

Avpr1a Cas9-CKO Strategy

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Design Date: 2019-8-23

Project Overview



Project Name

Avpr1a

Project type

Cas9-CKO

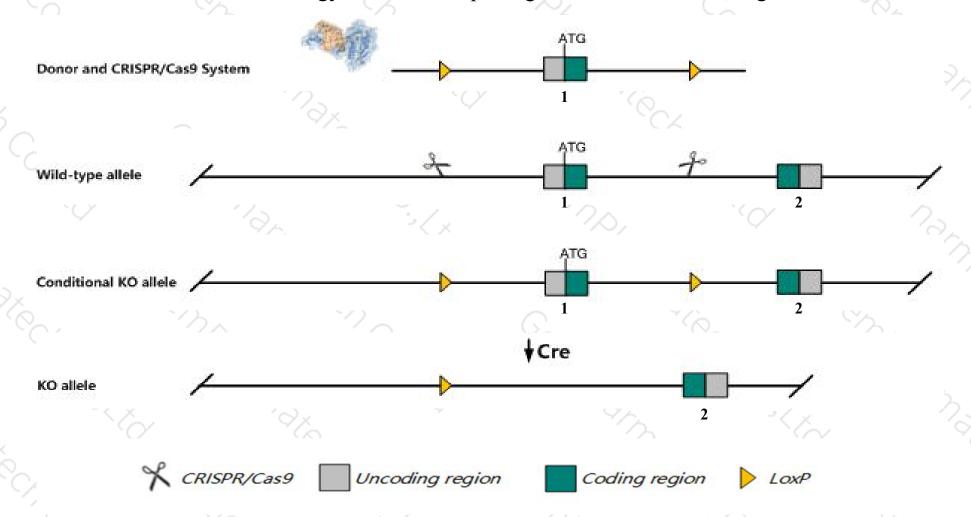
Strain background

C57BL/6JGpt

Conditional Knockout strategy



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



Technical routes



- The *Avpr1a* gene has 1 transcript. According to the structure of *Avpr1a* gene, exon1 of *Avpr1a-201* (ENSMUST00000020323.6) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Avpr1a* 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 disruptions in this gene display a stimulus processing deficit similar to that seen in schizophrenia. Anxiety-like behaviors are reduced in males but not females. B cell development is also affected.
- > The Avpr1a gene is located on the Chr10. 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)



Avpr1a arginine vasopressin receptor 1A [Mus musculus (house mouse)]

Gene ID: 54140, updated on 25-Mar-2019

Summary

↑ ?

Official Symbol Avpr1a provided by MGI

Official Full Name arginine vasopressin receptor 1A provided by MGI

Primary source MGI:MGI:1859216

See related Ensembl:ENSMUSG00000020123

Gene type protein coding
RefSeq status REVIEWED

Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as AVPR, Avpr1, V1a, V1aR

Summary This gene encodes a receptor for arginine vasopressin, a neurohypophyseal hormone involved in diuresis inhibition, smooth muscle

contraction, liver glycogenolysis stimulation and regulation of adrenocorticotropic hormone release from the pituitary. This receptor represents one of three G protein-coupled arginine vasopressin receptors which functions through a phosphotidylinositol-calcium second messenger

system in vascular and hepatic tissues [provided by RefSeq, Jul 2008]

Expression Biased expression in liver adult (RPKM 15.9), bladder adult (RPKM 11.1) and 13 other tissuesSee more

Orthologs <u>human</u> all

Transcript information (Ensembl)



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

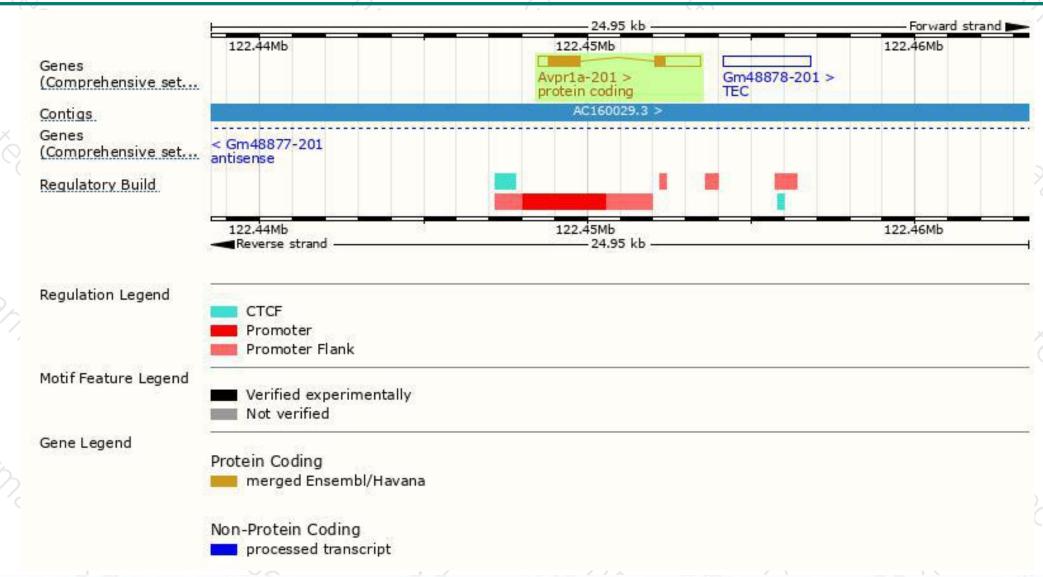
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	l
Avpr1a-201	ENSMUST00000020323.6	2670	423aa	Protein coding	CCDS24215	Q3U1H9 Q62463	TSL:1 GENCODE basic APPRIS P1	

The strategy is based on the design of Avpr1a-201 transcript, The transcription is shown below



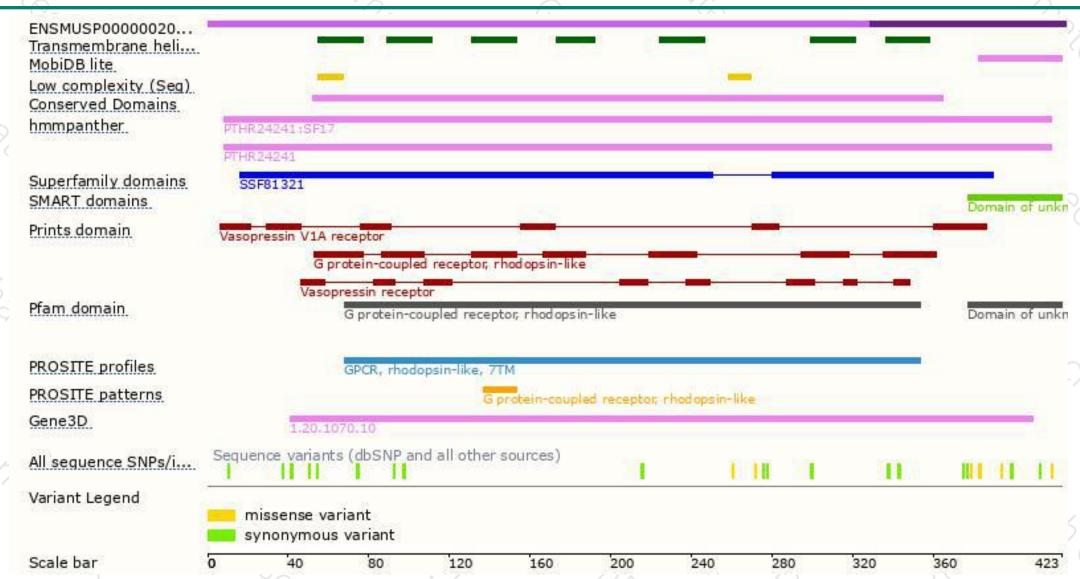
Genomic location distribution





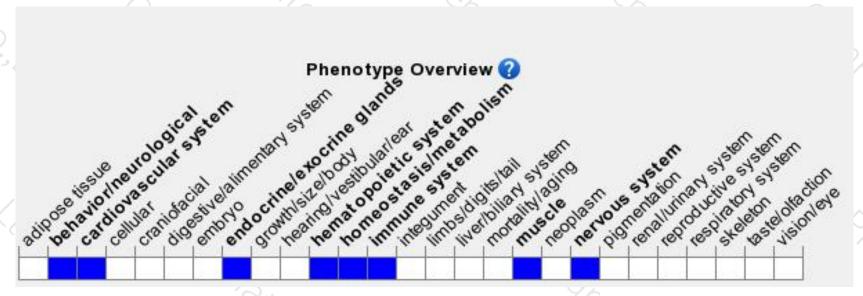
Protein domain





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 disruptions in this gene display a stimulus processing deficit similar to that seen in schizophrenia. Anxiety-like behaviors are reduced in males but not females. B cell development is also affected.



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





