

Scarf1 Cas9-KO Strategy

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Date:2020-02-25

Project Overview



Project Name

Scarf1

Project type

Cas9-KO

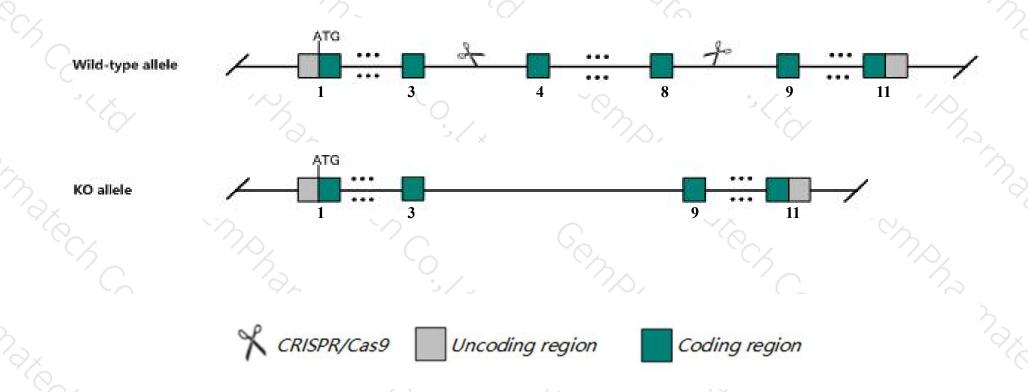
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Scarf1 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Scarf1* gene has 3 transcripts. According to the structure of *Scarf1* gene, exon4-exon8 of *Scarf1-201*(ENSMUST00000042808.12) transcript is recommended as the knockout region. The region contains 1093bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Scarf1* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Homozygous null mice are viable and fertile with no abnormalities detected in plasma glucose, cholesterol, or triglycerides, or in the brain, lung, heart, kidney, liver, or testes. Mice homozygous for a targeted allele exhibit impaired clearance of apoptotic cells and autoimmune disease.
- > The N-terminal of Scarf1 gene will remain several amino acids, it may remain the partial function of Scarf1 gene.
- The knockout region is near to the C-terminal of *Rilp* gene, this strategy may influence the regulatory function of the C-terminal of *Rilp* gene.
- The *Scarf1* gene is located on the Chr11. 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)



Scarf1 scavenger receptor class F, member 1 [Mus musculus (house mouse)]

Gene ID: 380713, updated on 12-Aug-2019

▲ Summary

☆ ?

Official Symbol Scarf1 provided by MGI

Official Full Name scavenger receptor class F, member 1 provided by MGI

Primary source MGI:MGI:2449455

See related Ensembl: ENSMUSG00000038188

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as SREC; SREC-I; AA986099; mKIAA0149

Expression Broad expression in lung adult (RPKM 24.0), adrenal adult (RPKM 13.3) and 24 other tissues See more

Orthologs human all

Genomic context



Location: 11; 11 B5

See Scarf1 in Genome Data Viewer

Exon count: 10

Annotation release	Status	Assembly		Location NC_000077.6 (7551348975526582) NC_000077.5 (7532704375340082)	
108currentBuild 37.2previous assembly		GRCm38.p6 (GCF_000001635.26)	11		
		MGSCv37 (GCF_000001635.18)	11		

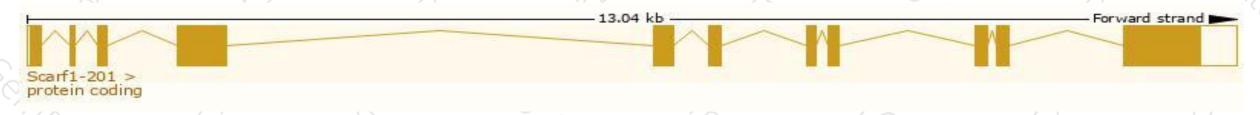
Transcript information (Ensembl)



The gene has 3 transcripts, all transcripts are shown below:

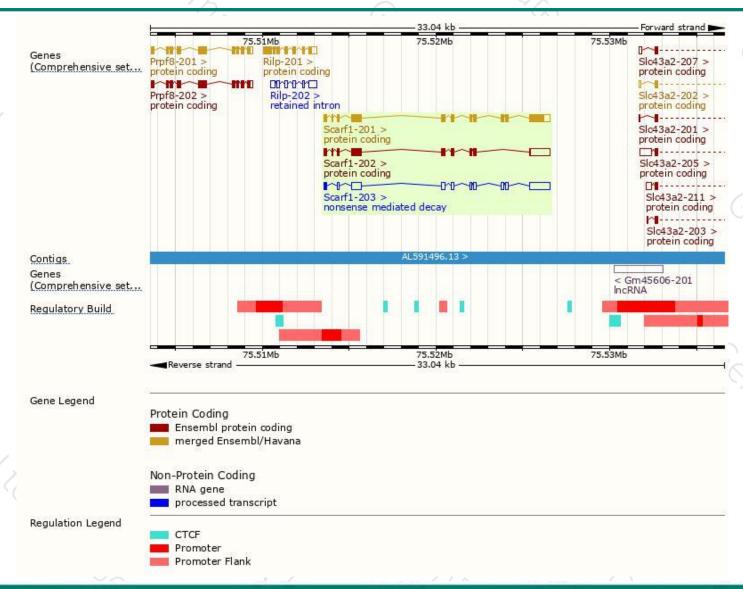
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Scarf1-201	ENSMUST00000042808.12	2887	820aa	Protein coding	CCDS25050	Q5ND28	TSL:1 GENCODE basic APPRIS P2
Scarf1-202	ENSMUST00000118243.1	2617	<u>478aa</u>	Protein coding	9 4 3	B7ZC28	TSL:1 GENCODE basic APPRIS ALT2
Scarf1-203	ENSMUST00000123819.1	2789	<u>40aa</u>	Nonsense mediated decay	020	D6RJK6	TSL:1

The strategy is based on the design of Scarf1-201 transcript, The transcription is shown below



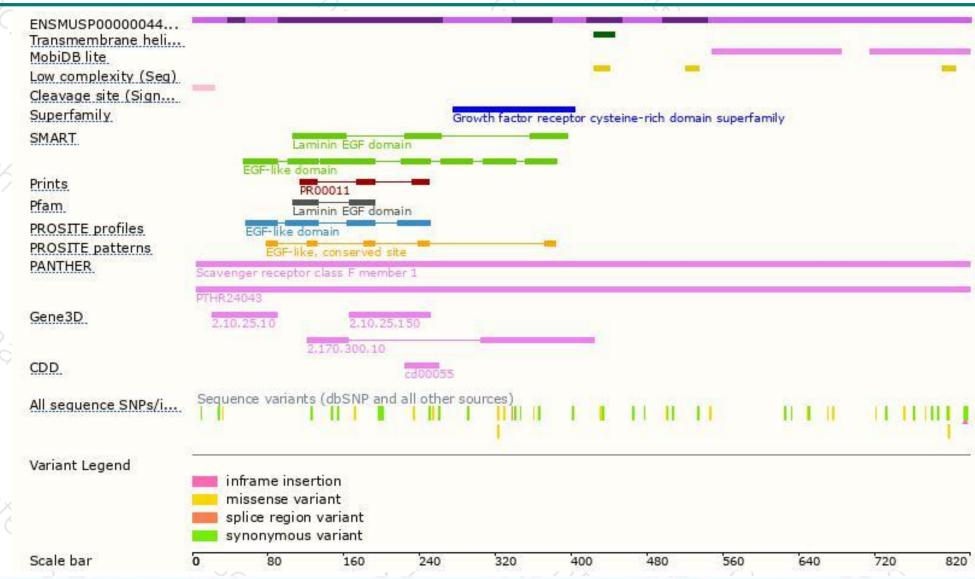
Genomic location distribution





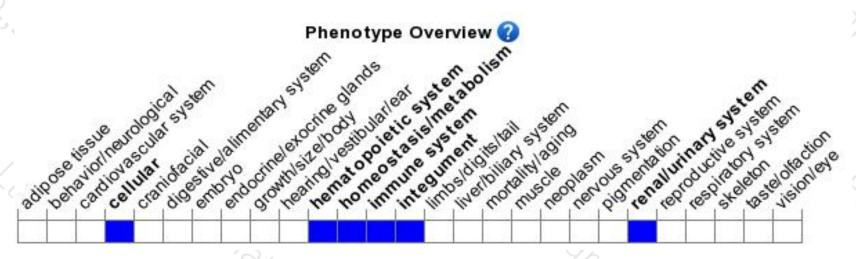
Protein domain





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, Homozygous null mice are viable and fertile with no abnormalities detected in plasma glucose, cholesterol, or triglycerides, or in the brain, lung, heart, kidney, liver, or testes. Mice homozygous for a targeted allele exhibit impaired clearance of apoptotic cells and autoimmune disease.



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





