

# *Ppp1r9a* Cas9-CKO Strategy

Designer: Xiaojing Li  
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Reviewer: Jia Yu

# Project Overview

**Project Name**

*Ppp1r9a*

**Project type**

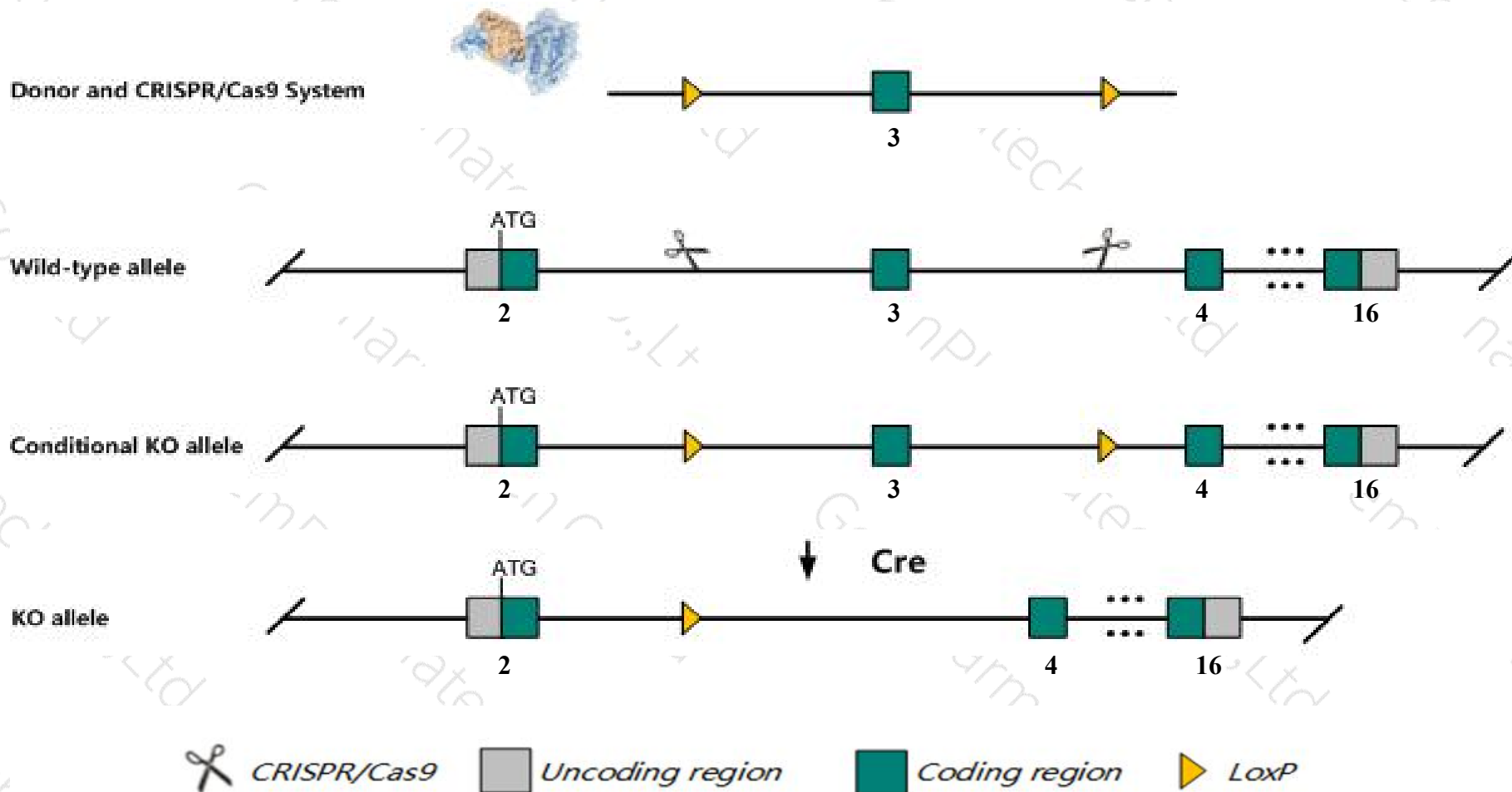
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Ppp1r9a* gene. The schematic diagram is as follows:



- The *Ppp1r9a* gene has 12 transcripts. According to the structure of *Ppp1r9a* gene, exon3 of *Ppp1r9a-201* (ENSMUST00000035813.8) transcript is recommended as the knockout region. The region contains 133bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Ppp1r9a* 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, Mice homozygous for a knock-out allele exhibit defects in dopamine-mediated neuromodulation, deficient long-term potentiation at corticostriatal synapses, increased spontaneous excitatory post-synaptic current frequency, and enhanced locomotor activation in response to cocaine treatment.
- The *Ppp1r9a* gene is located on the Chr6. 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)

## Ppp1r9a protein phosphatase 1, regulatory subunit 9A [ *Mus musculus* (house mouse) ]

Gene ID: 243725, updated on 24-Oct-2019

### Summary

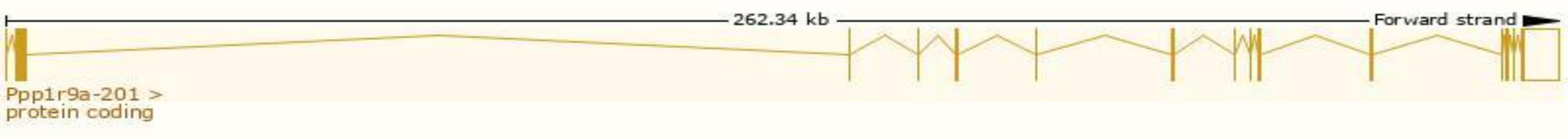
Official Symbol	Ppp1r9a provided by <a href="#">MGI</a>
Official Full Name	protein phosphatase 1, regulatory subunit 9A provided by <a href="#">MGI</a>
Primary source	<a href="#">MGI:MGI:2442401</a>
See related	<a href="#">Ensembl:ENSMUSG00000032827</a>
Gene type	protein coding
RefSeq status	VALIDATED
Organism	<a href="#">Mus musculus</a>
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	NRB; BB181831; 5330407E15; neurabin-I; 2810430P21Rik; 4930518N04Rik; A230094E16Rik
Expression	Broad expression in cortex adult (RPKM 12.5), frontal lobe adult (RPKM 11.0) and 22 other tissues <a href="#">See more</a>
Orthologs	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

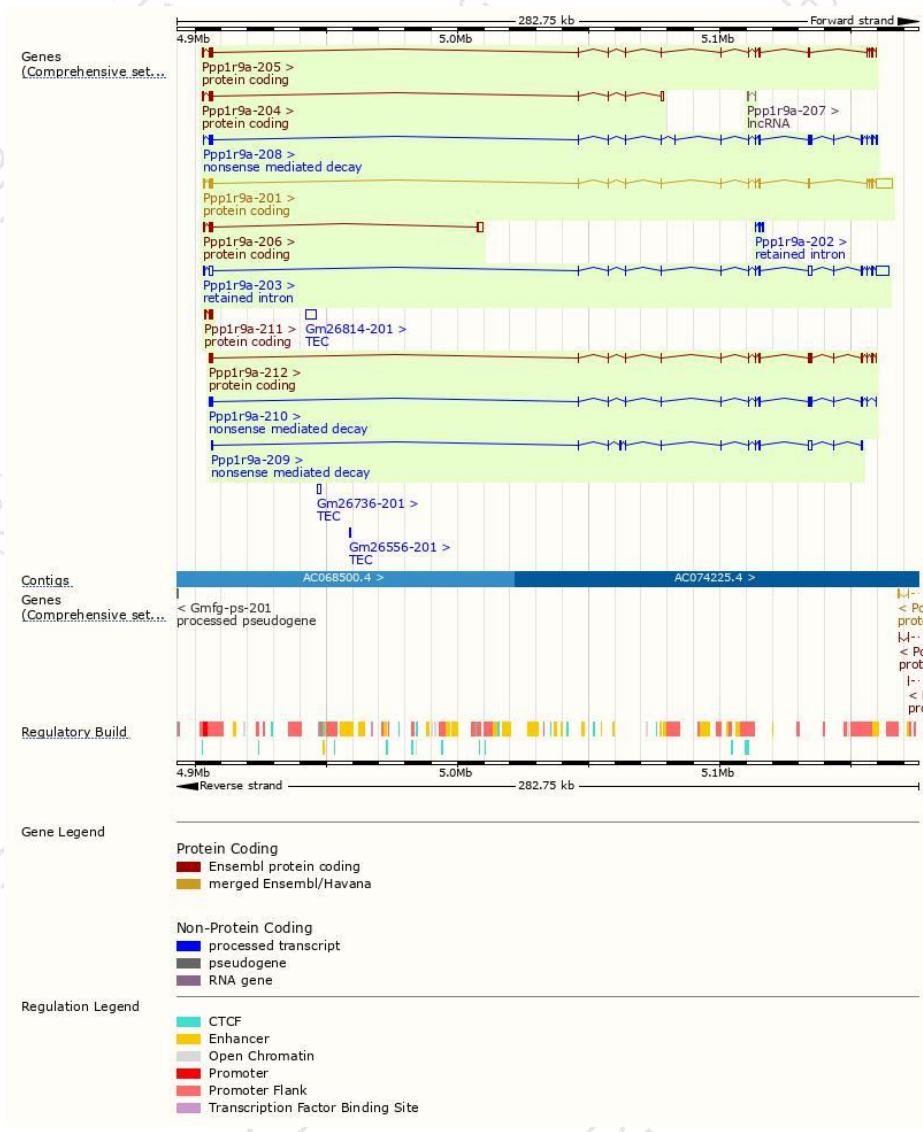
The gene has 12 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ppp1r9a-201	<a href="#">ENSMUST00000035813.8</a>	9547	<a href="#">1095aa</a>	Protein coding	<a href="#">CCDS19897</a>	<a href="#">Q7TN74</a>	TSL:1 GENCODE basic APPRIS P2
Ppp1r9a-212	<a href="#">ENSMUST00000177456.7</a>	3975	<a href="#">1292aa</a>	Protein coding	-	<a href="#">H3BJD6</a>	TSL:5 GENCODE basic APPRIS ALT2
Ppp1r9a-206	<a href="#">ENSMUST00000175962.1</a>	3769	<a href="#">533aa</a>	Protein coding	-	<a href="#">H3BKE7</a>	TSL:1 GENCODE basic
Ppp1r9a-205	<a href="#">ENSMUST00000175889.7</a>	3462	<a href="#">1042aa</a>	Protein coding	-	<a href="#">H3BL28</a>	CDS 3' incomplete TSL:1
Ppp1r9a-204	<a href="#">ENSMUST00000168998.8</a>	2966	<a href="#">642aa</a>	Protein coding	-	<a href="#">Q3UXW4</a>	TSL:1 GENCODE basic
Ppp1r9a-211	<a href="#">ENSMUST00000177338.1</a>	1680	<a href="#">447aa</a>	Protein coding	-	<a href="#">Q8BMP0</a>	CDS 3' incomplete TSL:1
Ppp1r9a-208	<a href="#">ENSMUST00000176263.7</a>	4810	<a href="#">977aa</a>	Nonsense mediated decay	-	<a href="#">H3BJD0</a>	TSL:5
Ppp1r9a-210	<a href="#">ENSMUST00000177153.7</a>	3834	<a href="#">955aa</a>	Nonsense mediated decay	-	<a href="#">H3BKQ7</a>	TSL:5
Ppp1r9a-209	<a href="#">ENSMUST00000176729.7</a>	3046	<a href="#">232aa</a>	Nonsense mediated decay	-	<a href="#">H3BJA6</a>	CDS 5' incomplete TSL:1
Ppp1r9a-203	<a href="#">ENSMUST00000164110.8</a>	9395	No protein	Retained intron	-	-	TSL:1
Ppp1r9a-202	<a href="#">ENSMUST00000065842.6</a>	885	No protein	Retained intron	-	-	TSL:1
Ppp1r9a-207	<a href="#">ENSMUST00000176136.1</a>	357	No protein	lncRNA	-	-	TSL:2

The strategy is based on the design of *Ppp1r9a-201* transcript,The transcription is shown below

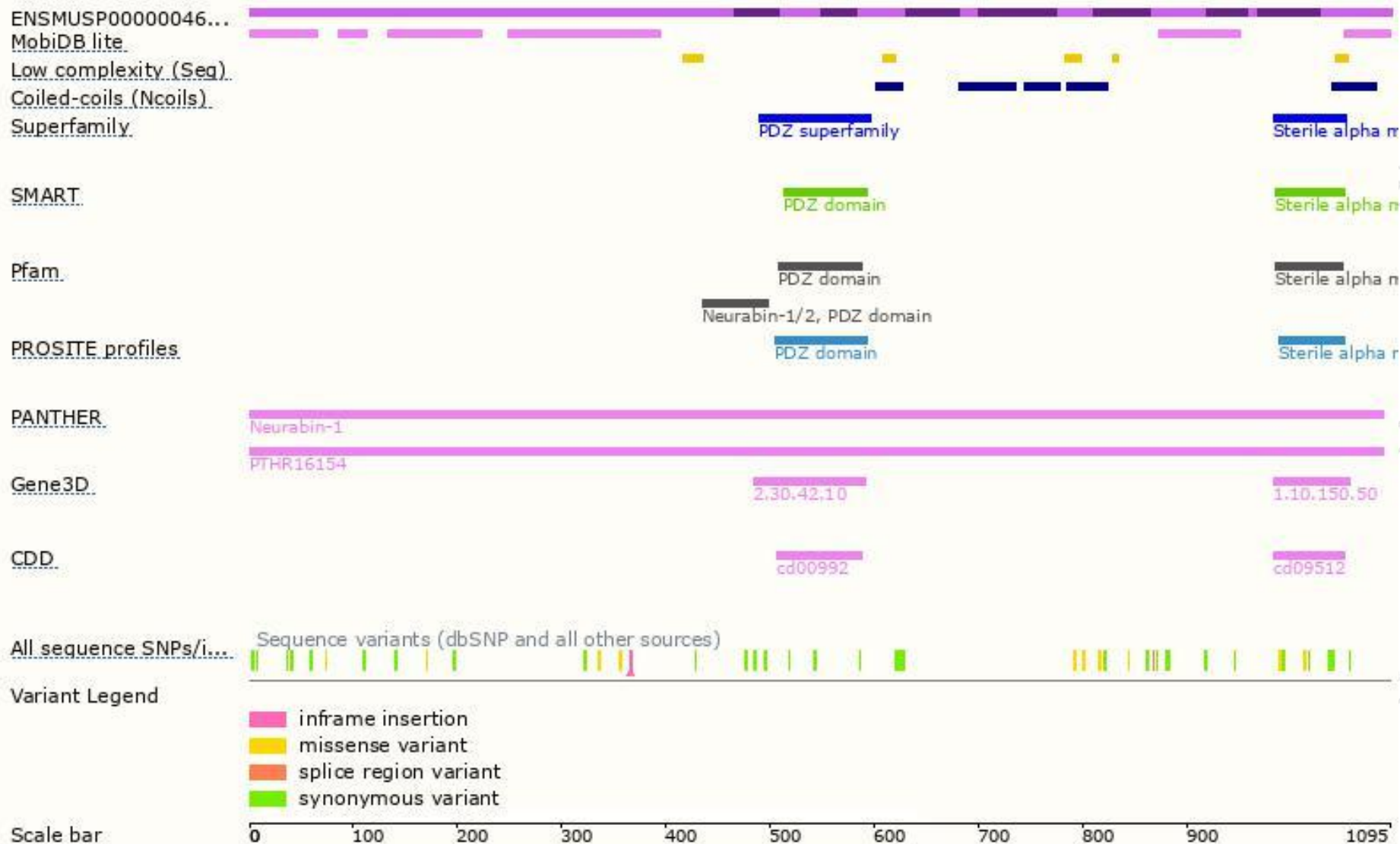


# Genomic location distribution

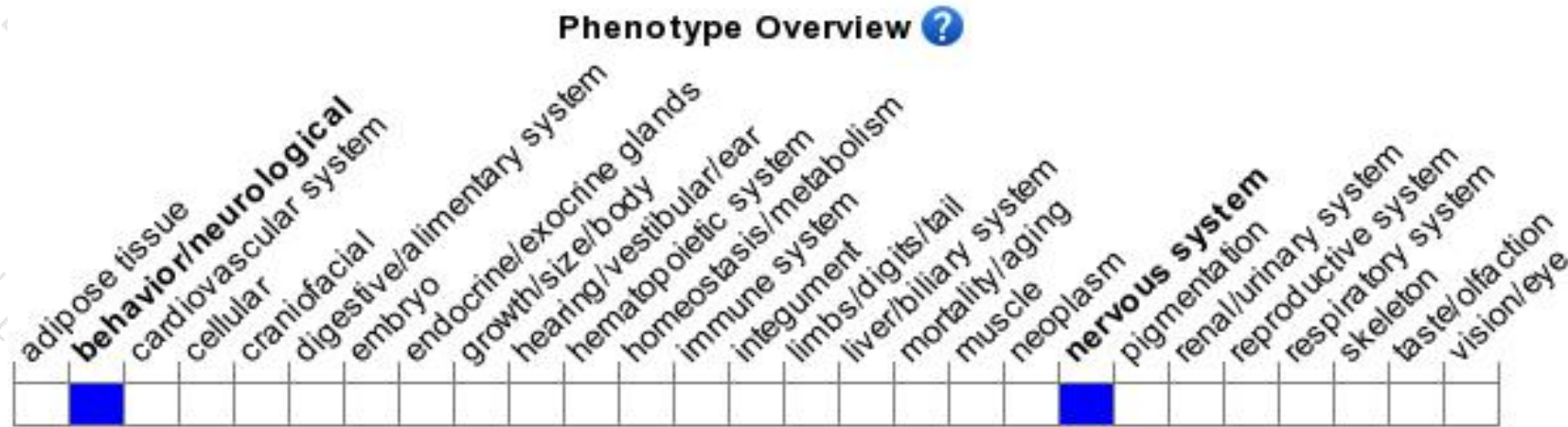




# Protein domain



# 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 knock-out allele exhibit defects in dopamine-mediated neuromodulation, deficient long-term potentiation at corticostriatal synapses, increased spontaneous excitatory post-synaptic current frequency, and enhanced locomotor activation in response to cocaine treatment.

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

Tel: 400-9660890

