

Ccnd2 Cas9-CKO Strategy

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Design Date: 2019-9-20

Project Overview



Project Name

Ccnd2

Project type

Cas9-CKO

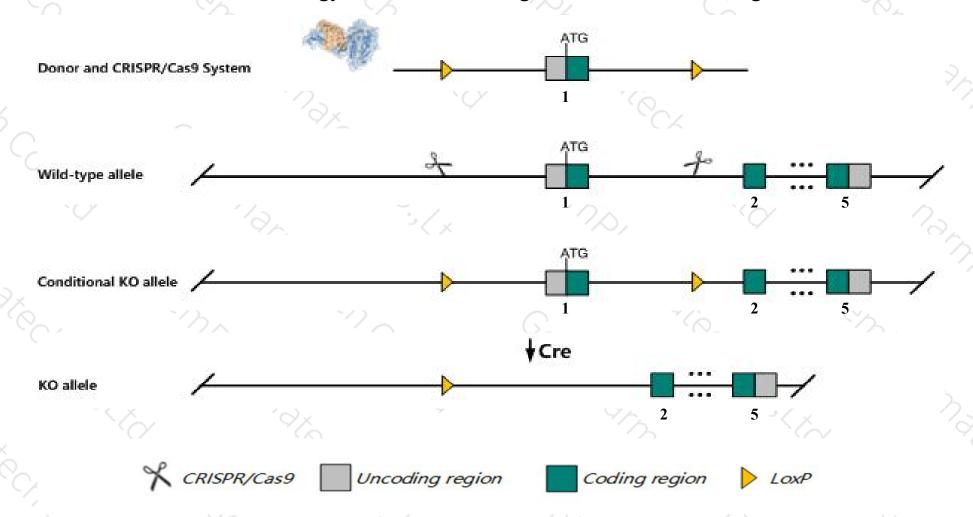
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Ccnd2 gene. The schematic diagram is as follows:



Technical routes



- The *Ccnd2* gene has 8 transcripts. According to the structure of *Ccnd2* gene, exon1 of *Ccnd2-201* (ENSMUST0000000188.11) 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 *Ccnd2* 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, Homozygotes for a targeted null mutation are sterile: females lack a granulosa cell response to follicle stimulating hormone, while males have hypoplastic testes. Mutants also show decreased cerebellar granule cell and stellate neuron populations.
- The KO region contains intron of the 9330179D12Rik gene. Knockout the region may affect the splicing function of 9330179D12Rik gene
- > The *Ccnd2* 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)



Ccnd2 cyclin D2 [Mus musculus (house mouse)]

Gene ID: 12444, updated on 11-Sep-2019

Summary

☆ ?

Official Symbol Ccnd2 provided by MGI

Official Full Name cyclin D2 provided by MGI

Primary source MGI:MGI:88314

See related Ensembl:ENSMUSG00000000184

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 cD2; Vin1; Vin-1; C86853; Al256817; BF642806; 2600016F06Rik

Expression Broad expression in CNS E11.5 (RPKM 77.3), limb E14.5 (RPKM 41.5) and 22 other tissues See more

Orthologs <u>human</u> <u>all</u>

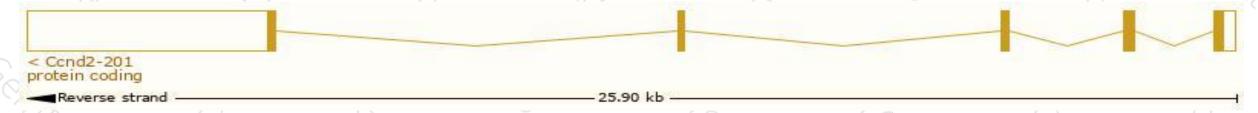
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

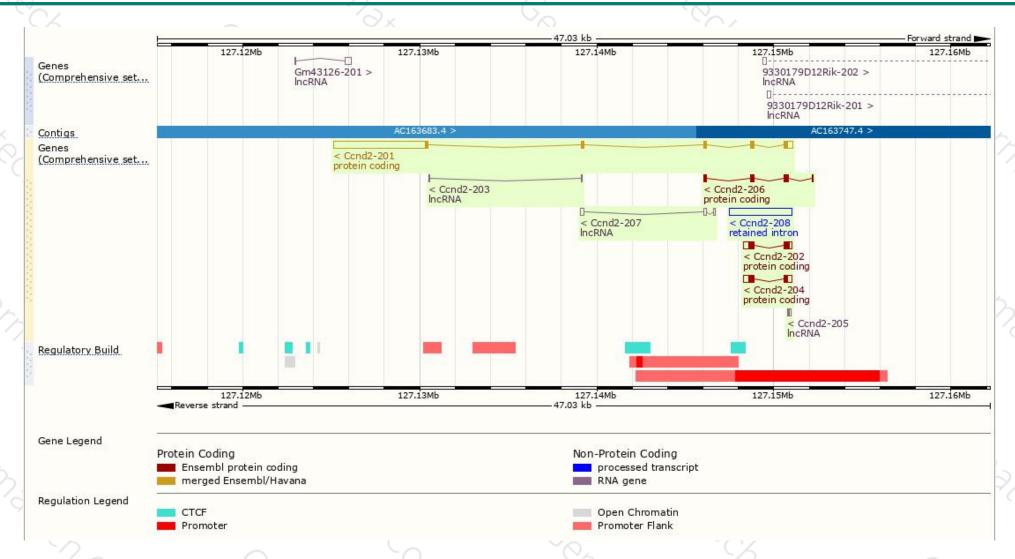
Name 🍦	Transcript ID	bp 🌲	Protein	Biotype	CCDS 🍦	UniProt	Flags
Ccnd2-201	ENSMUST00000000188.11	6330	289aa	Protein coding	CCDS20564₽	P30280 ₽ Q4FK45 ₽	TSL:1 GENCODE basic APPRIS P1
Ccnd2-204	ENSMUST00000201637.1	1006	<u>156aa</u>	Protein coding	-	Q9D8L9@	TSL:1 GENCODE basic
Ccnd2-202	ENSMUST00000201066.1	983	<u>156aa</u>	Protein coding	-	Q9D8L9₽	TSL:2 GENCODE basic
Ccnd2-206	ENSMUST00000201902.3	627	189aa	Protein coding	-	A0A0J9YU51₽	CDS 3' incomplete TSL:5
Ccnd2-208	ENSMUST00000202363.1	3506	No protein	Retained intron	-	-	TSL:NA
Ccnd2-207	ENSMUST00000201985.1	415	No protein	IncRNA	-	-	TSL:2
Ccnd2-203	ENSMUST00000201564.1	127	No protein	IncRNA	-	-	TSL:1
Ccnd2-205	ENSMUST00000201857.1	115	No protein	IncRNA	-	-	TSL:3

The strategy is based on the design of *Ccnd2-201* transcript, The transcription is shown below



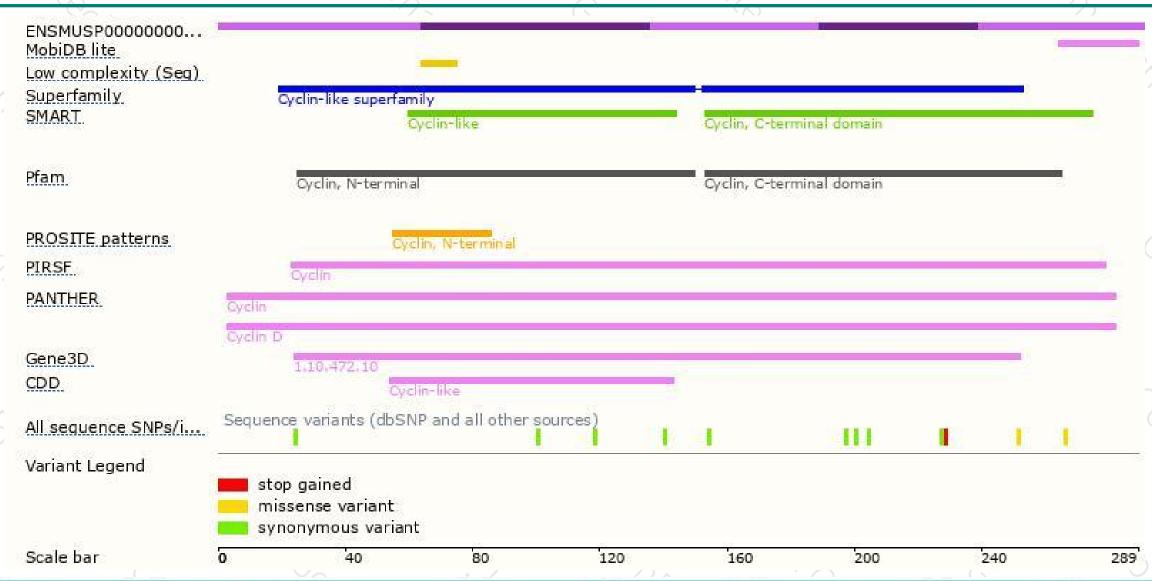
Genomic location distribution





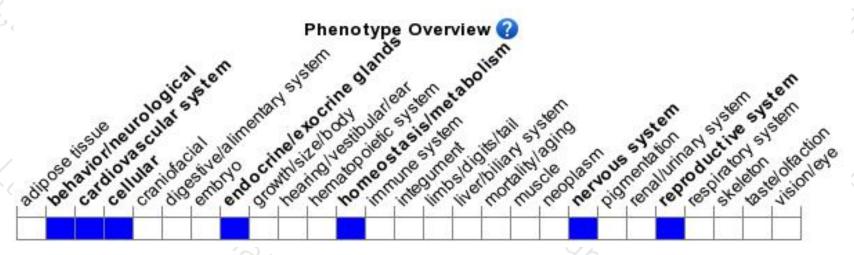
Protein domain





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, Homozygotes for a targeted null mutation are sterile: females lack a granulosa cell response to follicle stimulating hormone, while males have hypoplastic testes. Mutants also show decreased cerebellar granule cell and stellate neuron populations.



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





