

Nptx1 Cas9-CKO Strategy

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Project Overview



Project Name

Nptx1

Project type

Cas9-CKO

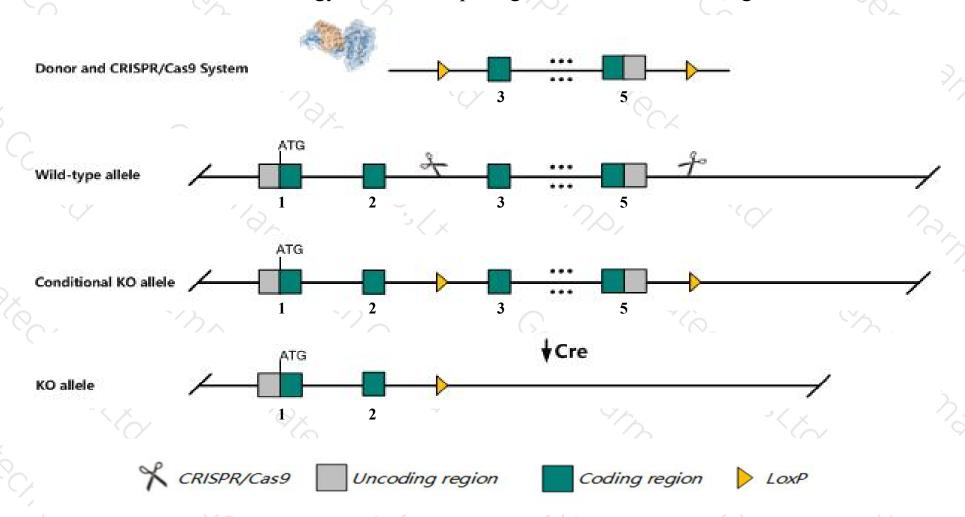
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Nptx1 gene. The schematic diagram is as follows:



Technical routes



- The *Nptx1* gene has 1 transcript. According to the structure of *Nptx1* gene, exon3-exon5 of *Nptx1-201* (ENSMUST00000026670.4) transcript is recommended as the knockout region. The region contains 647bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Nptx1* 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.

Notice



- > According to the existing MGI data, Mice homozygous for a null mutation of this gene display a mild alteration in retinal ganglion cell innervation but are fertile with no obvious behavioral abnormalities.
- ➤ This strategy may affect the 5-terminal regulation of the Gm11762 gene.
- The *Nptx1* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)



Nptx1 neuronal pentraxin 1 [Mus musculus (house mouse)]

Gene ID: 18164, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Nptx1 provided by MGI

Official Full Name neuronal pentraxin 1 provided by MGI

Primary source MGI:MGI:107811

See related Ensembl:ENSMUSG00000025582

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 D11Bwg1004e, Np1

Expression Biased expression in cerebellum adult (RPKM 59.4), cortex adult (RPKM 31.6) and 5 other tissuesSee more

Orthologs <u>human</u> all

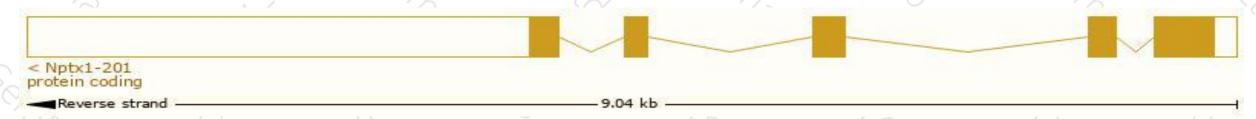
Transcript information (Ensembl)



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

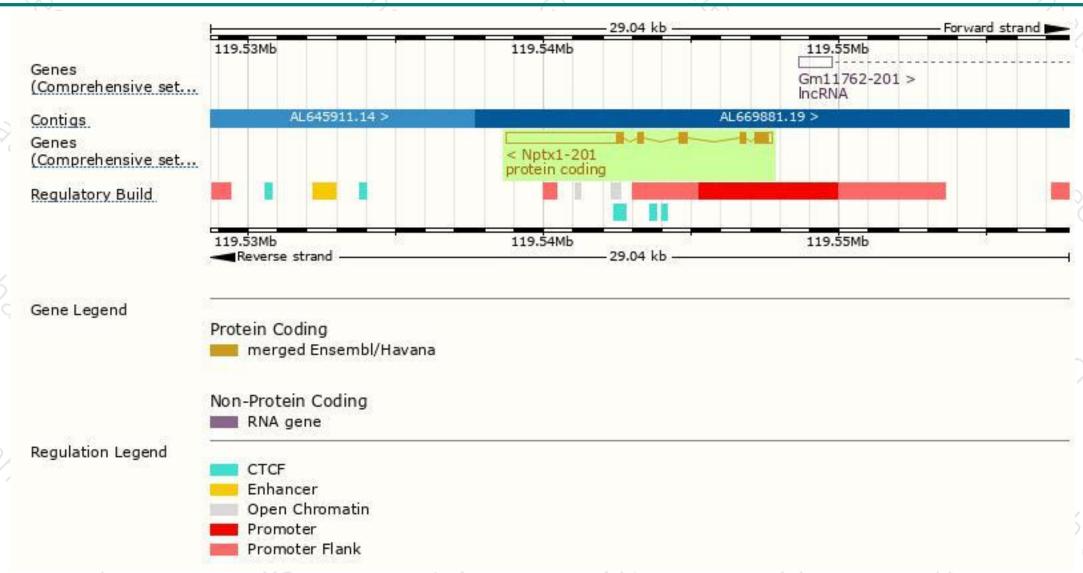
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Nptx1-201	ENSMUST00000026670.4	5217	432aa	Protein coding	CCDS25719	A2ACL9 Q62443	TSL:1 GENCODE basic APPRIS P1	

The strategy is based on the design of Nptx1-201 transcript, The transcription is shown below



Genomic location distribution





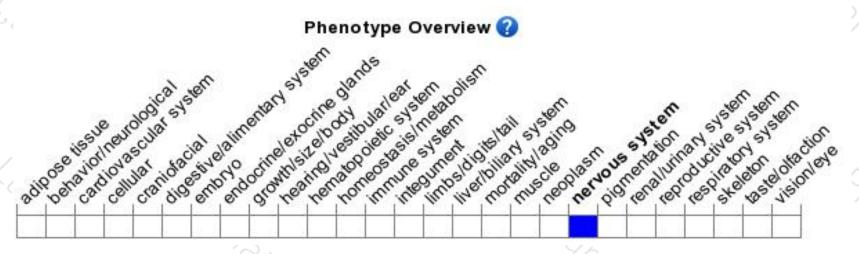
Protein domain





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 mutation of this gene display a mild alteration in retinal ganglion cell innervation but are fertile with no obvious behavioral abnormalities.



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





