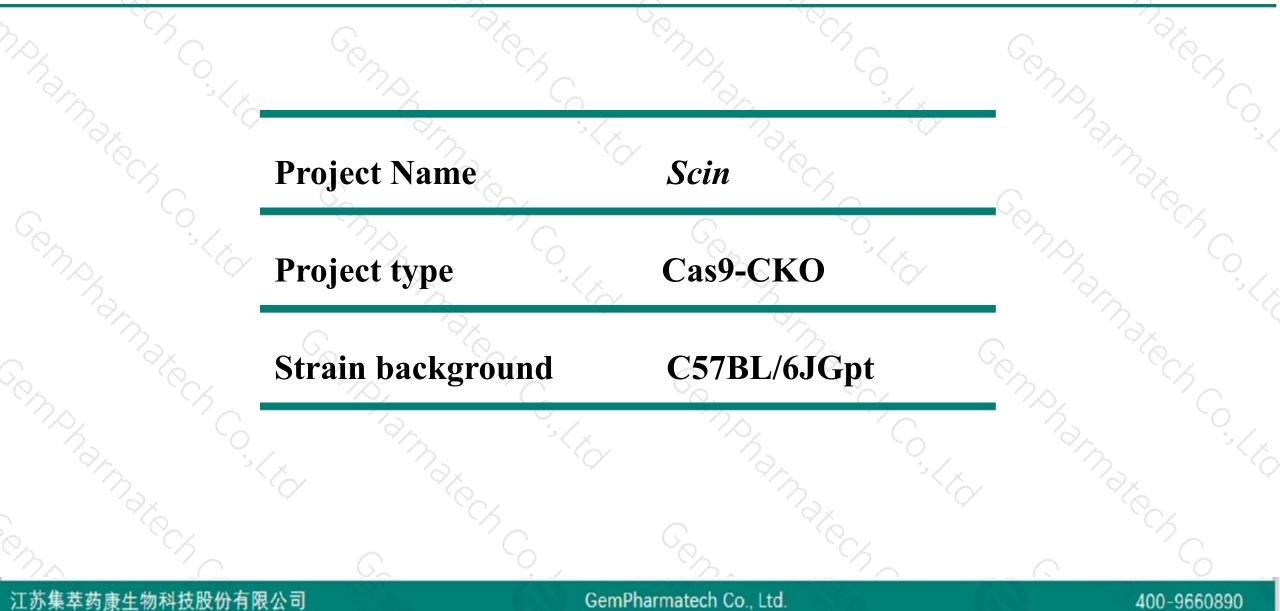


Scin Cas9-CKO Strategy

Designer: Reviewer: Design Date: JiaYu Xiaojing Li 2020-2-11

Project Overview



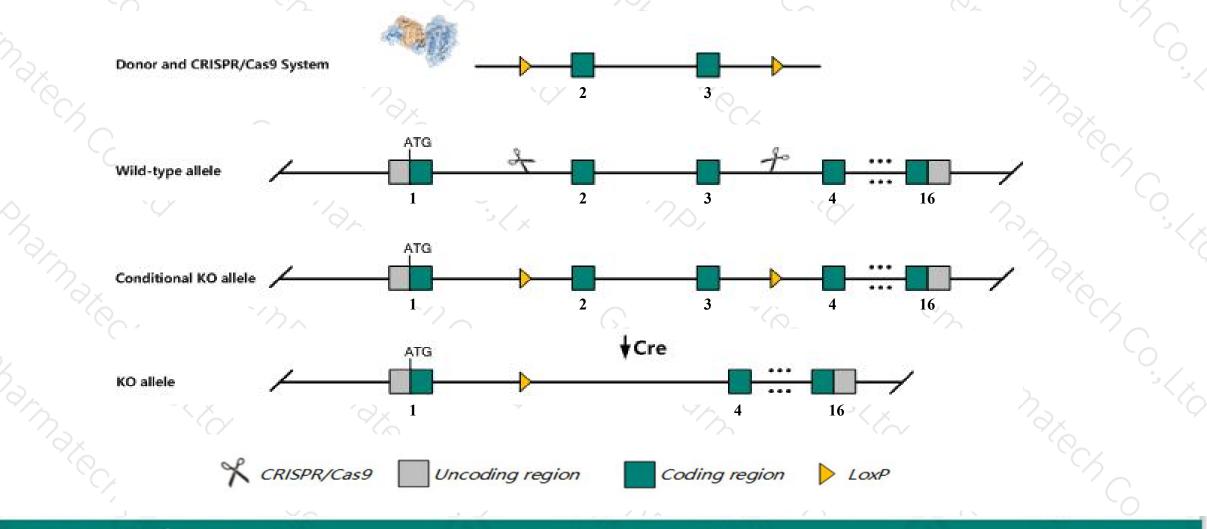


Conditional Knockout strategy



400-9660890

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



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The Scin gene has 2 transcripts. According to the structure of Scin gene, exon2-exon3 of Scin-201 (ENSMUST0000002640.5) transcript is recommended as the knockout region. The region contains 317bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Scin* 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.



- According to the existing MGI data, Mice homozygous for a conditional allele knocked-out in osteoclasts exhibit impaired osteoclast differentiation and reduced peridontal disease-mediated bone loss.
- The Scin gene is located on the Chr12. 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)



☆ ?

Scin scinderin [Mus musculus (house mouse)]

Gene ID: 20259, updated on 31-Jan-2019

Summary

Official SymbolScin provided by MGIOfficial Full Namescinderin provided by MGIPrimary sourceMGI:MGI:1306794See relatedEnsembl:ENSMUSG0000002565Gene typeprotein codingprotein codingVALIDATEDOrganismMus musculusLineageEukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Golires; Rodentia; Myomorpha;
Muroidea; Murinae; Mus; MusAlso knownasAW545522, adseverinExpressionBiased expression in colon adult (RPKM 56.3), large intestine adult (RPKM 22.6) and 4 other tissues
See more

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Transcript information (Ensembl)



The gene has 2 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Scin-201	ENSMUST0000002640.5	2995	<u>715aa</u>	Protein coding	CCDS49055	<u>Q60604</u>	TSL:1 GENCODE basic APPRIS P1
Scin-202	ENSMUST0000078481.13	2654	<u>615aa</u>	Protein coding	CCDS25891	<u>Q60604</u>	TSL:1 GENCODE basic

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

< Scin-201 protein coding

Reverse strand

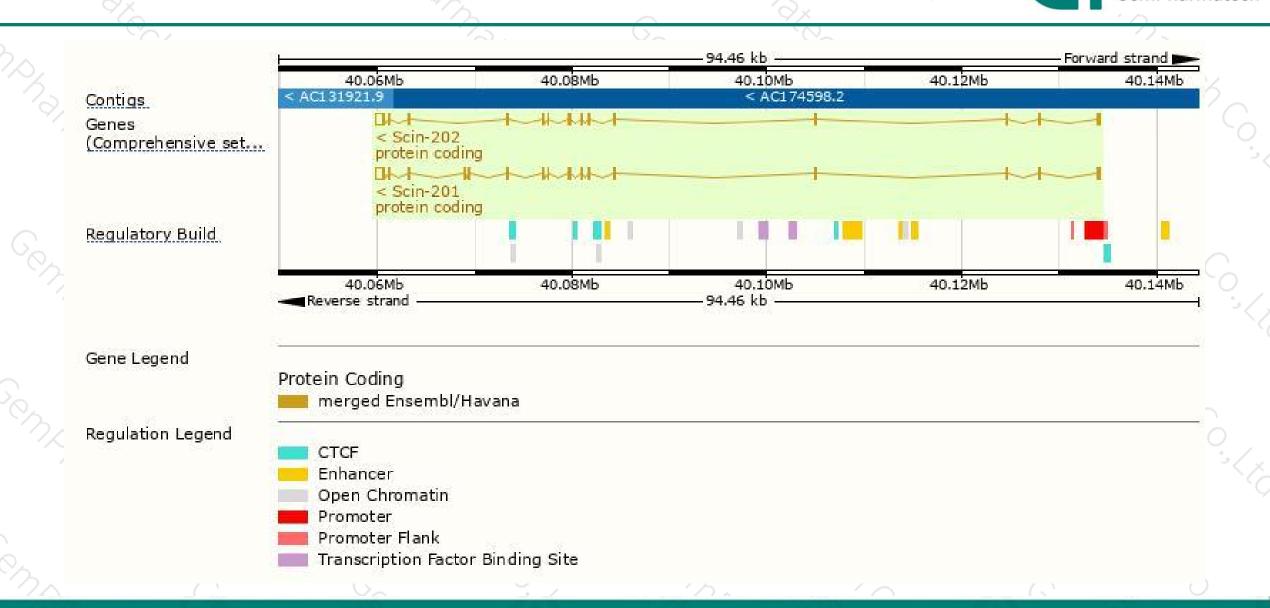
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Genomic location distribution



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Protein domain



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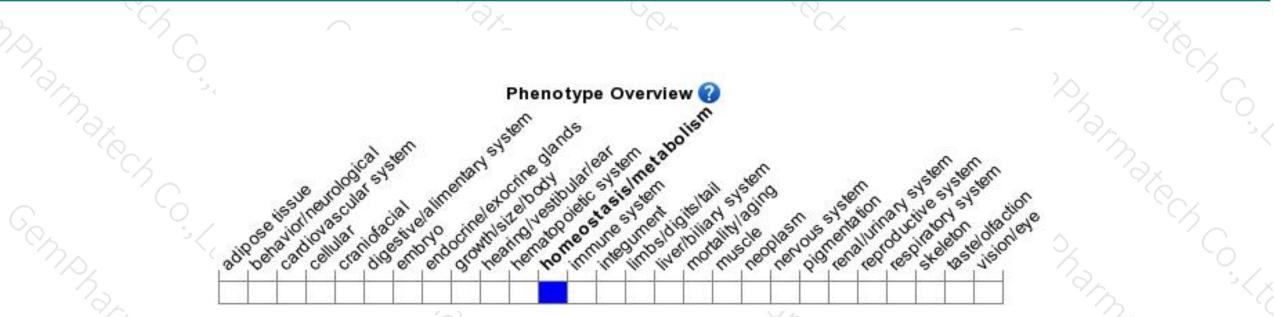
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Mouse phenotype description(MGI)





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, Mice homozygous for a conditional allele knocked-out in osteoclasts exhibit impaired osteoclast differentiation and reduced peridontal disease-mediated bone loss.



If you have any questions, you are welcome to inquire. Tel: 400-9660890



