

***Neurod1* Cas9-KO Strategy**

Designer: XiaoJing Li
Design Date:2019-11-27
Reviewer:JiaYu

Project Overview

Project Name

Neurod1

Project type

Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Neurod1* gene. The schematic diagram is as follows:



- The *Neurod1* gene has 1 transcript. According to the structure of *Neurod1* gene, exon2 of *Neurod1-201* (ENSMUST00000041099.4) 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 *Neurod1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygotes for targeted null mutations exhibit neonatal diabetes, pancreatic enteroendocrine cell deficits, impaired hearing and balance, retinal degeneration, and seizures. Survival past birth is dependent on genetic background.
- The knockout region overlaps with *Cerkl-210 lncrna* gene, which may affect *cerkl-210 lncrna* after knockout.
- The *Neurod1* gene is located on the Chr2. 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)

Neurod1 neurogenic differentiation 1 [*Mus musculus* (house mouse)]

Gene ID: 18012, updated on 26-Nov-2019

Summary

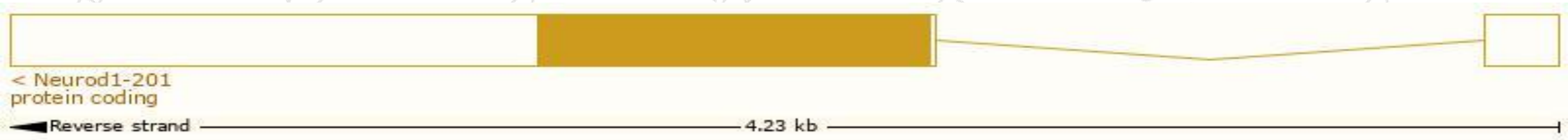
| | |
|--------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Official Symbol | Neurod1 provided by MGI |
| Official Full Name | neurogenic differentiation 1 provided by MGI |
| Primary source | MGI:MGI:1339708 |
| See related | Ensembl:ENSMUSG00000034701 |
| 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 | Nd1; BETA2; BHF-1; Neurod; bHLHa3 |
| Expression | Biased expression in cerebellum adult (RPKM 75.5), whole brain E14.5 (RPKM 12.5) and 3 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

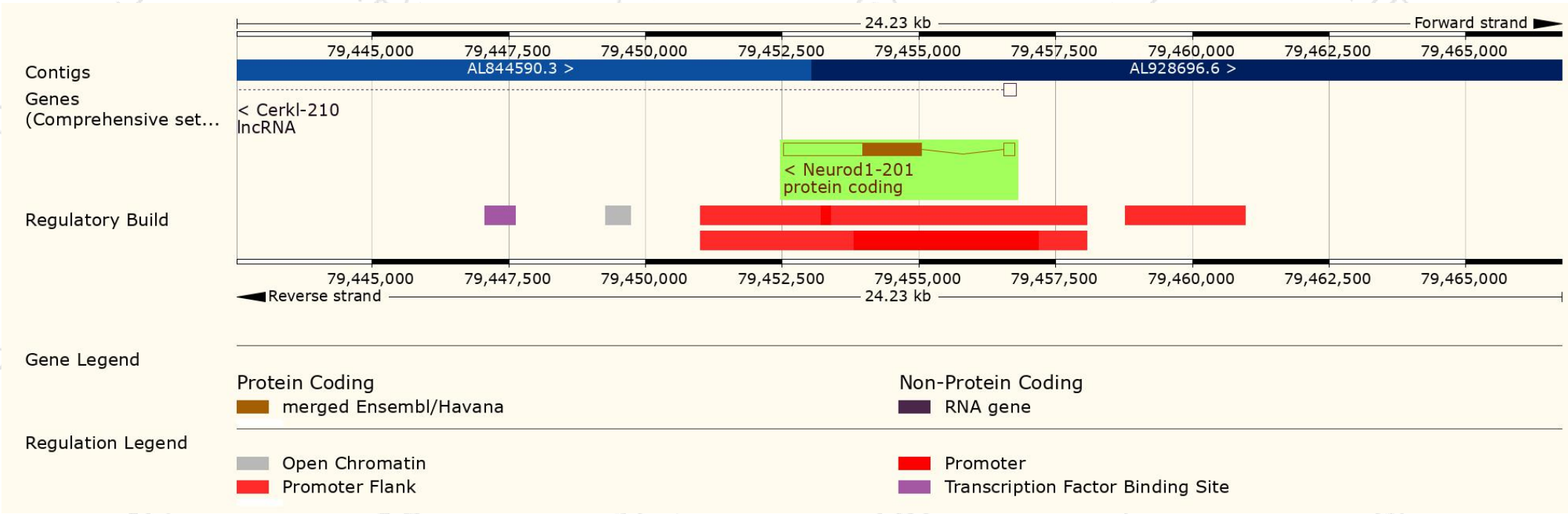
The gene has 1 transcript, and the transcript is shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|-------------|--------------------------------------|------|-----------------------|----------------|---------------------------|------------------------|-------------------------------|
| Neurod1-201 | ENSMUST00000041099.4 | 2729 | 357aa | Protein coding | CCDS16169 | Q60867 | TSL:1 GENCODE basic APPRIS P1 |

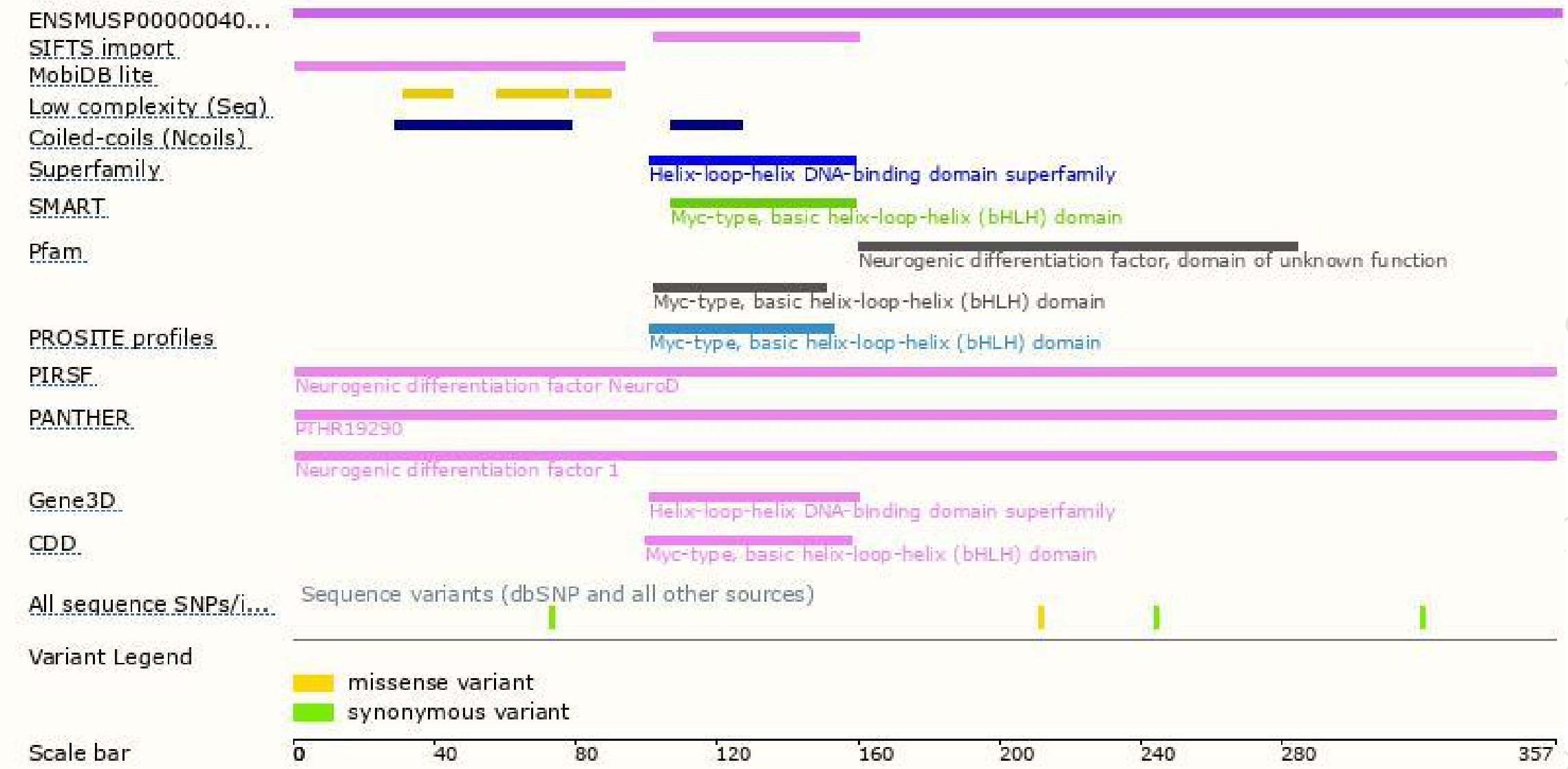
The strategy is based on the design of *Neurod1-201* transcript, The transcription is shown below



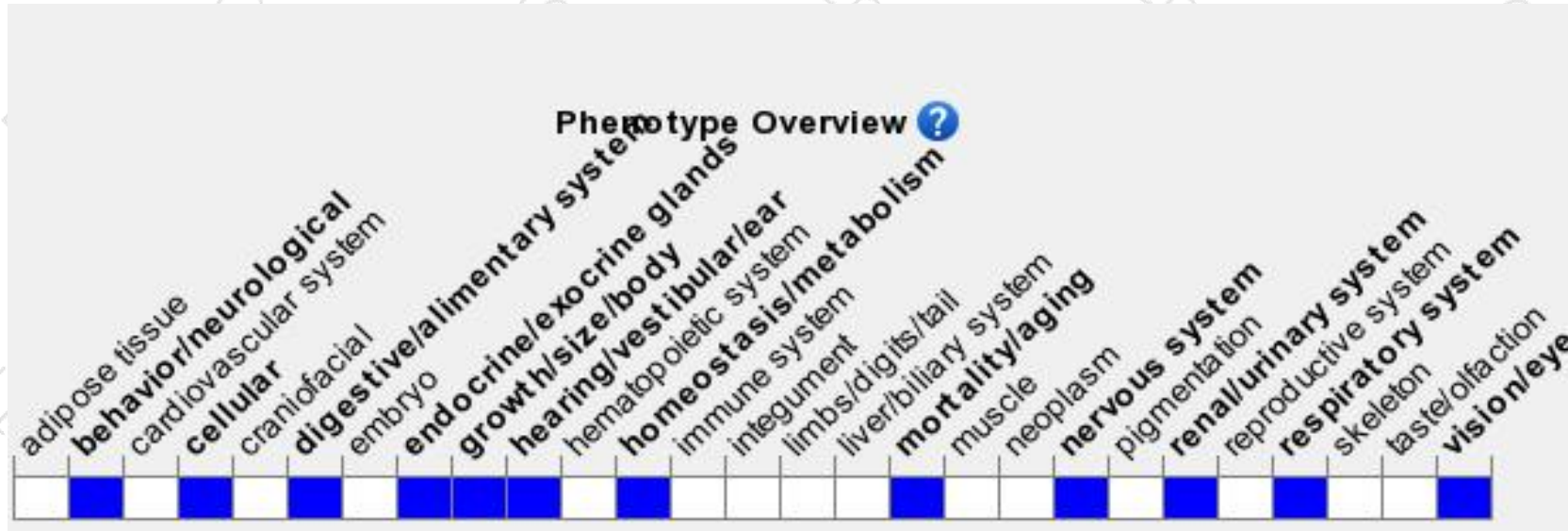
Genomic location distribution



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 targeted null mutations exhibit neonatal diabetes, pancreatic enteroendocrine cell deficits, impaired hearing and balance, retinal degeneration, and seizures. Survival past birth is dependent on genetic background.

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

Tel: 400-9660890

