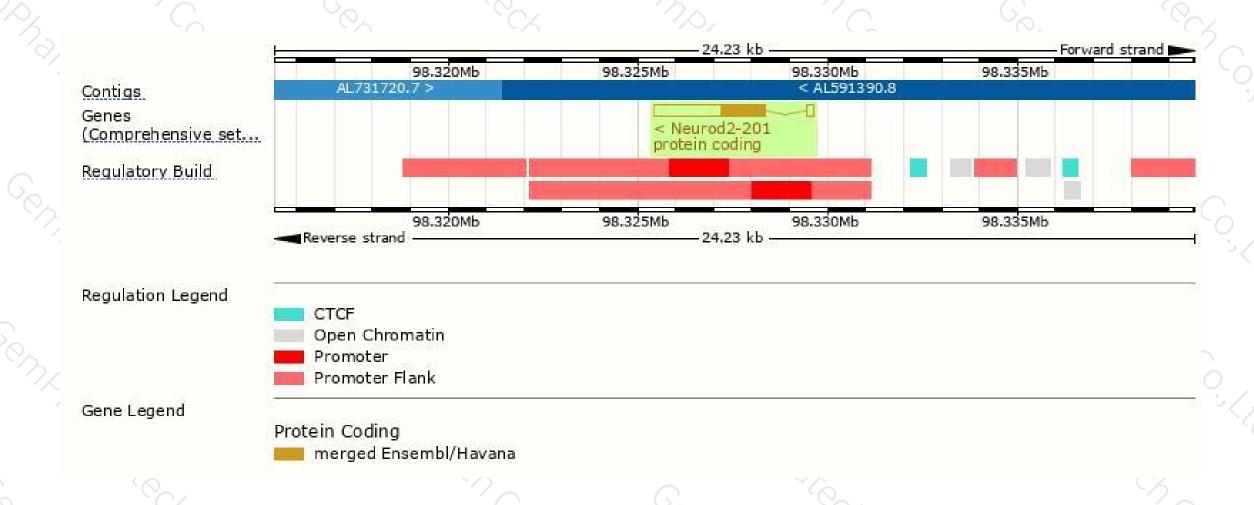
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Neurod2-201	ENSMUST00000041685.6	3142	<u>383aa</u>	Protein coding	CCDS25343	Q3TYB2 Q62414	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1

The strategy is based on the design of *Neurod2-201* transcript, the transcription is shown below:



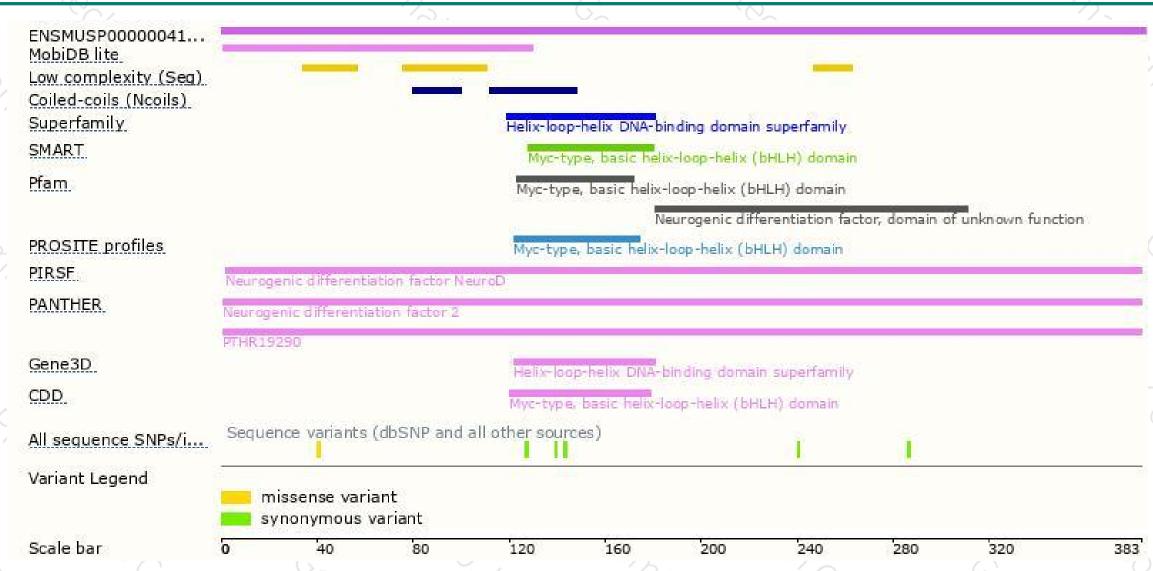
Genomic location distribution





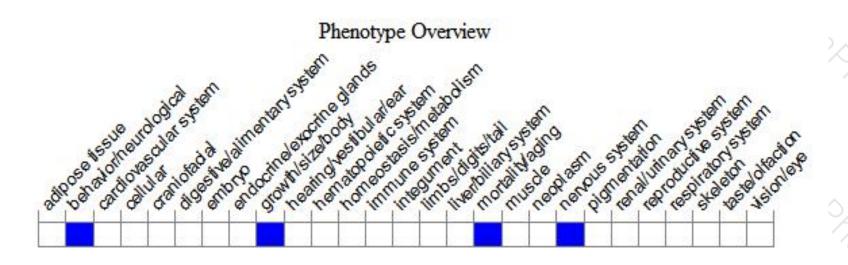
Protein domain





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, homozygotes for a targeted null mutation show elevated neuronal apoptosis resulting in ataxia, incoordination, elevated seizure susceptibility, and death by 35 days. Heterozygotes are moderately affected, and adults are subject to unexplained death.



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





