

Atp11b Cas9-KO Strategy

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Design Date: 2020-11-10

Project Overview

Project Name

Atp11b

Project type

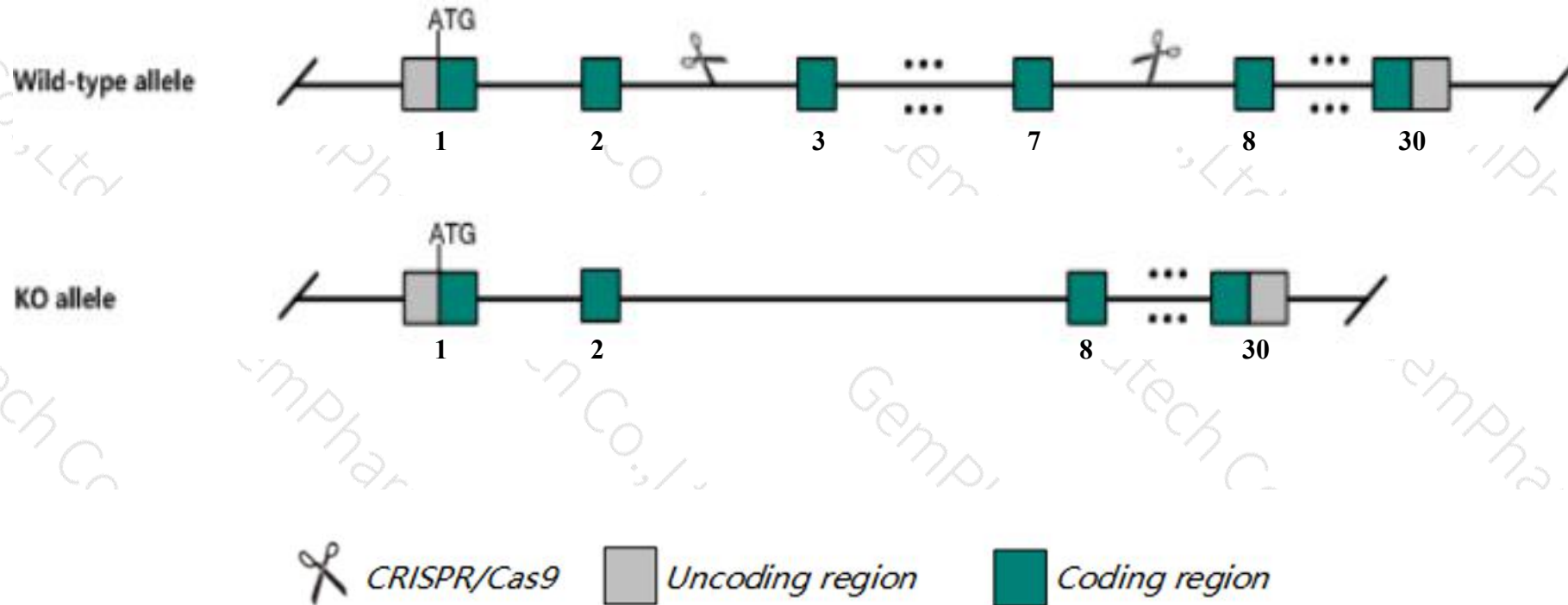
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Atp11b* gene. The schematic diagram is as follows:



- The *Atp11b* gene has 11 transcripts. According to the structure of *Atp11b* gene, exon3-exon7 of *Atp11b-201*(ENSMUST00000029257.14) 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 *Atp11b* 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.

- The *Atp11b* gene is located on the Chr3. 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 may not affect the *Atp11b*-206 transcript.
- 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)

Atp11b ATPase, class VI, type 11B [Mus musculus (house mouse)]

Gene ID: 76295, updated on 20-Mar-2020

Summary



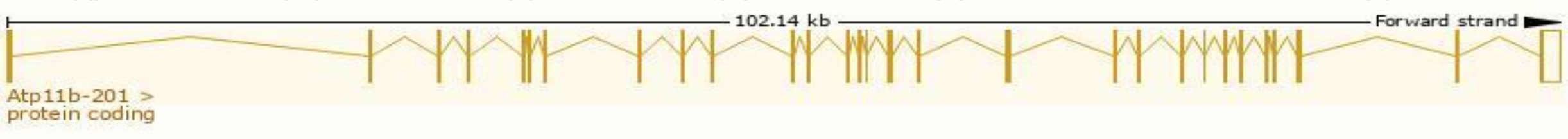
| | |
|---------------------------|---|
| Official Symbol | Atp11b provided by MGI |
| Official Full Name | ATPase, class VI, type 11B provided by MGI |
| Primary source | MGI:MGI:1923545 |
| See related | Ensembl:ENSMUSG00000037400 |
| Gene type | protein coding |
| RefSeq status | VALIDATED |
| Organism | Mus musculus |
| Lineage | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| Also known as | 1110019I14Rik, ATPIF, ATPIR, mKIAA0956 |
| Expression | Ubiquitous expression in frontal lobe adult (RPKM 18.5), CNS E18 (RPKM 15.8) and 28 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

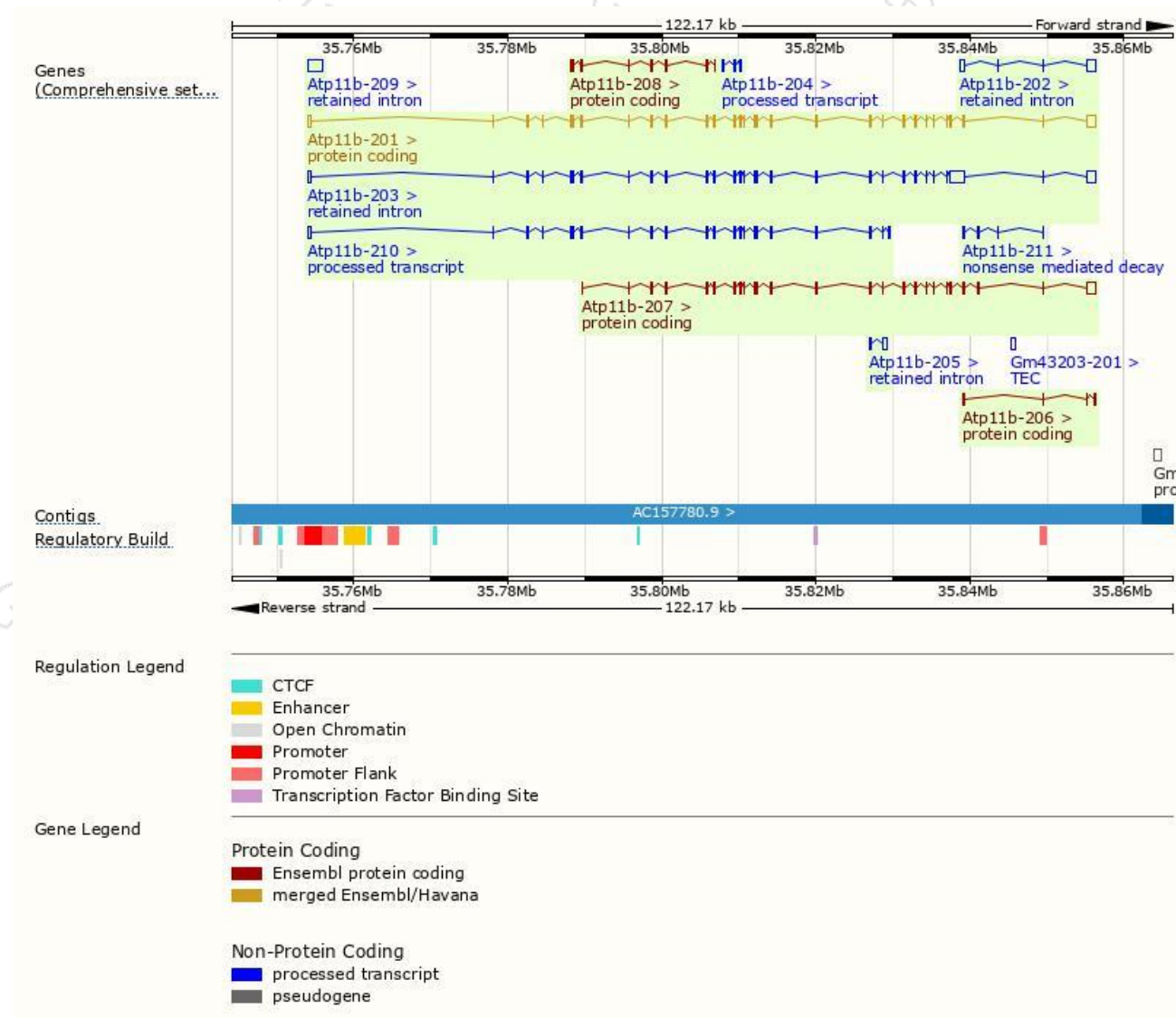
The gene has 11 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|------------|---------------------------------------|------|------------------------|-------------------------|---------------------------|----------------------------|--------------------------------|
| Atp11b-201 | ENSMUST00000029257.14 | 4868 | 1175aa | Protein coding | CCDS38414 | Q6DFW5 | TSL:1 GENCODE basic APPRIS P1 |
| Atp11b-207 | ENSMUST00000198599.1 | 4102 | 967aa | Protein coding | - | A0A0G2JE89 | CDS 5' incomplete TSL:5 |
| Atp11b-208 | ENSMUST00000199892.4 | 776 | 259aa | Protein coding | - | A0A0G2JED0 | CDS 5' and 3' incomplete TSL:3 |
| Atp11b-206 | ENSMUST00000197764.1 | 621 | 124aa | Protein coding | - | A0A0G2JF19 | CDS 5' incomplete TSL:3 |
| Atp11b-211 | ENSMUST00000211902.1 | 509 | 112aa | Nonsense mediated decay | - | A0A1D5RLK3 | CDS 5' incomplete TSL:5 |
| Atp11b-210 | ENSMUST00000200445.4 | 3028 | No protein | Processed transcript | - | - | TSL:1 |
| Atp11b-204 | ENSMUST00000196965.1 | 648 | No protein | Processed transcript | - | - | TSL:5 |
| Atp11b-203 | ENSMUST00000196700.4 | 6322 | No protein | Retained intron | - | - | TSL:5 |
| Atp11b-202 | ENSMUST00000196409.1 | 1918 | No protein | Retained intron | - | - | TSL:5 |
| Atp11b-209 | ENSMUST00000200351.1 | 1906 | No protein | Retained intron | - | - | TSL:NA |
| Atp11b-205 | ENSMUST00000197003.1 | 638 | No protein | Retained intron | - | - | TSL:3 |

The strategy is based on the design of *Atp11b-201* transcript,the transcription is shown below:



Genomic location distribution



Protein domain



集萃药康
GemPharmatech



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

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