

# Il34 Cas9-KO Strategy

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**Reviewer:** Yang Zeng

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# **Project Overview**



Project Name Il34

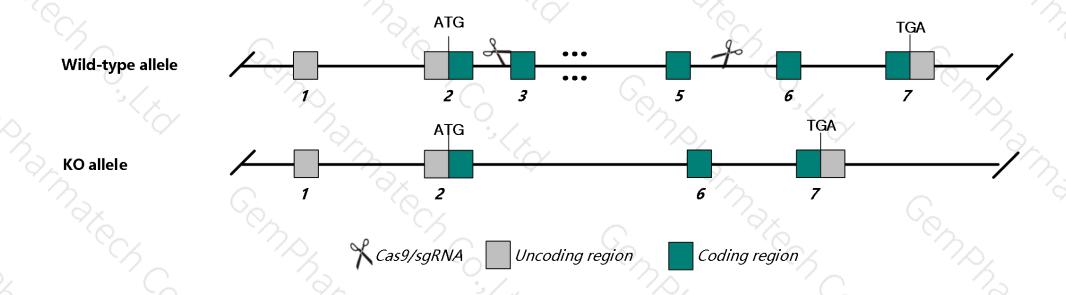
Project type Cas9-KO

Strain background C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Il34* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Il34* gene has 5 transcripts. According to the structure of *Il34* gene, exon3-exon5 of *Il34-201*(ENSMUST0000076846.10) transcript is recommended as the knockout region. The region contains 374bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Il34* gene. The brief process is as follows: CRISPR/Cas9 system 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.

### **Notice**



- > According to the existing MGI data, Mice homozygous for a knock-out allele exhibit reduced Langerhans cells and microglial cells in the skin and brain, respectively, with decreased susceptibility to type IV hypersensitivity reaction and fungal infection but increased susceptibility to viral infection.
- Transcript *Il34-202/203/205* may not be affected. The KO region is close to 5'UTR region of the *Gm15894* gene. Knockout the region may affect the regulation of *Gm15894* gene.
- The *Il34* gene is located on the Chr8. 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.

### Gene information (NCBI)



#### II34 interleukin 34 [ Mus musculus (house mouse) ]

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☆ ?

Gene ID: 76527, updated on 3-Aug-2021



Official Symbol II34 provided by MGI

Official Full Name interleukin 34 provided by MGI

Primary source MGI:MGI:1923777

See related Ensembl: ENSMUSG00000031750

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 Al593503; 2010004A03Rik

Expression Broad expression in adrenal adult (RPKM 23.9), frontal lobe adult (RPKM 16.7) and 18 other tissues See more

Orthologs <u>human</u> all



Try the new Gene table

Try the new Transcript table

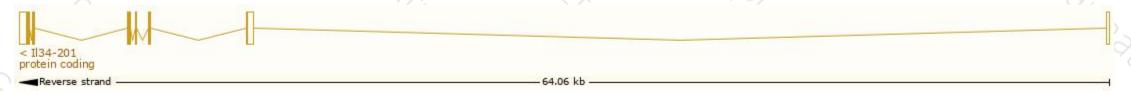
# Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

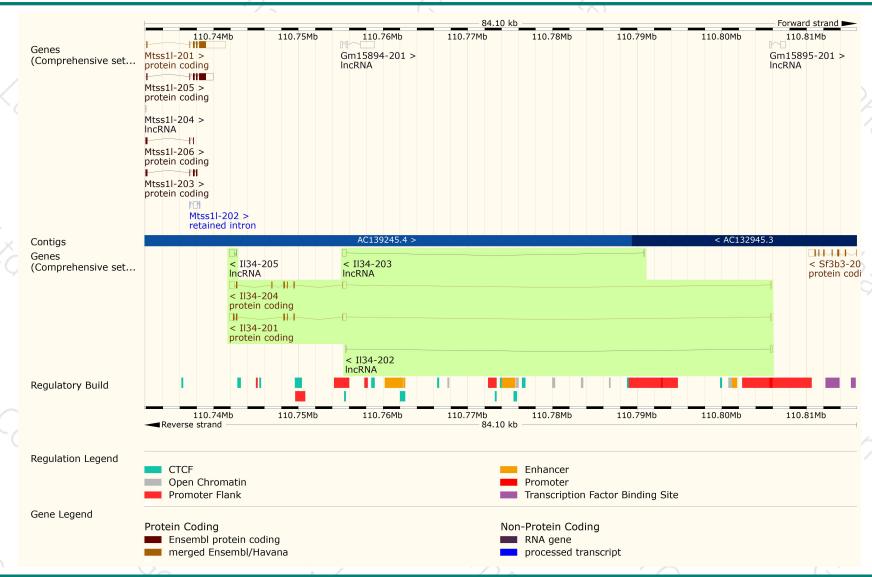
Name 🍦	Transcript ID	bp 🛊	Protein	Biotype	CCDS	UniProt Match	Flags
1134-204	ENSMUST00000150680.2	1796	219aa	Protein coding	CCDS52670 ₺	Q8R1R4-2₽	GENCODE basic   TSL:1
1134-201	ENSMUST00000076846.11	1709	<u>235aa</u>	Protein coding	CCDS52669 ₽	Q8R1R4-1@	GENCODE basic APPRIS P1 TSL:1
1134-205	ENSMUST00000154803.2	675	No protein	Processed transcript	-	-	TSL:3
1134-203	ENSMUST00000148732.2	515	No protein	Processed transcript	15	107	TSL:3
1134-202	ENSMUST00000137362.2	309	No protein	Processed transcript	-	-	TSL:3

The strategy is based on the design of *Il34-201* transcript, The transcription is shown below



### Genomic location distribution





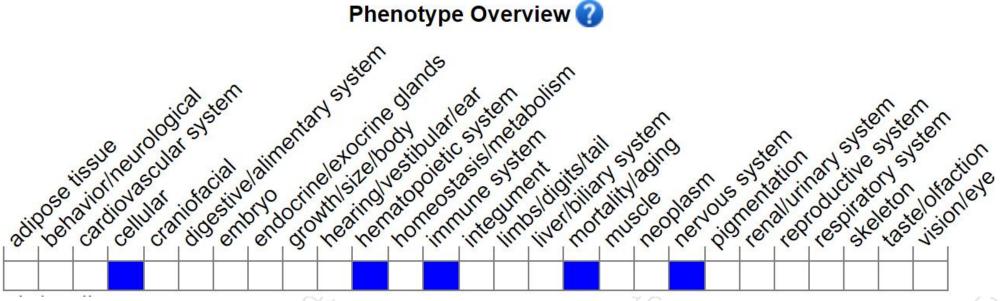
## Protein domain





## 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 knock-out allele exhibit reduced Langerhans cells and microglial cells in the skin and brain, respectively, with decreased susceptibility to type IV hypersensitivity reaction and fungal infection but increased susceptibility to viral infection.



If you have any questions, you are welcome to inquire. Tel: 400-966 0890





