

# *Grin3a* Cas9-KO Strategy

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# Project Overview



**Project Name**

***Grin3a***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Grin3a* gene. The schematic diagram is as follows:



- The *Grin3a* gene has 4 transcripts. According to the structure of *Grin3a* gene, exon2 of *Grin3a-202* (ENSMUST00000093859.10) transcript is recommended as the knockout region. The region contains 605bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Grin3a* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a disruption in this gene display increased current densities in some cerebrocortical neurons of the brain, increased levels of prepulse inhibition, and altered dendritic spine morphology. Otherwise, they display a normal phenotype.
- Transcript *Grin3a-204* may not be affected
- The N-terminal of *Grin3a* gene will remain 233aa, it may remain the partial function of *Grin3a* gene.
- The *Grin3a* gene is located on the Chr4. 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)

## Grin3a glutamate receptor ionotropic, NMDA3A [Mus musculus (house mouse)]

Gene ID: 242443, updated on 31-Jan-2019

### Summary



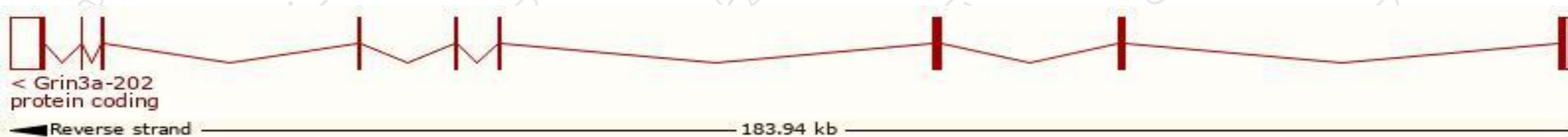
<b>Official Symbol</b>	Grin3a provided by <a href="#">MGI</a>
<b>Official Full Name</b>	glutamate receptor ionotropic, NMDA3A provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1933206</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000039579</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<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>	6430537F04, A830097C19Rik, NMDAR-L, NR3A, mKIAA1973
<b>Expression</b>	Biased expression in CNS E18 (RPKM 8.2), whole brain E14.5 (RPKM 3.6) and 6 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

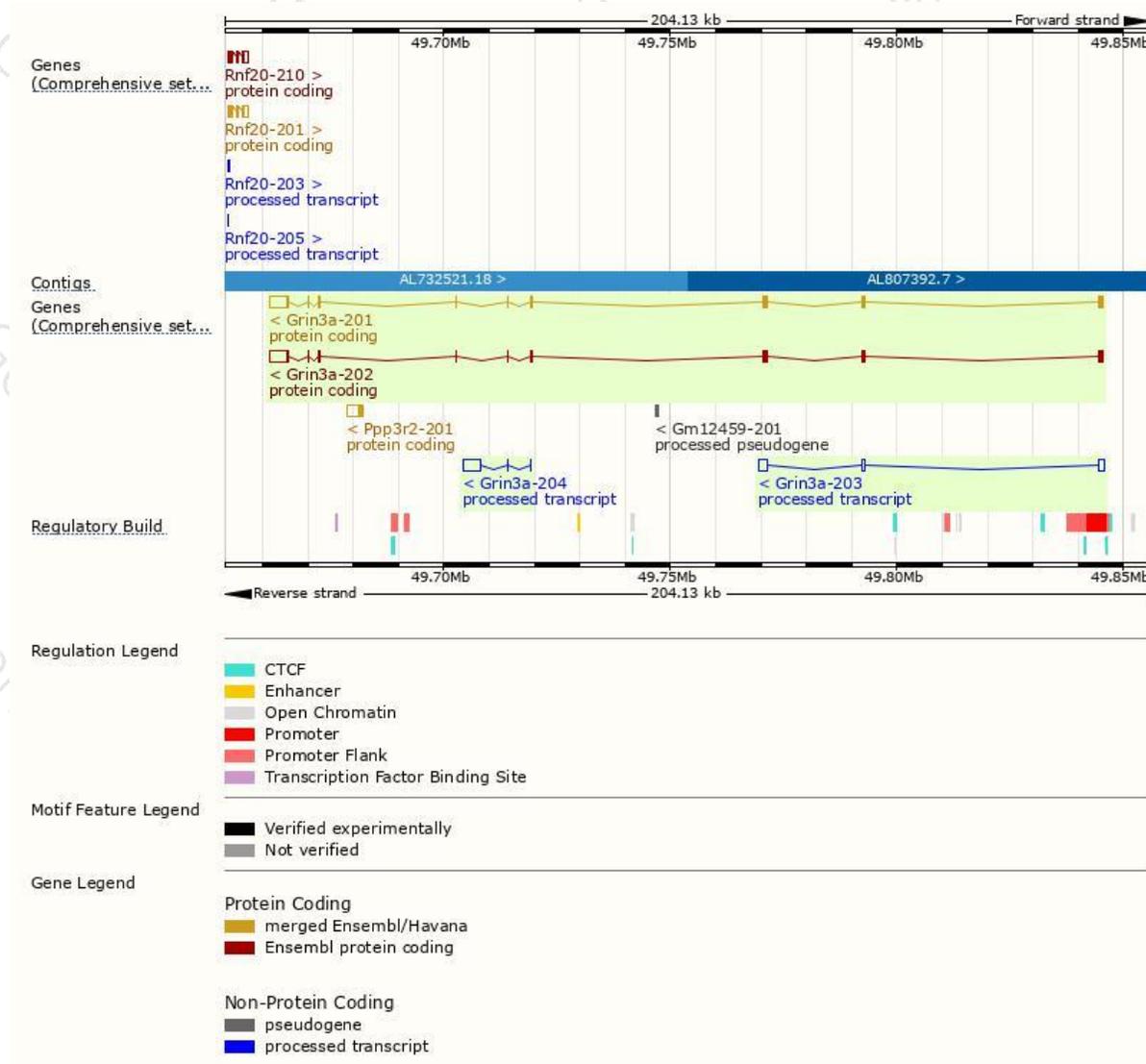
The gene has 4 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Grin3a-202	<a href="#">ENSMUST00000093859.10</a>	7491	<a href="#">1135aa</a>	Protein coding	<a href="#">CCDS71385</a>	<a href="#">A2AIR5</a>	TSL:1 GENCODE basic
Grin3a-201	<a href="#">ENSMUST00000076674.3</a>	7431	<a href="#">1115aa</a>	Protein coding	<a href="#">CCDS51175</a>	<a href="#">A2AIR4</a>	TSL:1 GENCODE basic APPRIS P1
Grin3a-204	<a href="#">ENSMUST00000149059.1</a>	3925	No protein	Processed transcript	-	-	TSL:5
Grin3a-203	<a href="#">ENSMUST00000131797.1</a>	3799	No protein	Processed transcript	-	-	TSL:1

The strategy is based on the design of *Grin3a-202* transcript, The transcription is shown below



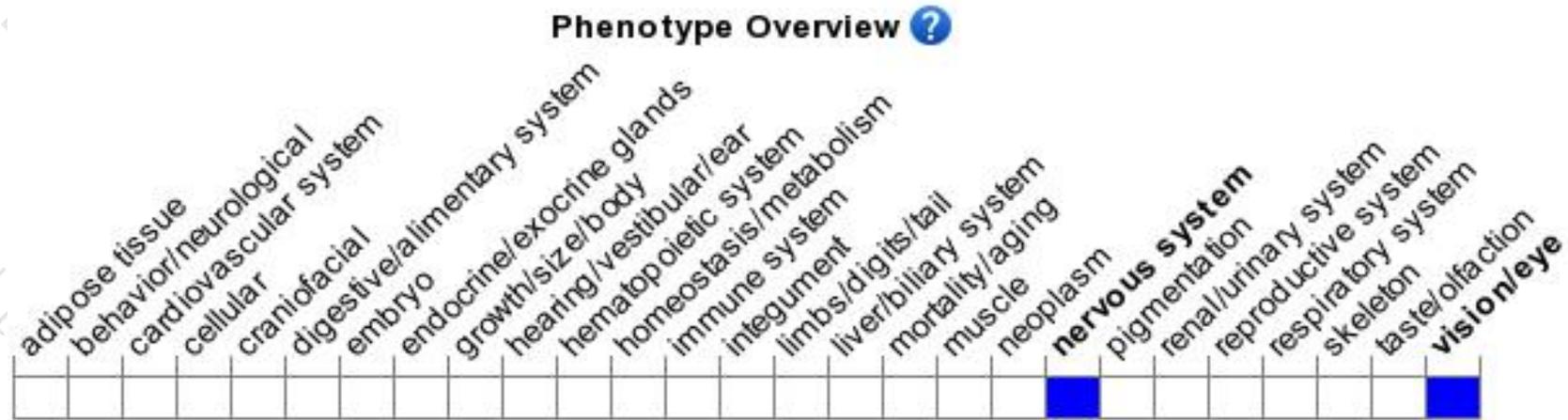
# Genomic location distribution



# 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 disruption in this gene display increased current densities in some cerebrocortical neurons of the brain, increased levels of prepulse inhibition, and altered dendritic spine morphology. Otherwise, they display a normal phenotype.

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

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