

# *Nln* Cas9-KO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

*Nln*

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

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



- The *Nln* gene has 8 transcripts. According to the structure of *Nln* gene, exon2-exon3 of *Nln-201* (ENSMUST00000109315.4) transcript is recommended as the knockout region. The region contains 409bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Nln* gene. The brief process is as follows: CRISPR/Cas9 system w

- According to the existing MGI data, Mice homozygous for a null allele exhibit increased glucose tolerance, insulin sensitivity, and gluconeogenesis. Mice also show decreased body weight and run less in a low intensity regime to exhaustion.
- Transcript 206 CDS 5' and 3' incomplete the influences is unknown.
- The *Nln* gene is located on the Chr13. 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)

## Nln neurolysin (metallopeptidase M3 family) [Mus musculus (house mouse)]

Gene ID: 75805, updated on 31-Jan-2019

### Summary



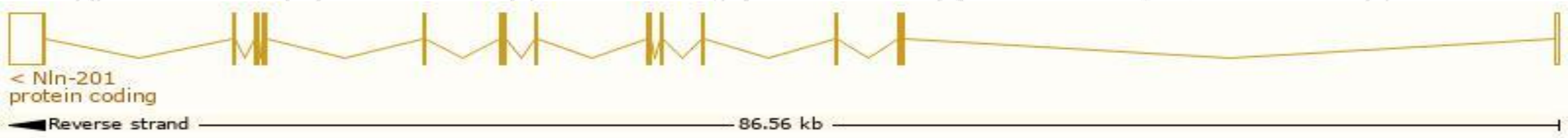
<b>Official Symbol</b>	Nln provided by <a href="#">MGI</a>
<b>Official Full Name</b>	neurolysin (metallopeptidase M3 family) provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1923055</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG000000021710</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	PROVISIONAL
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	4930472G13Rik, C79345
<b>Expression</b>	Ubiquitous expression in CNS E18 (RPKM 12.4), whole brain E14.5 (RPKM 7.3) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

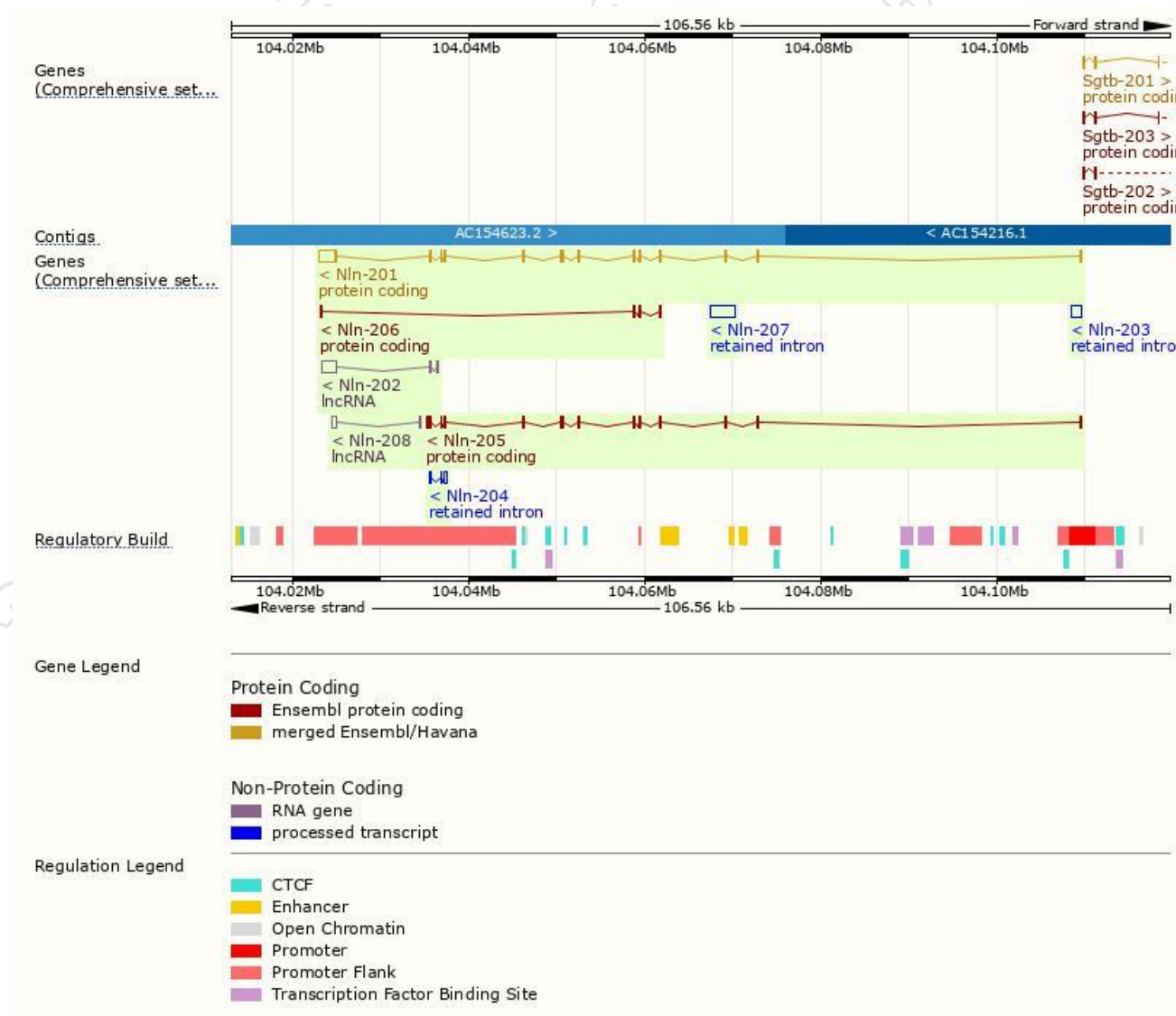
The gene has 8 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
NIn-201	<a href="#">ENSMUST00000109315.4</a>	4162	<a href="#">704aa</a>	Protein coding	<a href="#">CCDS36772</a>	<a href="#">Q91YP2</a>	TSL:1 GENCODE basic APPRIS P1
NIn-205	<a href="#">ENSMUST00000224945.1</a>	2486	<a href="#">683aa</a>	Protein coding	-	<a href="#">A0A286YD12</a>	GENCODE basic
NIn-206	<a href="#">ENSMUST00000225324.1</a>	453	<a href="#">151aa</a>	Protein coding	-	<a href="#">A0A286YD77</a>	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete
NIn-207	<a href="#">ENSMUST00000225478.1</a>	2720	No protein	Retained intron	-	-	
NIn-203	<a href="#">ENSMUST00000224086.1</a>	1213	No protein	Retained intron	-	-	
NIn-204	<a href="#">ENSMUST00000224475.1</a>	560	No protein	Retained intron	-	-	
NIn-202	<a href="#">ENSMUST00000224058.1</a>	1924	No protein	lncRNA	-	-	
NIn-208	<a href="#">ENSMUST00000225704.1</a>	564	No protein	lncRNA	-	-	

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

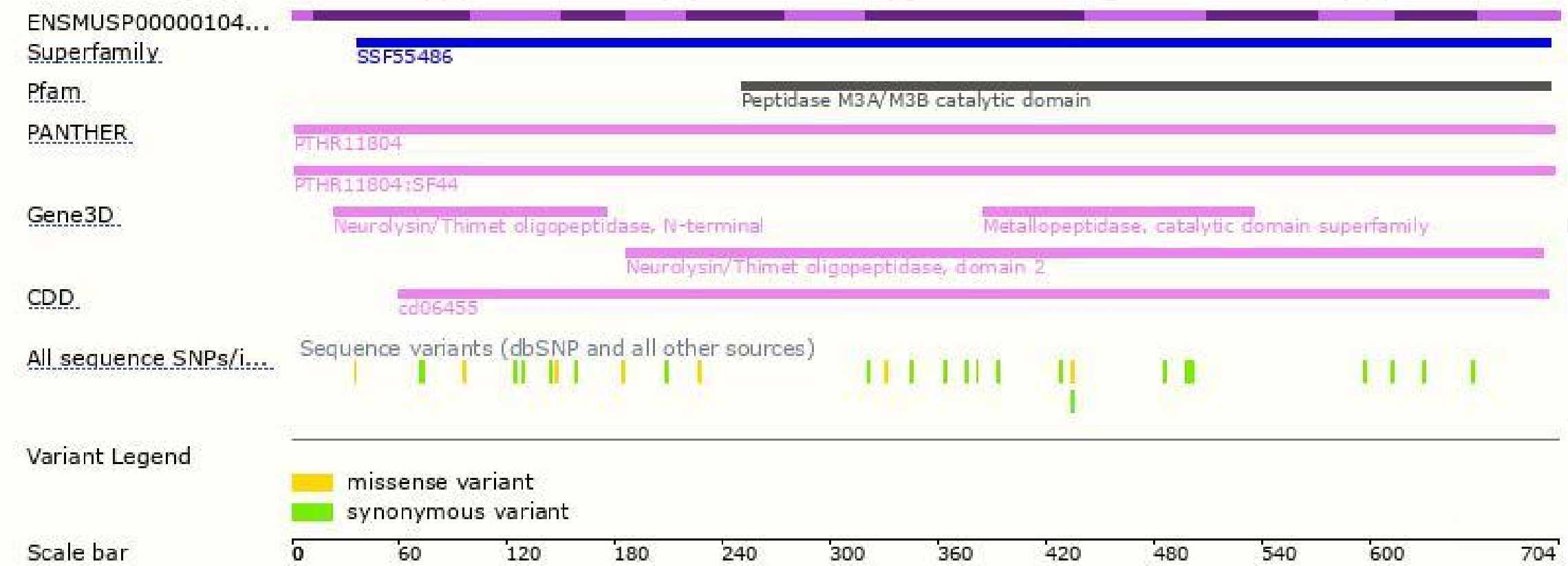


# Genomic location distribution

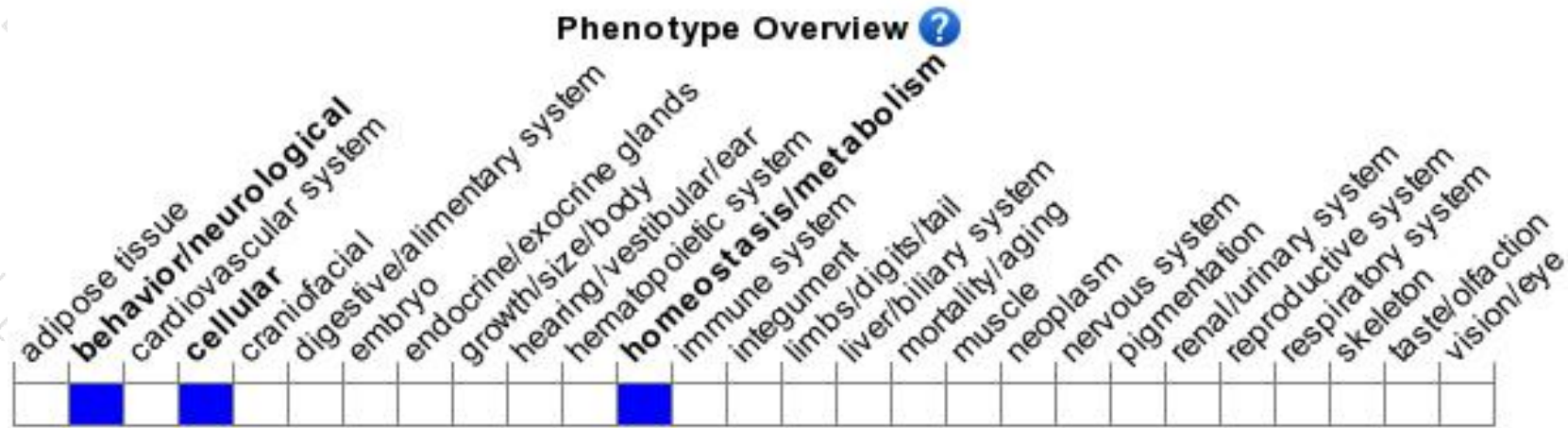




# Protein domain



# 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, Mice homozygous for a null allele exhibit increased glucose tolerance, insulin sensitivity, and gluconeogenesis. Mice also show decreased body weight and run less in a low intensity regime to exhaustion.

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

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