

Il33 Cas9-KO Strategy

Designer: QiongZhou

Project Overview

Project Name

Il33

Project type

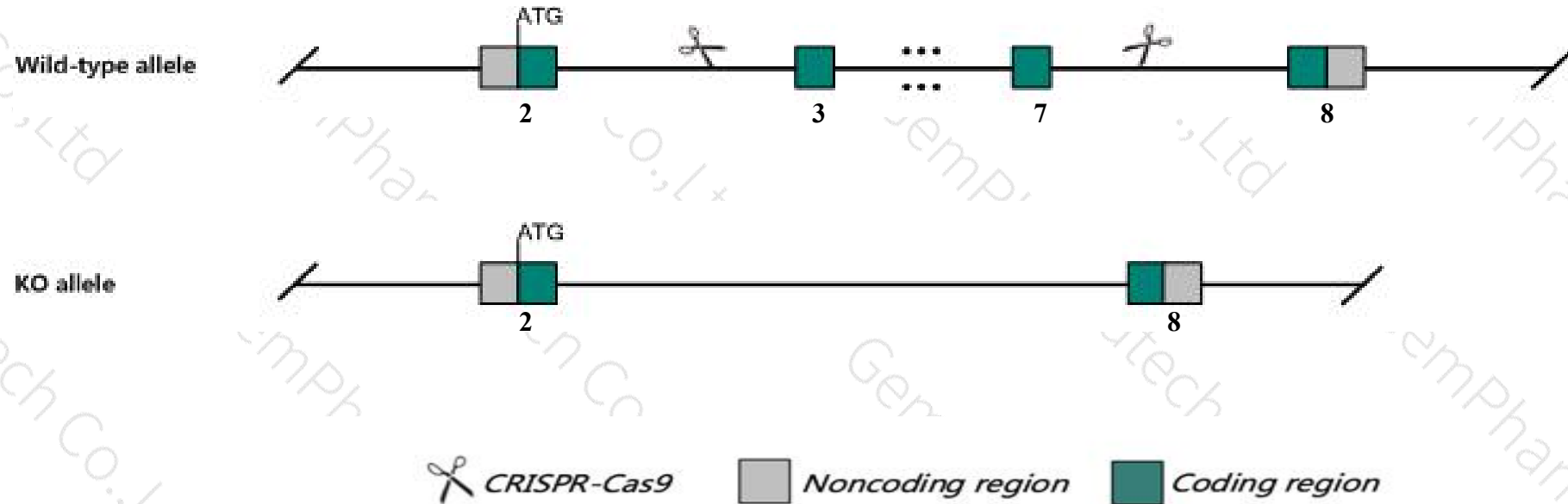
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Il33* gene. The schematic diagram is as follows:



- The *Il33* gene has 5 transcripts. According to the structure of *Il33* gene, exon3-exon7 of *Il33-201* (ENSMUST00000025724.8) transcript is recommended as the knockout region. The region contains 512bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Il33* gene. The brief process is as follows: gRNA was transcribed in vitro. Cas9 and gRNA 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.

- According to the existing MGI data, Nullizygous mutations lead to altered Type 2 immunity and increased susceptibility to parasite infection. Homozygotes for a null allele show accelerated ovarian functional decline and early reproductive aging due to impaired migration of ovarian macrophages and failed disposal of atretic follicles.
- The *Il33* gene is located on the Chr19. 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)

Il33 interleukin 33 [Mus musculus (house mouse)]

Gene ID: 77125, updated on 19-Mar-2019

Summary



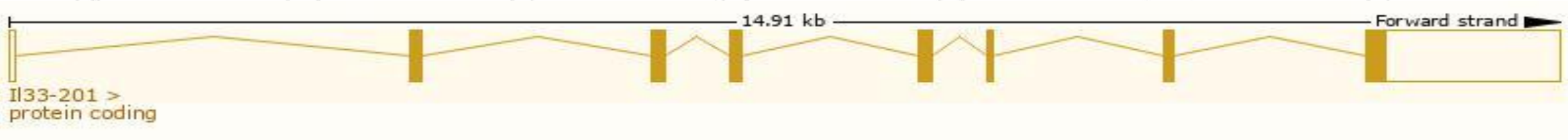
| | |
|---------------------------|---|
| Official Symbol | Il33 provided by MGI |
| Official Full Name | interleukin 33 provided by MGI |
| Primary source | MGI:MGI:1924375 |
| See related | Ensembl:ENSMUSG00000024810 |
| Gene type | protein coding |
| RefSeq status | VALIDATED |
| Organism | Mus musculus |
| Lineage | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| Also known as | 9230117N10Rik, Il-33, Il1f11, NF-HEV |
| Expression | Broad expression in lung adult (RPKM 13.9), cerebellum adult (RPKM 10.6) and 19 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

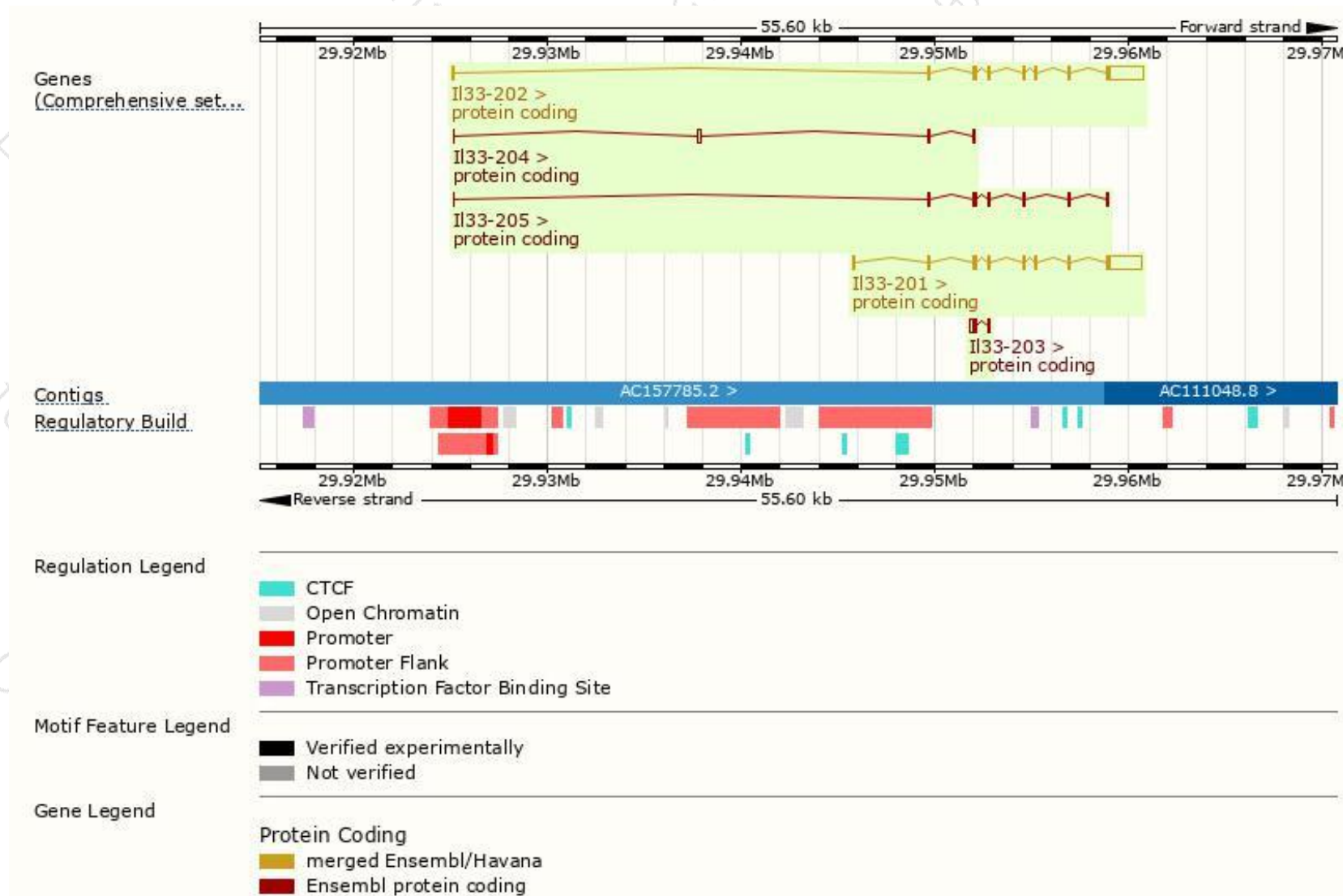
The gene has 5 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|----------|--------------------------------------|------|-----------------------|----------------|---------------------------|------------------------|-------------------------------|
| II33-201 | ENSMUST00000025724.8 | 2544 | 266aa | Protein coding | CCDS29740 | Q8BVZ5 | TSL:1 GENCODE basic APPRIS P1 |
| II33-202 | ENSMUST00000120388.8 | 2537 | 266aa | Protein coding | CCDS29740 | Q8BVZ5 | TSL:1 GENCODE basic APPRIS P1 |
| II33-205 | ENSMUST00000177518.7 | 725 | 228aa | Protein coding | - | H3BLN5 | CDS 3' incomplete TSL:3 |
| II33-204 | ENSMUST00000144528.7 | 377 | 66aa | Protein coding | - | D3Z6T7 | CDS 3' incomplete TSL:3 |
| II33-203 | ENSMUST00000136850.1 | 370 | 83aa | Protein coding | - | H3BKB5 | CDS 3' incomplete TSL:2 |

The strategy is based on the design of *II33-201* transcript,The transcription is shown below



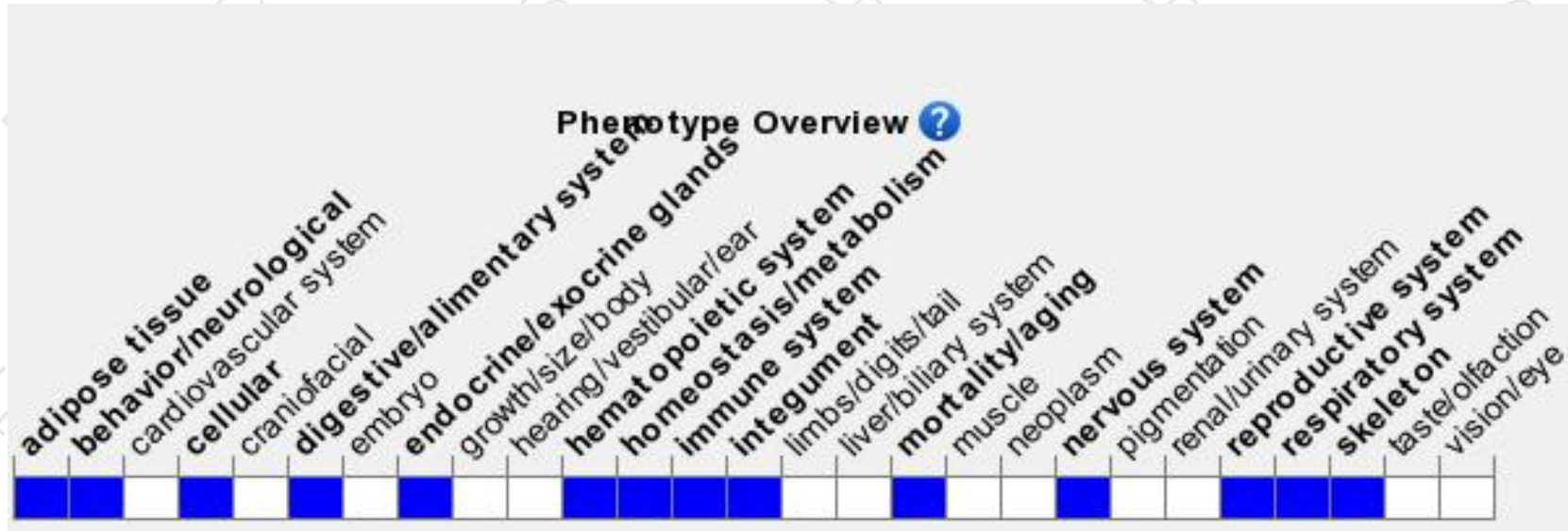
Genomic location distribution



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, Nullizygous mutations lead to altered Type 2 immunity and increased susceptibility to parasite infection. Homozygotes for a null allele show accelerated ovarian functional decline and early reproductive aging due to impaired migration of ovarian macrophages and failed disposal of atretic follicles.

If you have any questions, you are welcome to inquire.

Tel:400-9660890

