

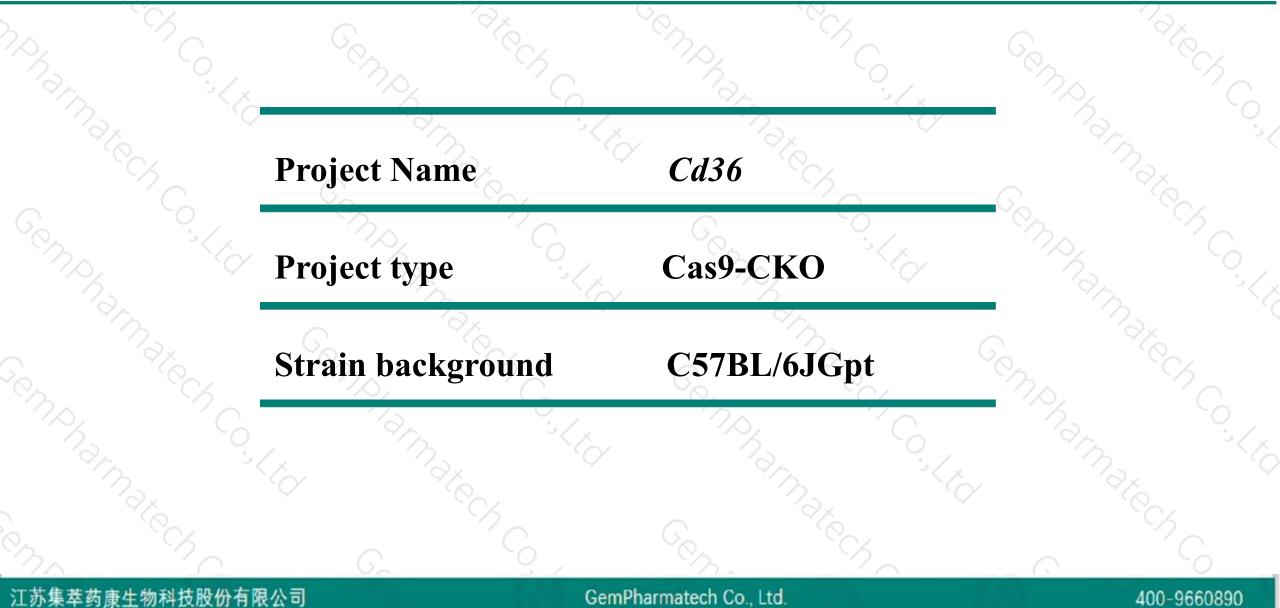
# ND AMAR AC Cd36 Cas9-CKO Strategy Romphamater Control

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# **Project Overview**

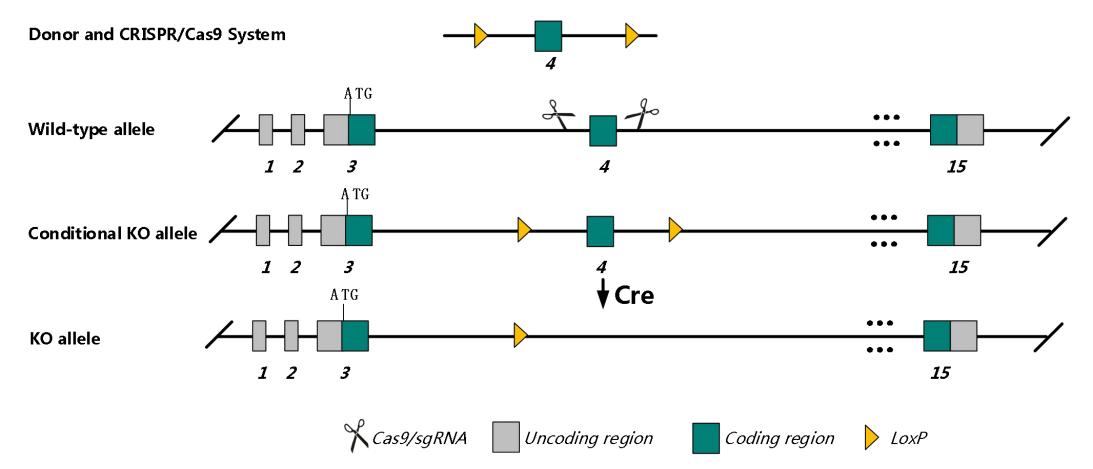




# **Conditional Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Cd36 gene. The schematic diagram is as follows:





The *Cd36* gene has 6 transcripts. According to the structure of *Cd36* gene, exon4 of *Cd36-204* (ENSMUST00000170051.7) transcript is recommended as the knockout region. The region contains 161bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Cd36* 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.



According to the existing MGI data, Homozygous mutant mice exhibit an immunodeficiency phenotype, are susceptible to S. aureus infection and develop ocular pterygium. Mice homozygous for disruptions in this gene display abnormal lipid homeostasis which affects energy utilization in the heart.

The *Cd36* gene is located on the Chr5. 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**



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#### Cd36 CD36 molecule [Mus musculus (house mouse)]

Gene ID: 12491, updated on 2-Apr-2019

#### Summary

Official SymbolCd36 provided by MGIOfficial Full NameCD36 molecule provided byMGIPrimary sourceMGI:MGI:107899See relatedEnsembl:ENSMUSG0000002944Gene typeprotein codingRefSeq statusVALIDATEDOrganismMus musculusLineageEukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Rodentia; Myomorpha;<br/>Muroidea; Murinae; Mus; MusAlso knownasFAT, GPIV, Scarb3ExpressionBiased expression in subcutaneous fat pad adult (RPKM 144.0), heart adult (RPKM 110.7) and 5 other tissues<br/>See more<br/>human all

**NCBI** 

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# **Transcript information**





The gene has 6 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cd36-204	ENSMUST00000170051.7	3399	<u>472aa</u>	Protein coding	CCDS19100	Q08857 Q3UAI3	TSL:1 GENCODE basic APPRIS P1
Cd36-201	ENSMUST0000082367.12	3016	<u>472aa</u>	Protein coding	CCDS19100	Q08857 Q3UAI3	TSL:1 GENCODE basic APPRIS P1
Cd36-202	ENSMUST00000165232.7	2708	<u>472aa</u>	Protein coding	CCDS19100	Q08857 Q3UAI3	TSL:1 GENCODE basic APPRIS P1
Cd36-203	ENSMUST00000169095.5	2677	<u>472aa</u>	Protein coding	CCDS19100	Q08857 Q3UAI3	TSL:1 GENCODE basic APPRIS P1
Cd36-206	ENSMUST00000197890.4	2564	<u>472aa</u>	Protein coding	CCDS19100	Q08857 Q3UAI3	TSL:1 GENCODE basic APPRIS P1
Cd36-205	ENSMUST00000197574.1	702	<u>142aa</u>	Protein coding	8 <del>.</del>	A0A0G2JFB7	CDS 3' incomplete TSL:5

The strategy is based on the design of Cd36-204 transcript, The transcription is shown below

< Cd36-204 protein coding

Reverse strand

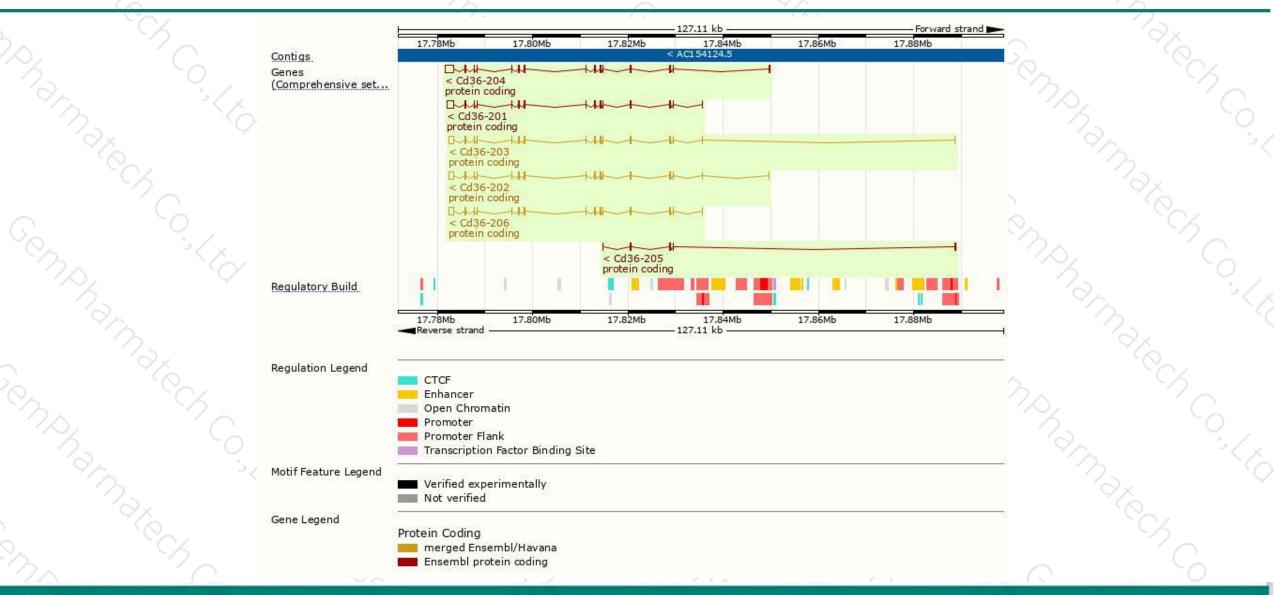
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## **Genomic location distribution**



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# **Protein domain**

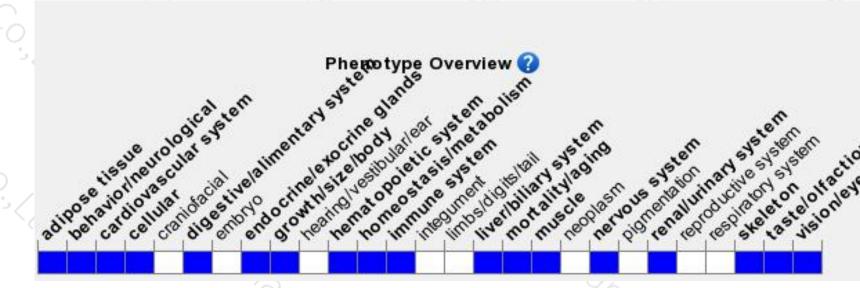
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	Variant Legend		sense vari onymous v						6
	Pfam domain All sequence SNPs/i	CD 36 f Sequeno			all other source	es)	n m		
3	Prints domain	CD36 fami CD36/sca	venger rece	ptor class B	member 1				G
	ENSMUSP00000133 Transmembrane heli Low complexity (Seg) hmmpanther	CD 36	τ <b>υ</b>	-11		in air			

# 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, Homozygous mutant mice exhibit an immunodeficiency phenotype, are susceptible to S. aureus infection and develop ocular pterygium. Mice homozygous for disruptions in this gene display abnormal lipid homeostasis which affects energy utilization in the heart.

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



