

Stard7 Cas9-CKO Strategy

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Project Overview



Project Name

Stard7

Project type

Cas9-CKO

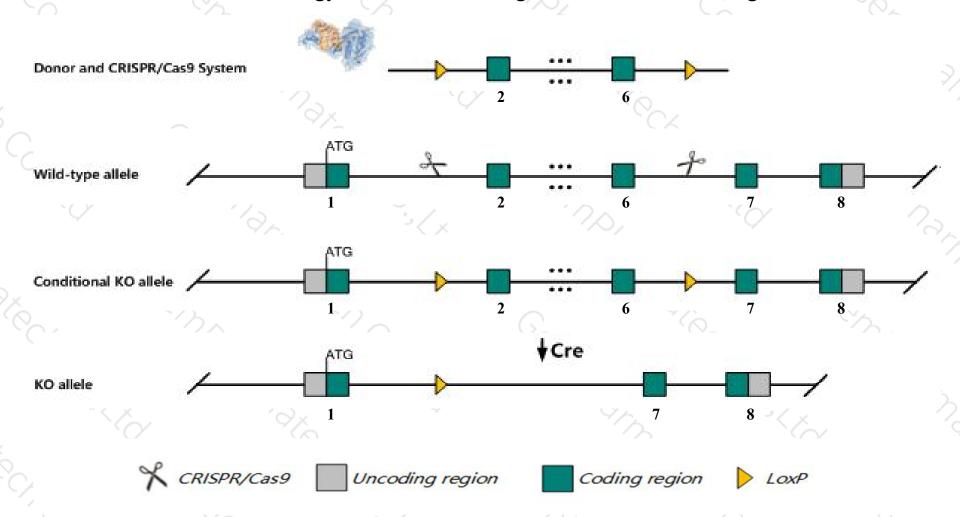
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Stard7 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Stard7* gene has 8 transcripts. According to the structure of *Stard7* gene, exon2-exon6 of *Stard7-202*(ENSMUST00000110375.8) transcript is recommended as the knockout region. The region contains 553bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Stard7* 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.

Notice



- ➤ According to the existing MGI data, Heterozygous KO results in exaggerated allergic response (lung inflammation, increased epithelial barrier permeability and airway responsiveness) and atopic dermatitis.
- The *Stard7* gene is located on the Chr2. 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)



Stard7 START domain containing 7 [Mus musculus (house mouse)]

Gene ID: 99138, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Stard7 provided by MGI

Official Full Name START domain containing 7 provided by MGI

Primary source MGI:MGI:2139090

See related Ensembl: ENSMUSG00000027367

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 Al852671, AL022671, AW544915

Expression Ubiquitous expression in CNS E18 (RPKM 28.6), heart adult (RPKM 28.6) and 28 other tissuesSee more

Orthologs <u>human</u> all

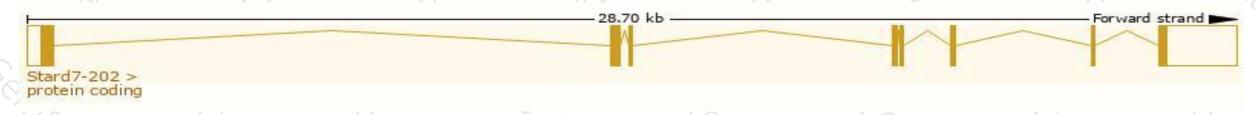
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

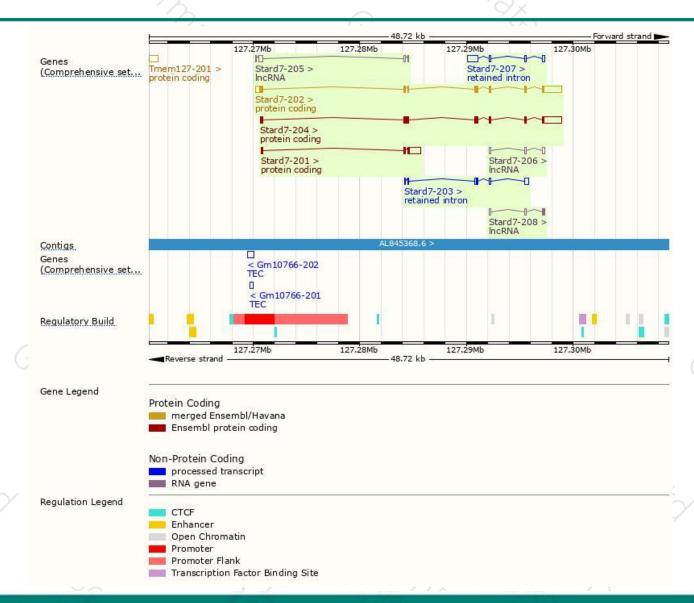
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Stard7-202	ENSMUST00000110375.8	3116	<u>373aa</u>	Protein coding	CCDS16698	Q8R1R3	TSL:1 GENCODE basic APPRIS P1
Stard7-204	ENSMUST00000125049.1	2869	<u>409aa</u>	Protein coding	89	F7BBA2	CDS 5' incomplete TSL:2
Stard7-201	ENSMUST00000110374.1	1530	<u>122aa</u>	Protein coding	91	Q8BXP7	TSL:1 GENCODE basic
Stard7-207	ENSMUST00000144687.7	1358	No protein	Retained intron	e e	-	TSL:2
Stard7-203	ENSMUST00000123152.7	773	No protein	Retained intron	85		TSL:2
Stard7-205	ENSMUST00000131196.1	679	No protein	IncRNA	89	-	TSL:5
Stard7-206	ENSMUST00000135616.7	459	No protein	IncRNA	91	-	TSL:1
Stard7-208	ENSMUST00000154549.1	400	No protein	IncRNA	62	2	TSL:5

The strategy is based on the design of Stard7-202 transcript, The transcription is shown below



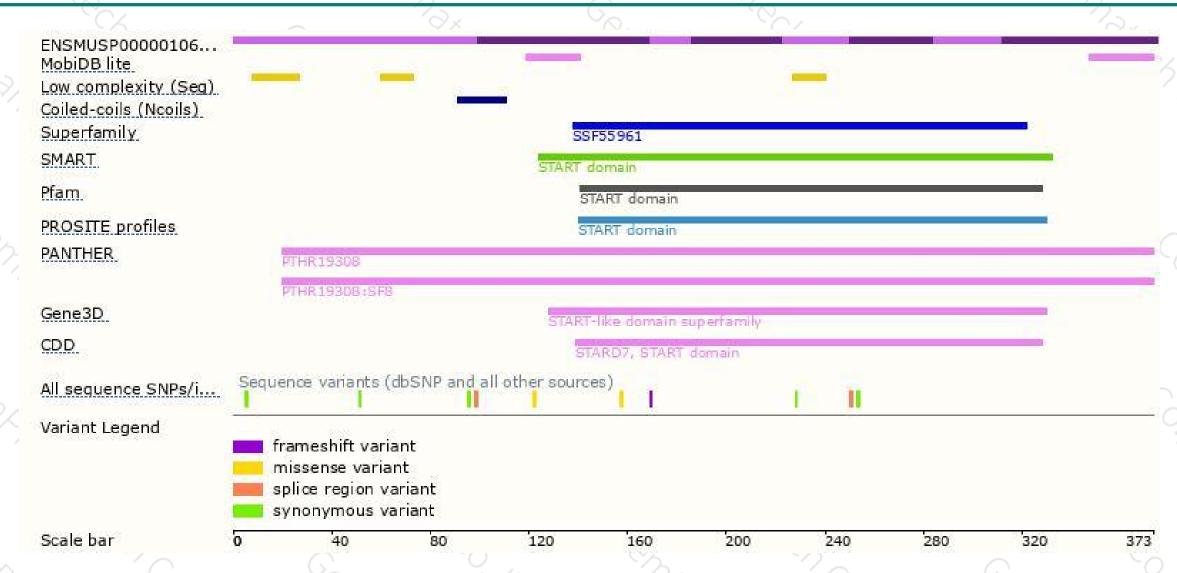
Genomic location distribution





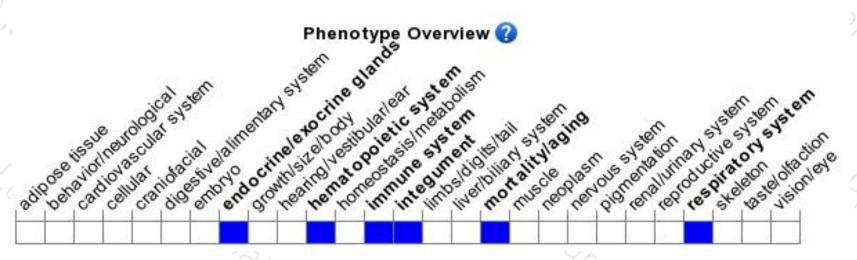
Protein domain





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, Heterozygous KO results in exaggerated allergic response (lung inflammation, increased epithelial barrier permeability and airway responsiveness) and atopic dermatitis.



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





