

Adnp Cas9-KO Strategy

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Design Date:2020-2-27

Project Overview



Project Name

Adnp

Project type

Cas9-KO

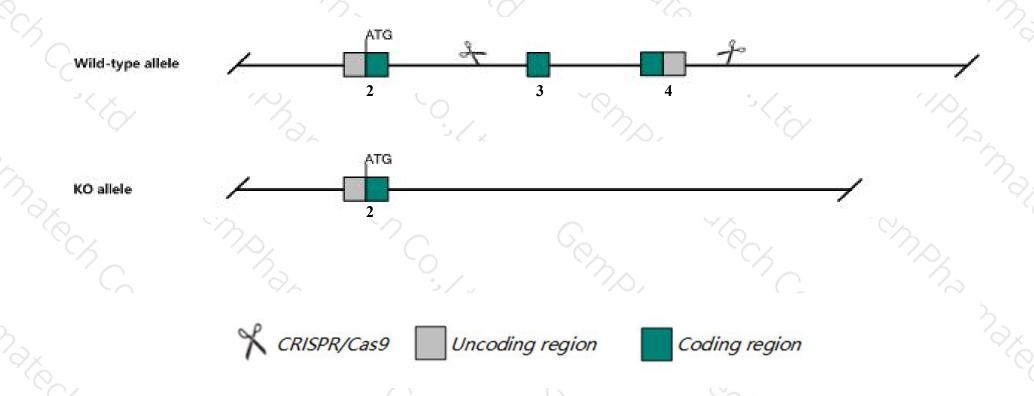
Strain background

C57BL/6JGpt

Knockout strategy



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



Technical routes



- The *Adnp* gene has 4 transcripts. According to the structure of *Adnp* gene, exon3-exon4 of *Adnp-202* (ENSMUST00000088001.5) transcript is recommended as the knockout region. The region contains most 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 *Adnp* gene. The brief process is as follows: gRNA was transcribed in vitro.Cas9 and gRNA 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.

Notice



- ➤ According to the existing MGI data, Developmental defects including the failure of the cranial neural tube to close lead to embryonic death between E8.5 and E9.
- > The *Adnp* 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)



Adnp activity-dependent neuroprotective protein [Mus musculus (house mouse)]

Gene ID: 11538, updated on 17-Feb-2019

Summary

Official Symbol Adnp provided by MGI

Official Full Name activity-dependent neuroprotective protein provided by MGI

MGI:MGI:1338758 Primary source

> Ensembl:ENSMUSG00000051149 See related

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 AA589558, mKIAA0784

This gene encodes a member of a protein family characterized by nine zinc finger motifs followed by a homeobox domain. In vitro studies Summary

demonstrate that the encoded protein interacts with the brahma-related gene1-associated or hBRM factors (BAF) gene expression regulating

complex, components of the protein translation machinery, and microtubule-associated proteins. This gene has been implicated in neuroprotection through various processes that include chromatin remodeling, splicing, cytoskeletal reorganization, and autophagy. Homozygous mutant knockout mice display embryonic lethality with defects in neural tube closure. Alternative splicing results in multiple

transcript variants. [provided by RefSeq, Nov 2016]

Expression Ubiquitous expression in CNS E11.5 (RPKM 36.2), CNS E14 (RPKM 35.6) and 28 other tissuesSee more

Orthologs human all

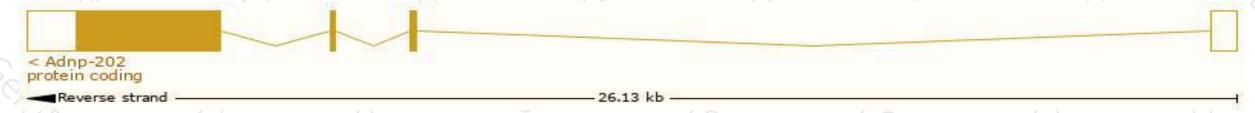
Transcript information (Ensembl)



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

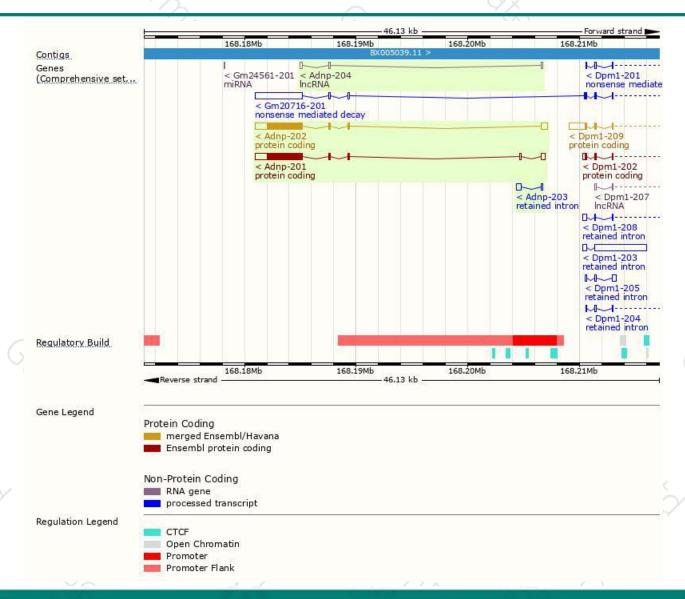
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Adnp-202	ENSMUST00000088001.5	4932	1108aa	Protein coding	CCDS38342	Q9Z103	TSL:1 GENCODE basic APPRIS P1
Adnp-201	ENSMUST00000057793.10	4917	1108aa	Protein coding	CCDS38342	Q9Z103	TSL:5 GENCODE basic APPRIS P1
Adnp-203	ENSMUST00000131200.1	554	No protein	Retained intron		20	TSL:2
Adnp-204	ENSMUST00000139984.7	434	No protein	IncRNA	20	29	TSL:2

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



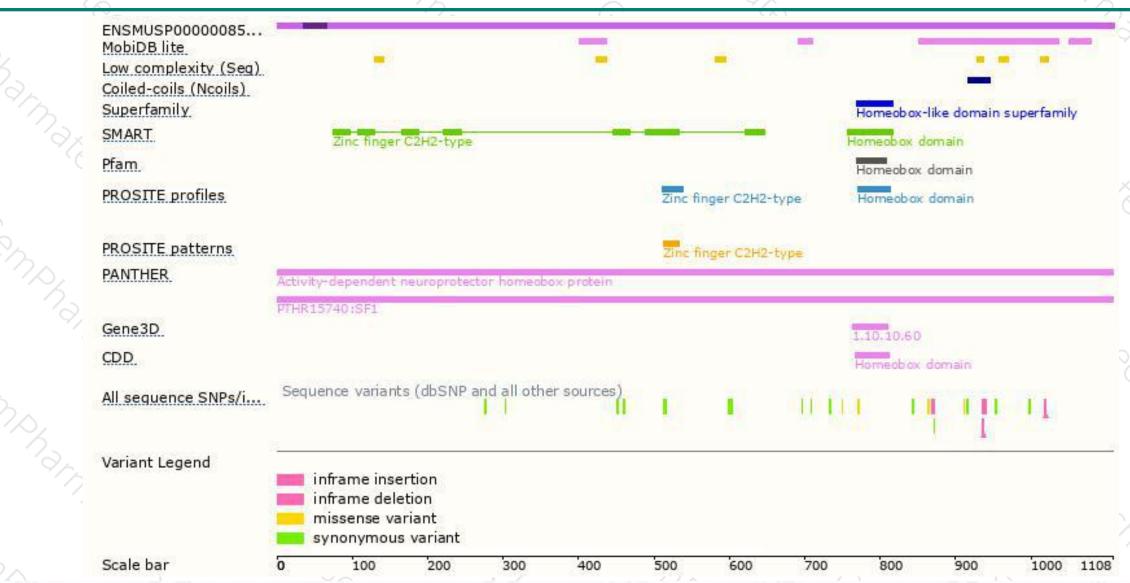
Genomic location distribution





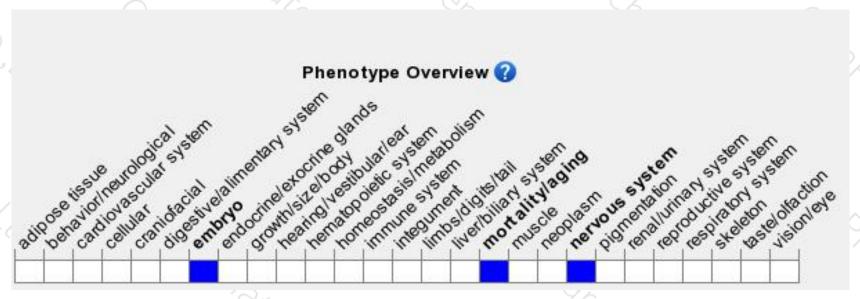
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

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





