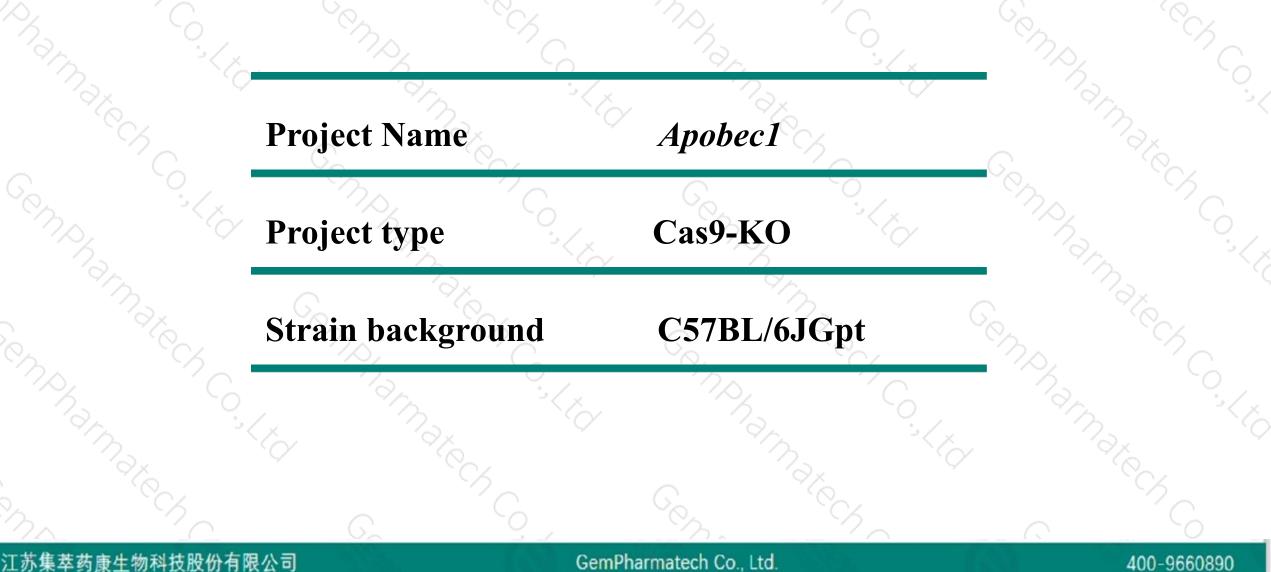


# Apobec1 Cas9-KO Strategy

Designer:Xueting Zhang Reviewer:Yanhua Shen Date:2019-11-13

## **Project Overview**



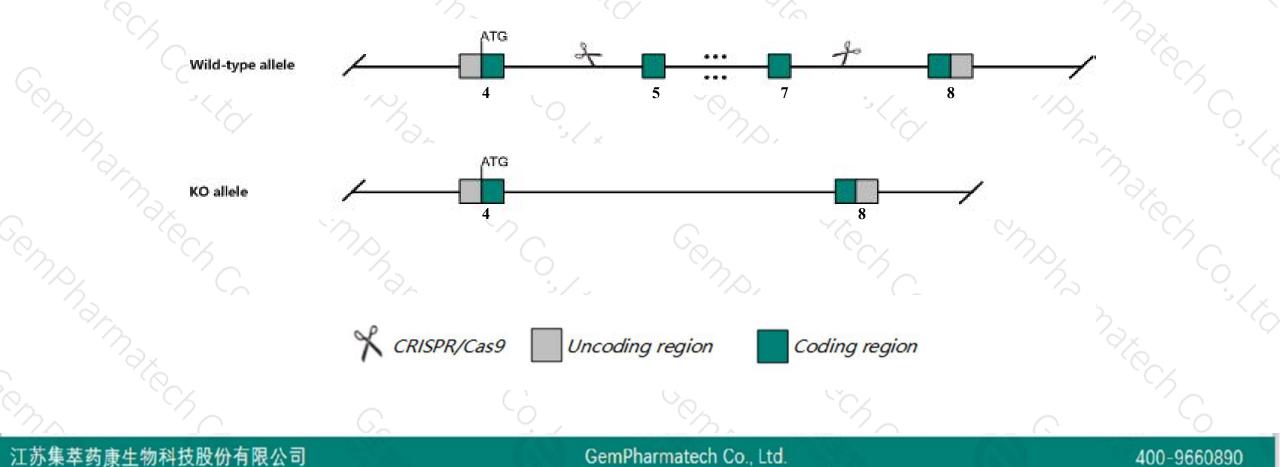


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# **Knockout** strategy



This model will use CRISPR/Cas9 technology to edit the *Apobec1* gene. The schematic diagram is as follows:





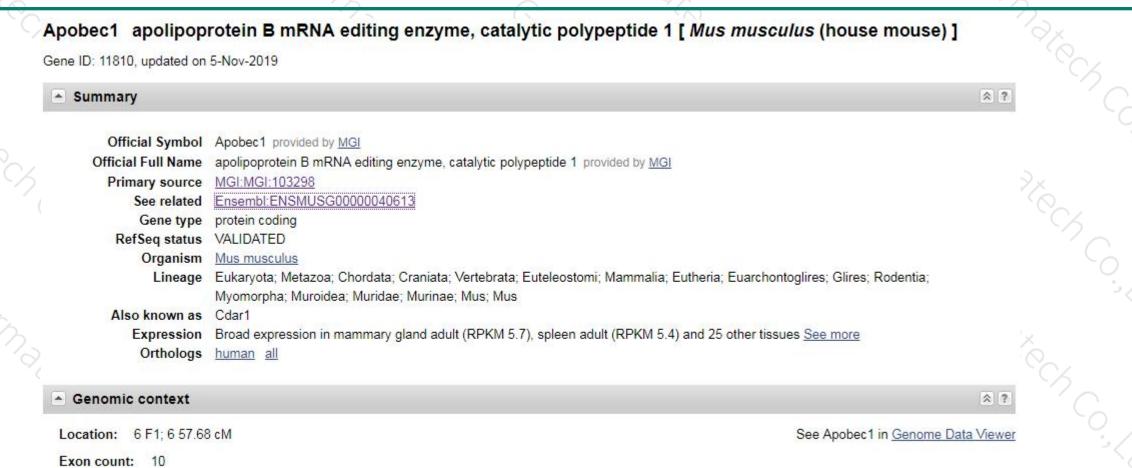
- The Apobec1 gene has 9 transcripts. According to the structure of Apobec1 gene, exon5-exon7 of Apobec1-202 (ENSMUST00000112586.7) transcript is recommended as the knockout region. The region contains 545bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Apobec1 gene. The brief process is as follows: CRISPR/Cas9 systemeters are appreciately ap

- > According to the existing MGI data, Mice homozygous for a null allele exhibit abnormal lipid homeostasis.
- ➤ Transcript *Apobec1*-209 may not be affected.
- The Apobec1 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Notice

# **Gene information** (NCBI)





Annotation release Status		Assembly		Location			
108	current	GRCm38.p6 (GCF_000001635.26)	6	NC_000072.6 (122577792122603024, complement)			
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	6	NC_000072.5 (122527810122552462, complement)			

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# **Transcript information (Ensembl)**



#### The gene has 9 transcripts, all transcripts are shown below:

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Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Apobec1-202	ENSMUST00000112586.7	2265	<u>229aa</u>	Protein coding	CCDS20498	P51908 Q3U9G8	TSL:1 GENCODE basic APPRIS P1
Apobec1-201	ENSMUST00000112585.7	2127	<u>229aa</u>	Protein coding	CCDS20498	P51908 Q3U9G8	TSL:1 GENCODE basic APPRIS P1
Apobec1-203	ENSMUST00000112587.10	2072	<u>229aa</u>	Protein coding	CCDS20498	P51908 Q3U9G8	TSL:5 GENCODE basic APPRIS P1
Apobec1-207	ENSMUST00000203204.2	775	<u>113aa</u>	Protein coding	4	A0A0N4SVL6	CDS 3' incomplete TSL:2
Apobec1-208	ENSMUST00000203309.2	666	<u>121aa</u>	Protein coding	10	A0A0N4SW85	CDS 3' incomplete TSL:2
Apobec1-206	ENSMUST00000203197.2	372	<u>43aa</u>	Protein coding	÷	A0A0N4SV86	CDS 3' incomplete TSL:3
Apobec1-205	ENSMUST00000147760.7	360	<u>25aa</u>	Protein coding		D3Z675	CDS 3' incomplete TSL:5
Apobec1-204	ENSMUST00000143356.1	560	No protein	Retained intron	4	<u>42</u>	TSL:3
Apobec1-209	ENSMUST00000204035.1	520	No protein	Retained intron	5	65	TSL:3
		2				5 X	

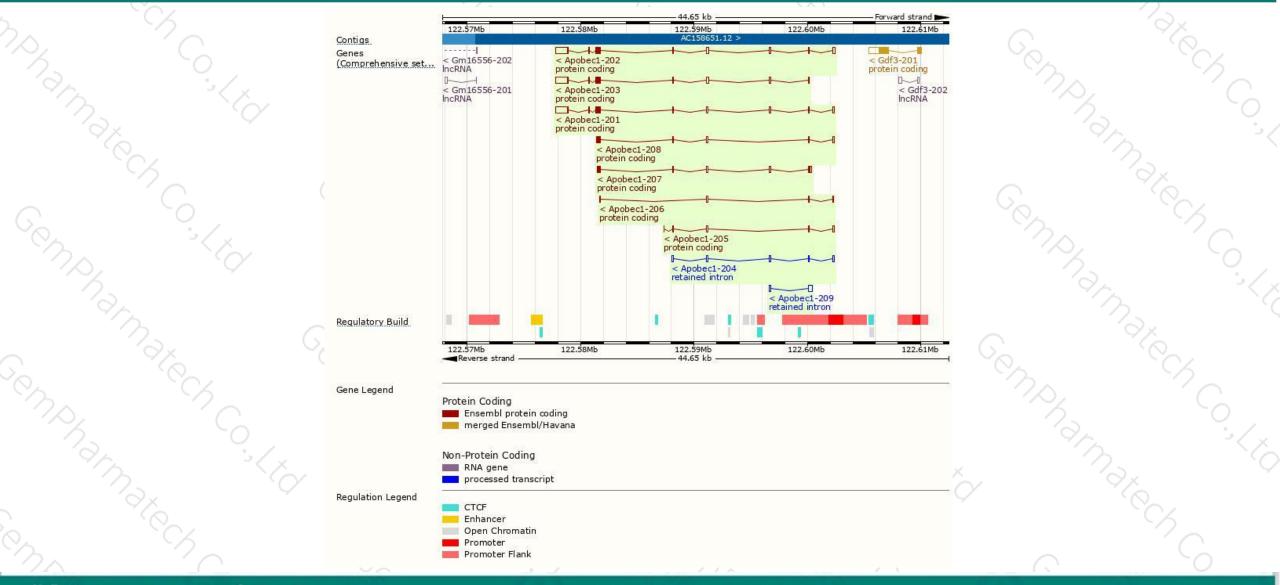
The strategy is based on the design of Apobec1-202 transcript, The transcription is shown below



- 24.65 kb -

#### **Genomic location distribution**





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## **Protein domain**



ENSMUSP00000108							
Superfamily Steen		Cytidine dean					
Pfam	C->U-/	editing enzyme APOBE	C-1				
PROSITE profiles	Cytidine and de	oxycytidylate deamina:	se domain				
PROSITE patterns	3	APOBEC/	CMP deaminase, zinc	-binding			
PANTHER	PTHR13857						C
	PTHR13857:SF26						
Gene3D	3.40.140.10			5.1			
	cd01283						
All sequence SNPs/i	Sequence variants	(dbSNP and all other	sources)			1	
Variant Legend							
	📒 missense vari						C.
	splice region synonymous						
Scale bar	0 20	40 60	80 100	120 140	160	180	200 229
1.0°C4							$\sim 2$

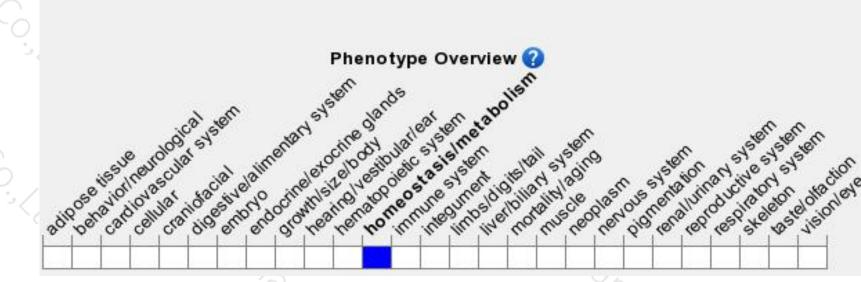
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## Mouse phenotype description(MGI)





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

According to the existing MGI data, Mice homozygous for a null allele exhibit abnormal lipid homeostasis.



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



