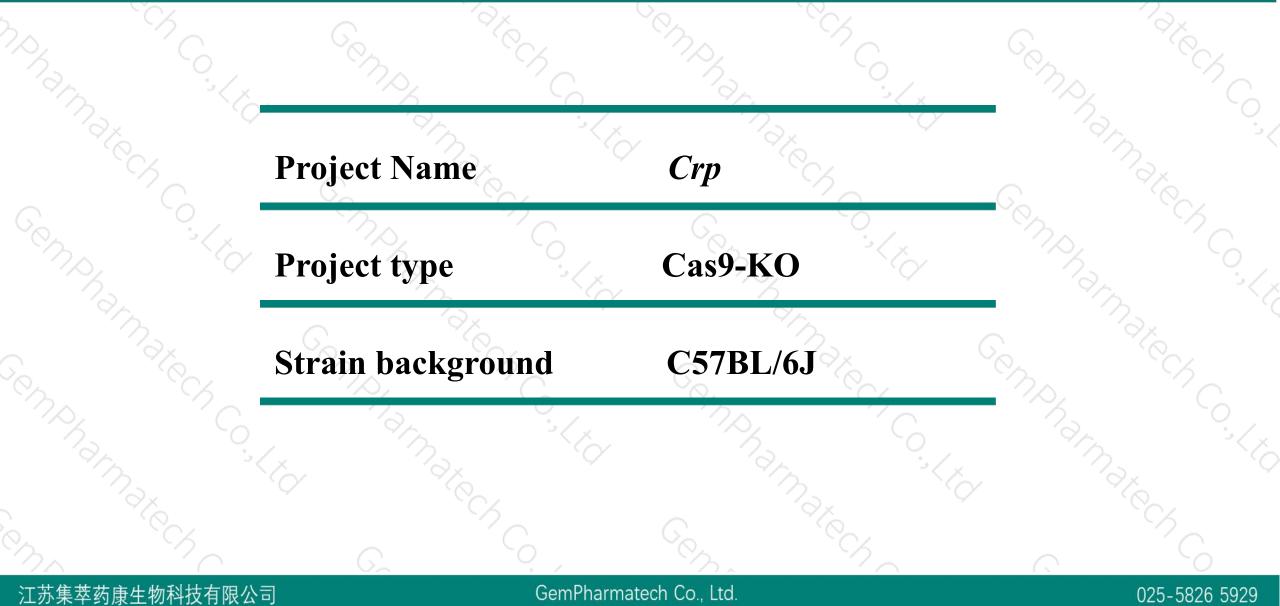


# Crp Cas9-KO Strategy Andraker Costy Cemphamatech (

Comphannated Co Designer:Qiong Zhou Semphamatech C

## **Project Overview**

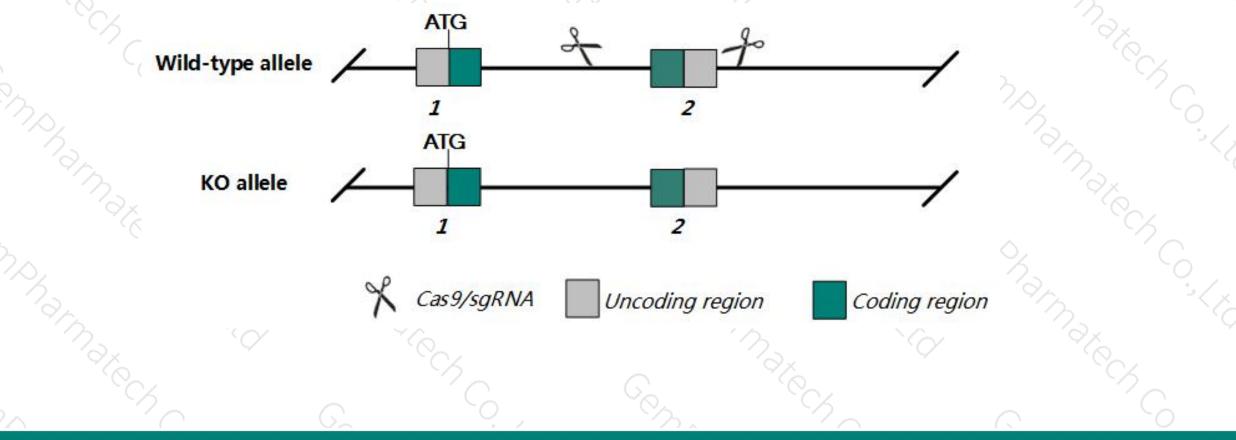






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This model will use CRISPR/Cas9 technology to edit the Crp gene. The schematic diagram is as follows:



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- The Crp gene has 10 transcripts. According to the structure of Crp gene, exon2- of Crp-201 (ENSMUST00000038495.4) 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 *Crp* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

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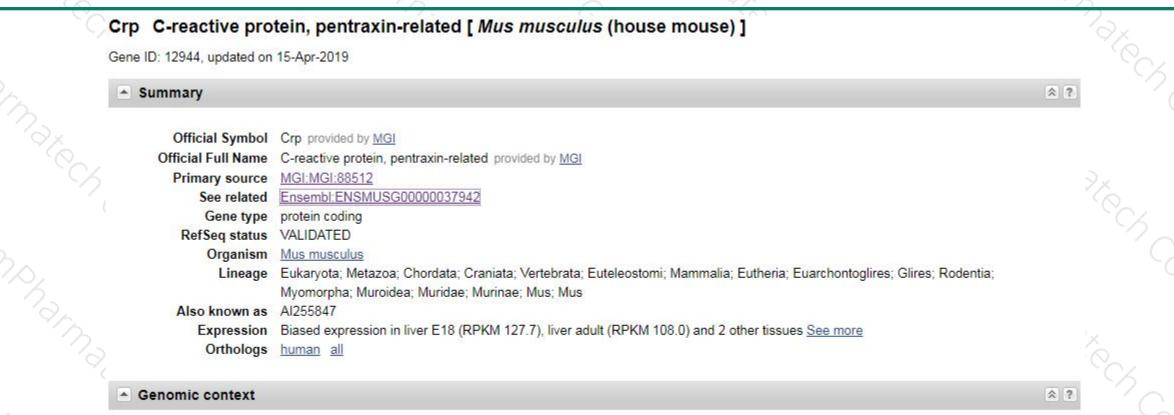
- According to the existing MGI data, Mice homozygous for a knock-out allele lack detectable C-reactive protein in the serum but are otherwise healthy and fertile.
- The Crp gene is located on the Chr1. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous 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.

Notice

# **Gene information (NCBI)**



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Location: 1 H3; 1 80.13 cM

See Crp in Genome Data Viewer

Exon count: 2

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Annotation release	Status	Assembly	Chr	Location	
106	current	GRCm38.p4 (GCF_000001635.24)	1	NC_000067.6 (172698056172699966)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	1	NC_000067.5 (174628187174630097)	

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# **Transcript information (Ensembl)**



The gene has 2 transcripts, all transcripts are shown below:

Name 🖕	Transcript ID	bp 🖕	Protein 🖕	Biotype 🖕	CCDS 🖕	UniProt 🖕	Flags			
Crp-201	ENSMUST0000038495.4	1695	<u>225aa</u>	Protein coding	<u>CCDS35787</u> @	P14847@	TSL:2	GENCODE basic	APPRIS P1	
Crp-202	ENSMUST00000194251.1	1894	No protein	Processed transcript		-		TSL:1		

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

			1.91 kb —			Forward strand
Crp-201 > protein coding						
2	. 0	02	×. /	125.	10	Q

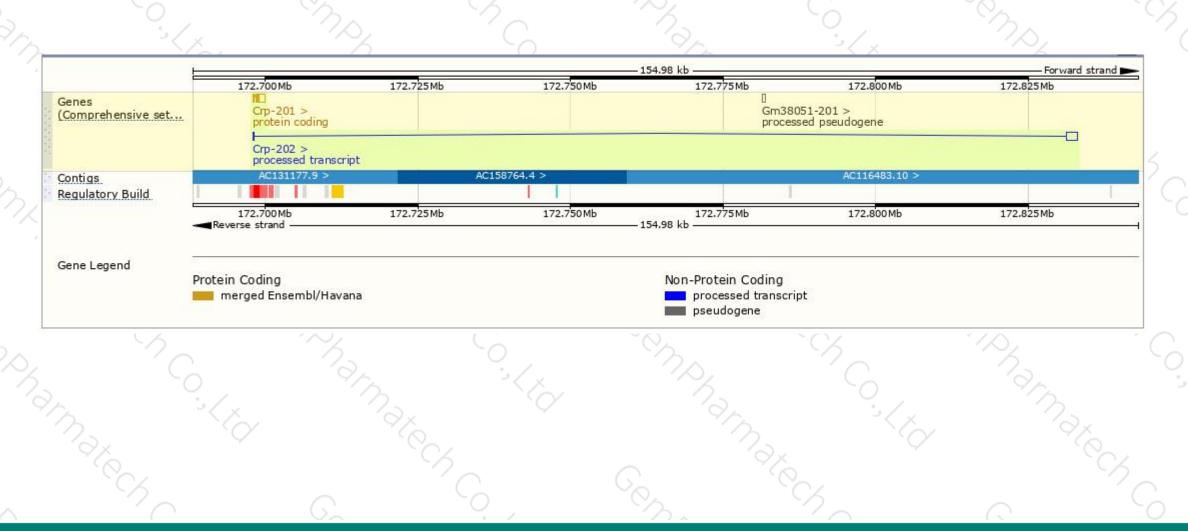
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## **Genomic location distribution**





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## **Protein domain**



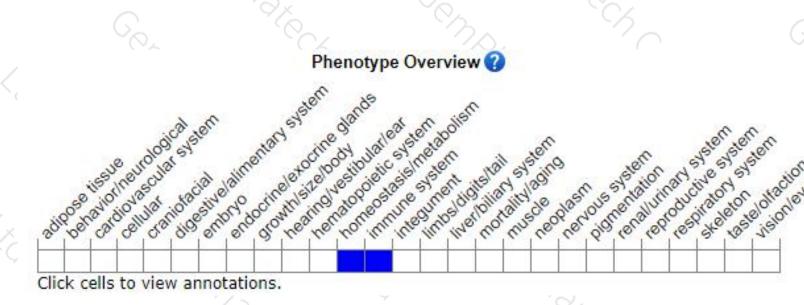
ENCHUG DODDODD	-										
ENSMUSP0000044 Conserved Domains											
Cleavage site (Sign hmmpanther	PTHR19277:SF119										_
	PTHR19277										
Superfamily domains	G	oncanavalin A-like leo	tin/glucanase doma	in superfamily							
SMART domains	Per	ntraxin-related									
<ul> <li>Prints domain</li> <li>Pfam domain</li> </ul>			Pentraxin-related								
PROSITE profiles		Pentraxin-related									
PROSITE patterns		Pentraxin-related				-	1.24				
Gene3D		0.120,200				Pentaxin, conse	erved site				
		s (dbSNP and all ot	her sources)								
All sequence SNPs/i					1		-		10.11		
Variant Legend	missense va	riant				synonym	ous variant				
Scale bar	0 20	40	60	80	100	120	140	160	180	200	22
				1.1		70.		0		15	
				$\sim$ $\times$							

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## Mouse phenotype description(MGI)





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

Mice homozygous for a knock-out allele lack detectable C-reactive protein in the serum but are otherwise healthy and fertile.

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If you have any questions, you are welcome to inquire. Tel: 025-5864 1534



