



Amer1 Cas9-CKO Strategy

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Reviewer:

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Design Date:

2019-11-26

Project Overview

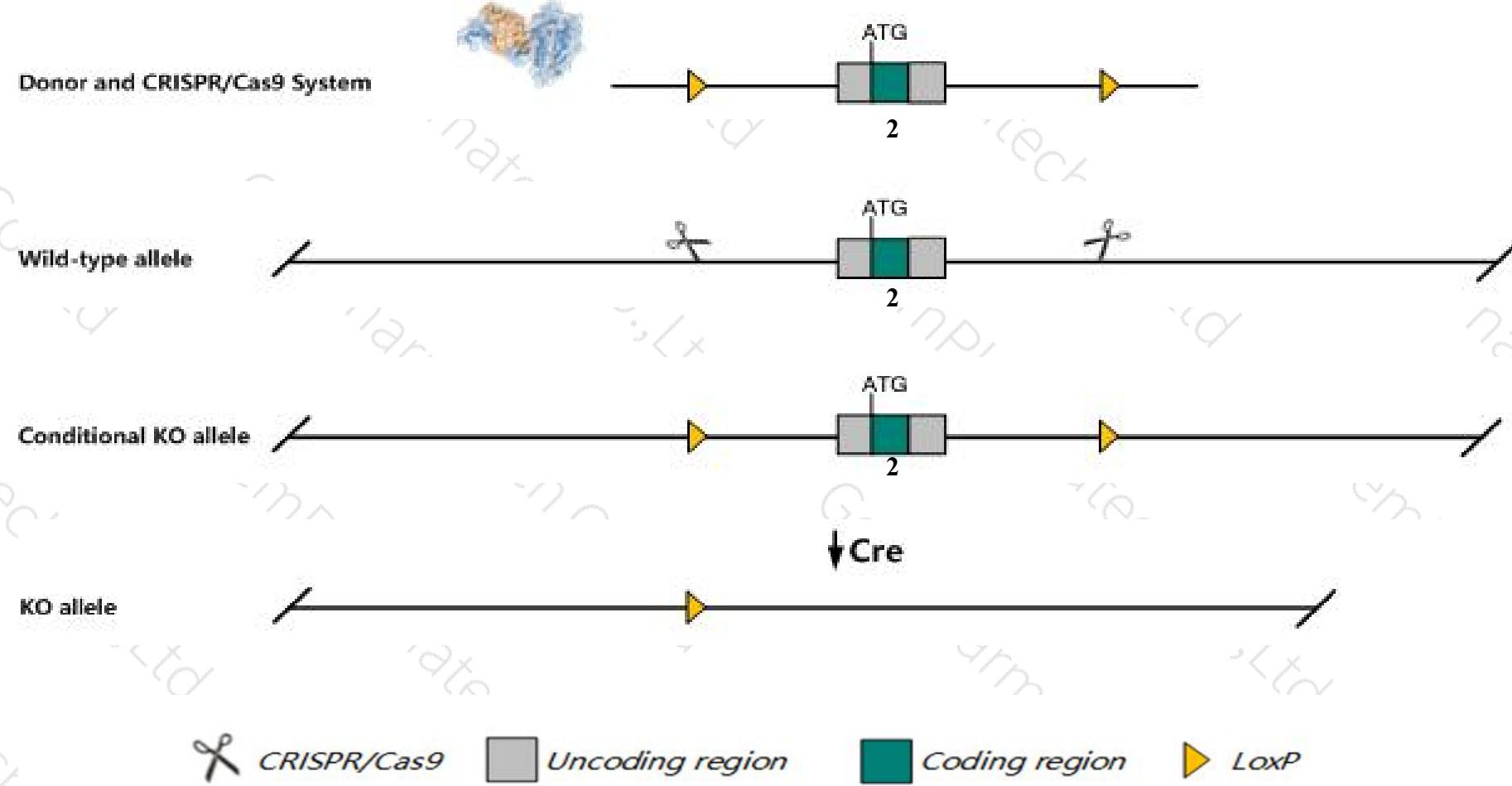
Project Name*Amer1*

Project type**Cas9-CKO**

Strain background**C57BL/6JGpt**

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Amer1* gene. The schematic diagram is as follows:



Technical routes

- The *Amer1* gene has 1 transcript. According to the structure of *Amer1* gene, exon2 of *Amer1-201* (ENSMUST00000084535.5) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Amer1* gene. The brief process is as follows:CRISPR/Cas9 system and Donor 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 flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.



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Notice

- According to the existing MGI data, Male mice hemizygous for a null mutation display neonatal lethality with cardiac overgrowth, bone overgrowth, bilateral or unilateral renal agenesis coupled with renal overgrowth, adipocyte and spleen hypoplasia, and altered mesenchymal progenitor cell fate specification.
- The *Amer1* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.



Gene information (NCBI)

Amer1 APC membrane recruitment 1 [Mus musculus (house mouse)]

Gene ID: 72345, updated on 31-Jan-2019

Summary



Official Symbol Amer1 provided by [MGI](#)

Official Full Name APC membrane recruitment 1 provided by [MGI](#)

Primary source [MGI:MGI:1919595](#)

See related [Ensembl:ENSMUSG00000050332](#)

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 2810002O09Rik, AW492303, Fam123b, Wtx

Expression Ubiquitous expression in bladder adult (RPKM 3.7), limb E14.5 (RPKM 3.4) and 28 other tissues [See more](#)

Orthologs [human](#) [all](#)

Transcript information (Ensembl)

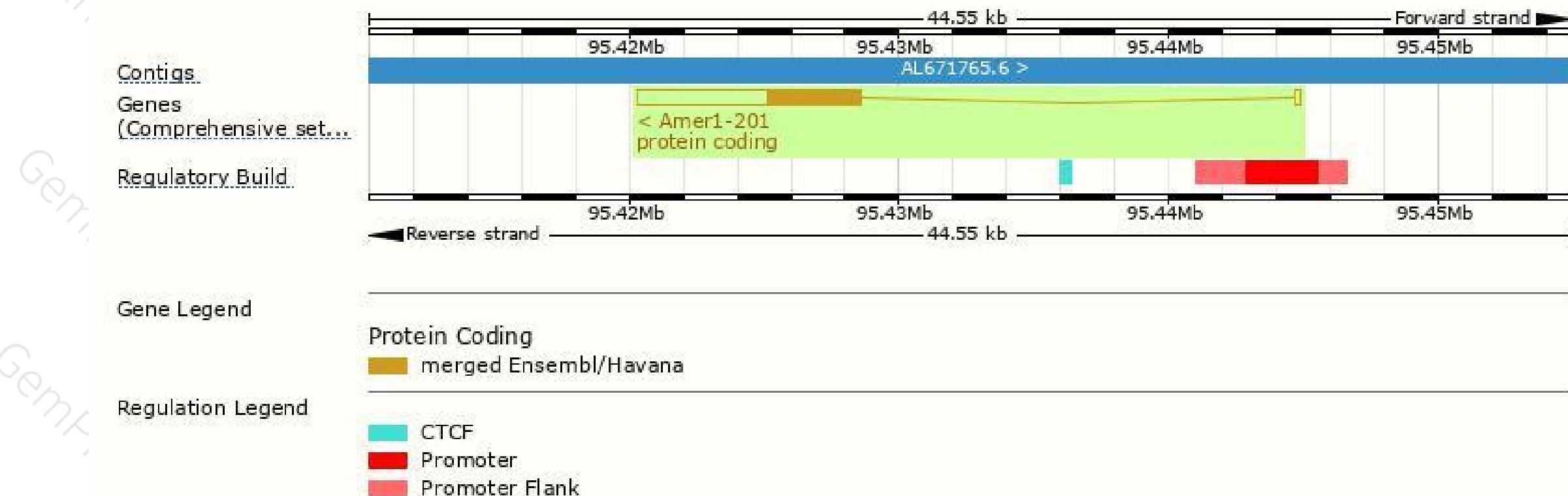
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Amer1-201	ENSMUST00000084535.5	8496	1132aa	Protein coding	CCDS41067	Q7TS75	TSL:1 GENCODE basic APPRIS P1

The strategy is based on the design of *Amer1-201* transcript. The transcription is shown below



Genomic location distribution



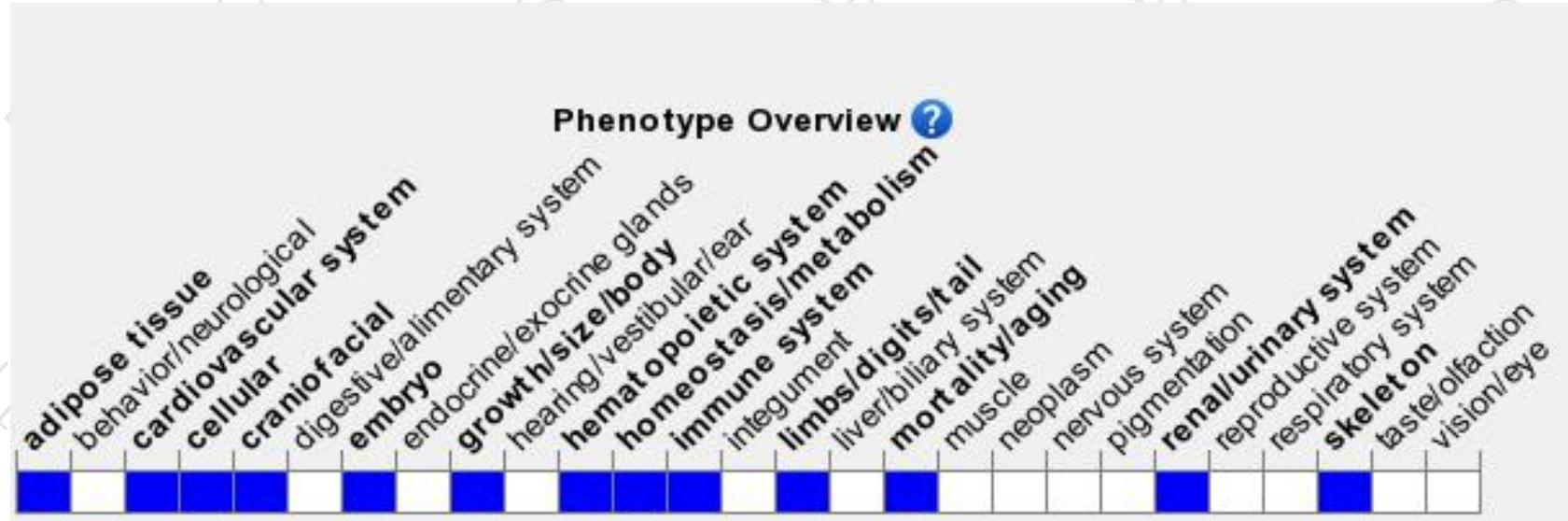
Protein domain





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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, Male mice hemizygous for a null mutation display neonatal lethality with cardiac overgrowth, bone overgrowth, bilateral or unilateral renal agenesis coupled with renal overgrowth, adipocyte and spleen hypoplasia, and altered mesenchymal progenitor cell fate specification.



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

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