

# ***Ammecr1* Cas9-KO Strategy**

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**Design Date: 2020-9-16**

# Project Overview

**Project Name**

*Ammecr1*

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Ammecr1* gene. The schematic diagram is as follows:



- The *Ammecr1* gene has 1 transcript. According to the structure of *Ammecr1* gene, exon3 of *Ammecr1*-201(ENSMUST00000041317.2) transcript is recommended as the knockout region. The region contains 115bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Ammecr1* 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, male chimeras hemizygous for a gene trapped allele appear normal at E9.5.
- The N-terminal of *Ammecr1* gene will remain several amino acids, it may remain the partial function of *Ammecr1* gene.
- The *Ammecr1* gene is located on the ChrX. 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)

## Ammecr1 Alport syndrome, mental retardation, midface hypoplasia and elliptocytosis chromosomal region gene 1 [Mus musculus (house mouse)]

Gene ID: 56068, updated on 13-Mar-2020

### Summary



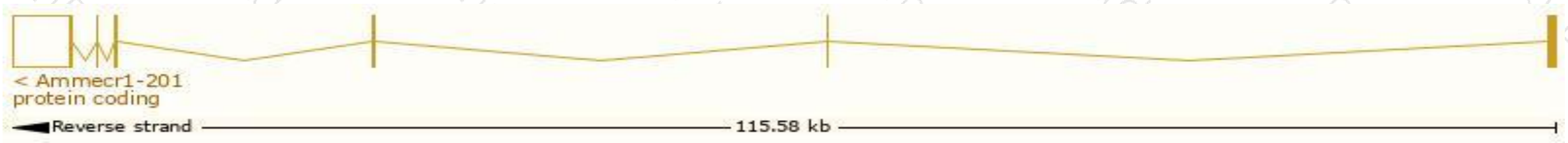
<b>Official Symbol</b>	Ammecr1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	Alport syndrome, mental retardation, midface hypoplasia and elliptocytosis chromosomal region gene 1 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1860206</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000042225</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	6230420G18Rik
<b>Expression</b>	Ubiquitous expression in liver E14 (RPKM 3.7), liver E14.5 (RPKM 3.0) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

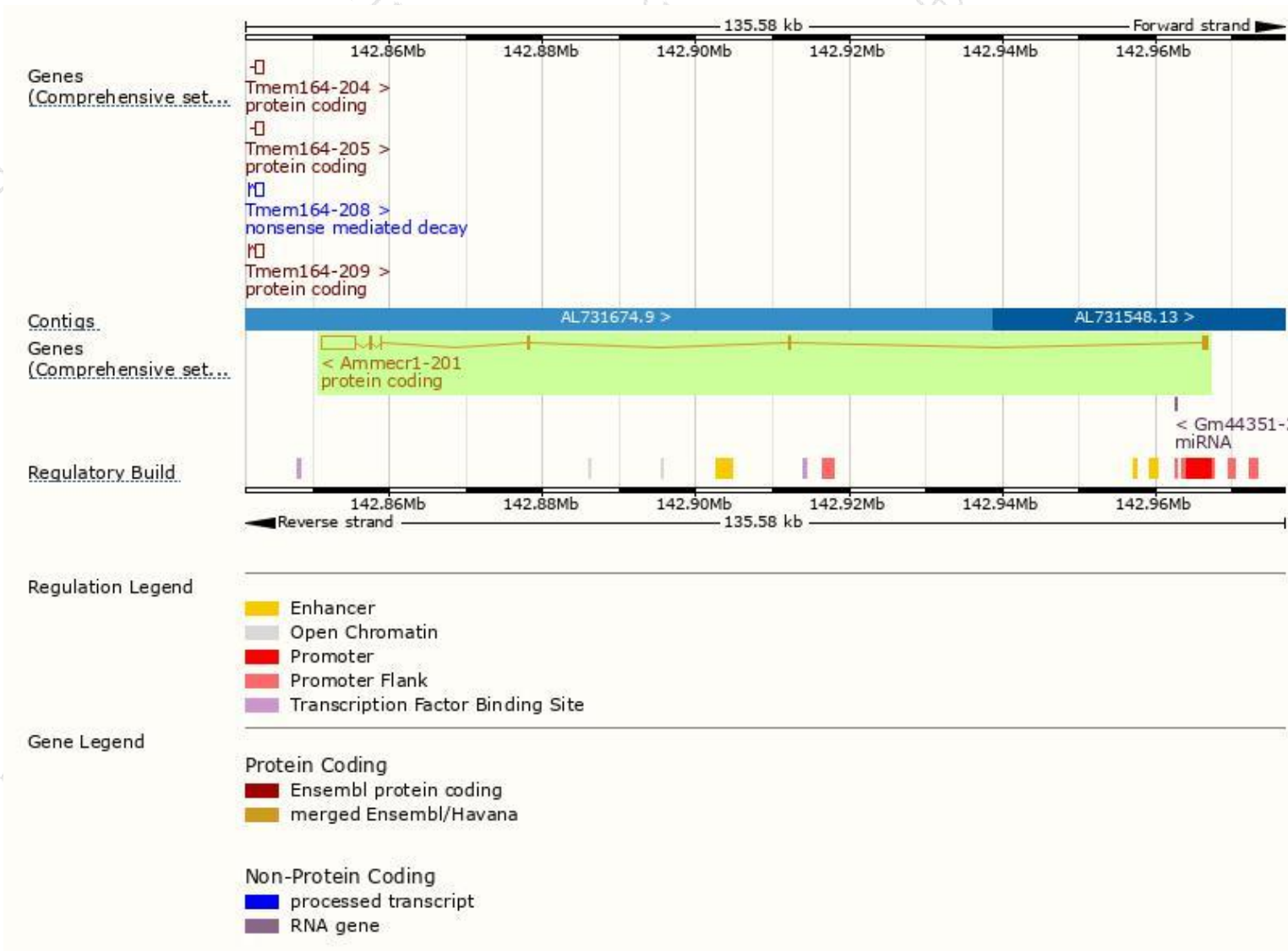
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
<b>Ammecr1-201</b>	<a href="#">ENSMUST00000041317.2</a>	5437	<a href="#">344aa</a>	Protein coding	<a href="#">CCDS30452</a>	<a href="#">Q9JHT5</a>	TSL:1 GENCODE basic APPRIS P1

The strategy is based on the design of *Ammecr1-201* transcript, the transcription is shown below:

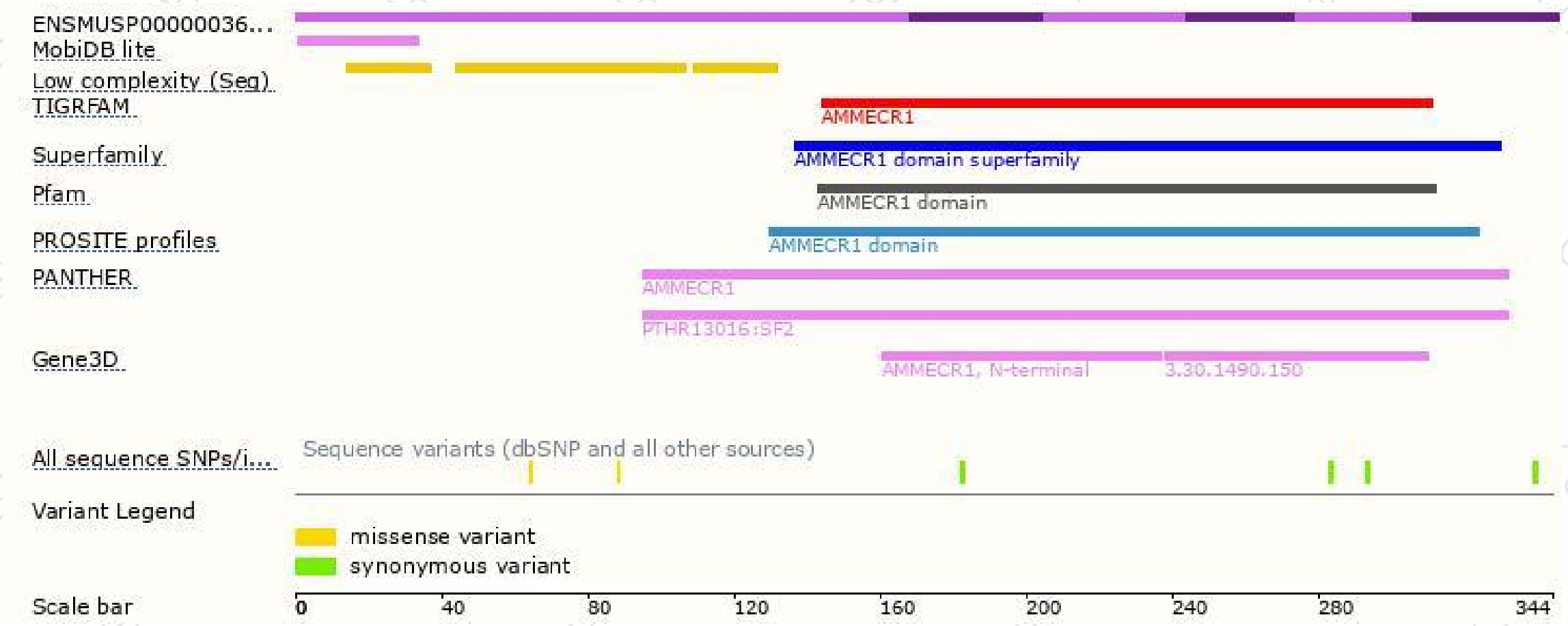


# Genomic location distribution





# Protein domain



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

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