

# Cdk1 Cas9-CKO Strategy

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# **Project Overview**



Project Name Cdk1

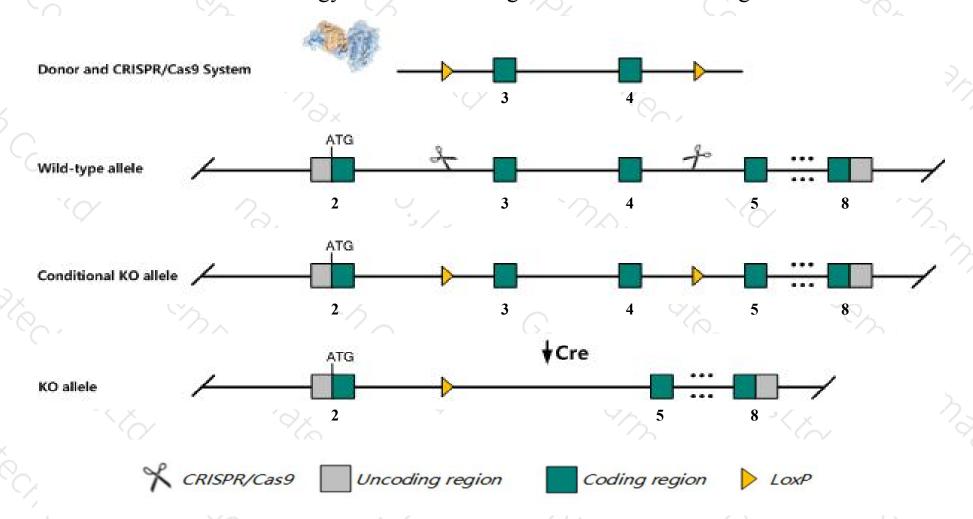
Project type Cas9-CKO

Strain background C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Cdk1* gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Cdk1* gene has 7 transcripts. According to the structure of *Cdk1* gene, exon3-exon4 of *Cdk1-201* (ENSMUST00000020099.12) transcript is recommended as the knockout region. The region contains 281bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cdk1* 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 allele die prior to e1.5.
- The *Cdk1* 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)



#### Cdk1 cyclin-dependent kinase 1 [Mus musculus (house mouse)]

Gene ID: 12534, updated on 13-Mar-2020

#### Summary

^ ?

Official Symbol Cdk1 provided by MGI

Official Full Name cyclin-dependent kinase 1 provided by MGI

Primary source MGI:MGI:88351

See related Ensembl: ENSMUSG00000019942

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 Cdc2, Cdc2a, p34

Expression Biased expression in liver E14 (RPKM 47.0), placenta adult (RPKM 44.1) and 11 other tissuesSee more

Orthologs human all

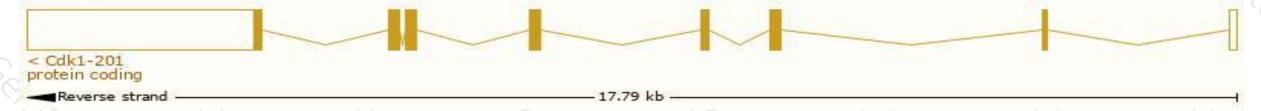
# Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

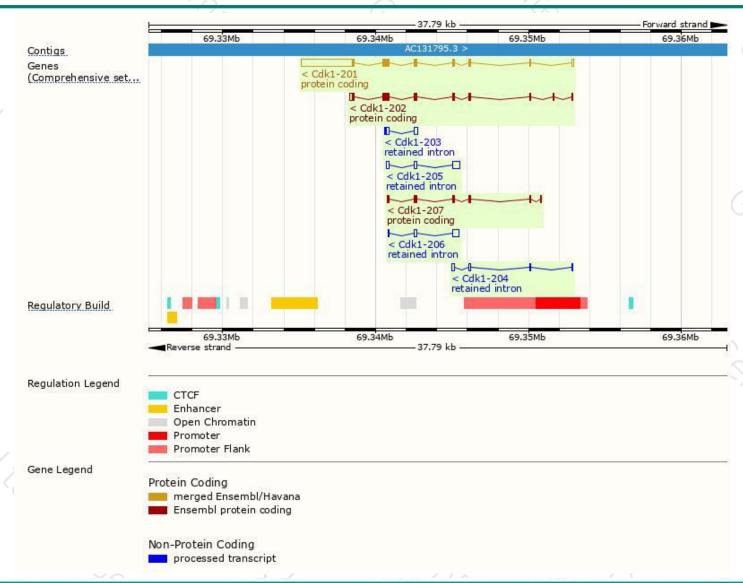
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cdk1-201	ENSMUST00000020099.12	4375	297aa	Protein coding	CCDS23908	P11440	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Cdk1-202	ENSMUST00000119827.7	1253	297aa	Protein coding	CCDS23908	P11440	TSL:5 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Cdk1-207	ENSMUST00000152448.1	695	200aa	Protein coding	923	D3Z2T9	CDS 3' incomplete TSL:3
Cdk1-205	ENSMUST00000129444.7	770	No protein	Retained intron	-	2	TSL:3
Cdk1-206	ENSMUST00000149474.1	616	No protein	Retained intron	170	-	TSL:5
Cdk1-204	ENSMUST00000124054.1	475	No protein	Retained intron		-	TSL:2
Cdk1-203	ENSMUST00000123458.1	378	No protein	Retained intron	(4)	-	TSL:2

The strategy is based on the design of Cdk1-201 transcript, the transcription is shown below:



### Genomic location distribution





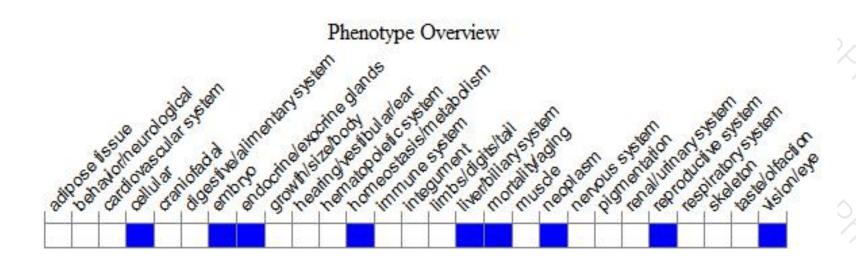
### Protein domain





# 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 allele die prior to E1.5.



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





