

Socs3 Cas9-CKO Strategy

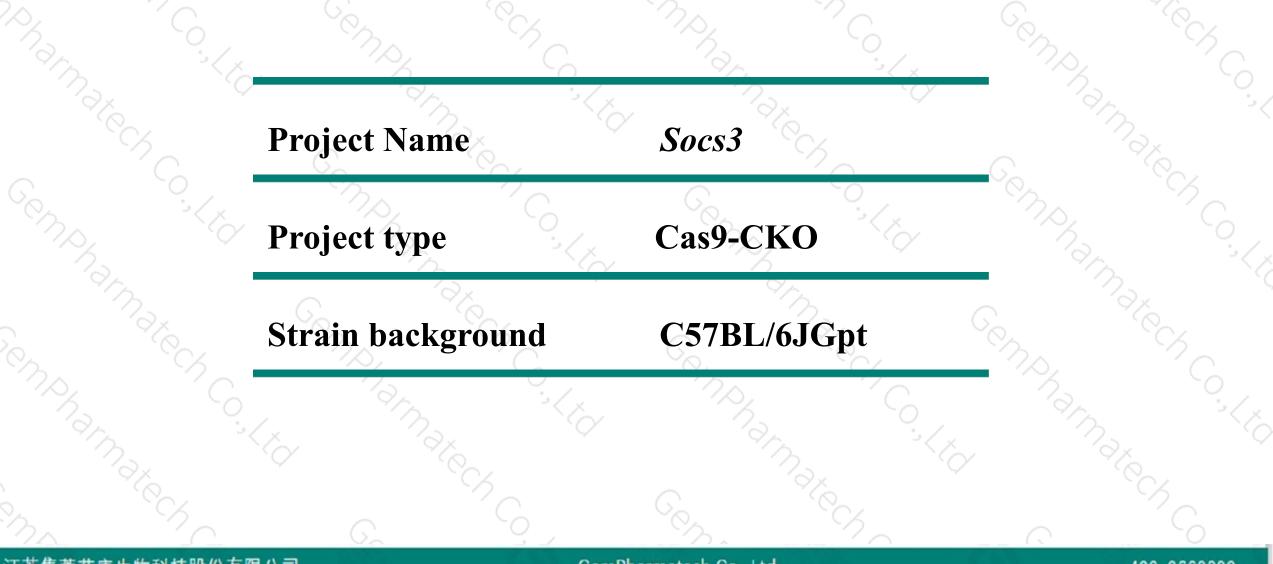
Designer: Reviewer:

Design Date:

Daohua Xu Huimin Su 2019-9-28

Project Overview





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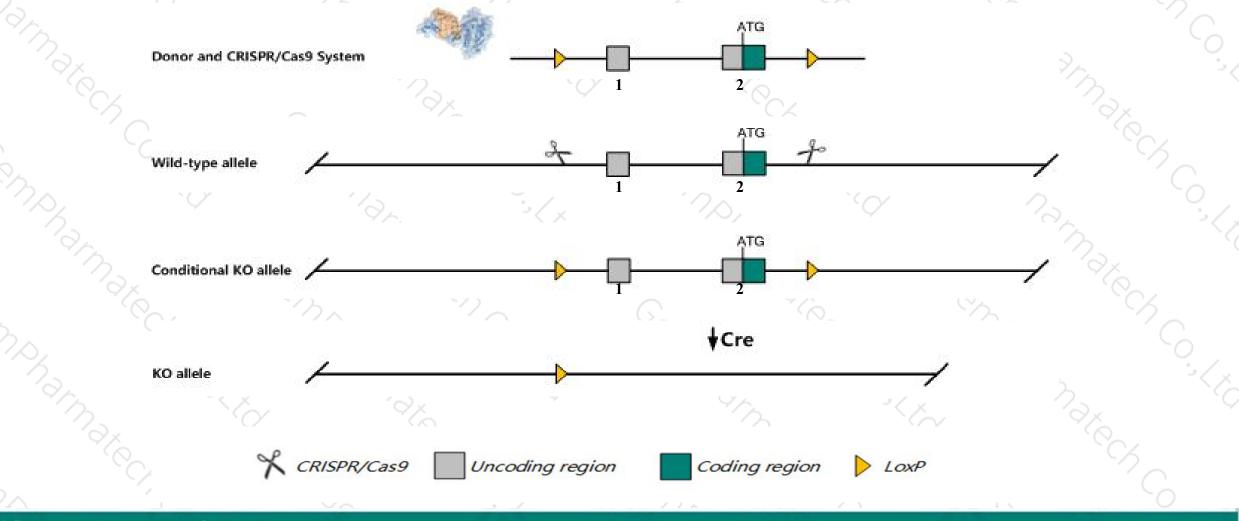
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Conditional Knockout strategy



400-9660890

This model will use CRISPR/Cas9 technology to edit the Socs3 gene. The schematic diagram is as follows:



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The Socs3 gene has 3 transcripts. According to the structure of Socs3 gene, exon1-exon2 of Socs3-201 (ENSMUST00000054002.3) 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 *Socs3* 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 mutations cause impairment if gene expression show immunological and hematopoetic abnormalities. Complete gene disruption causes lethality.
- The Socs3 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)



< ?

Socs3 suppressor of cytokine signaling 3 [Mus musculus (house mouse)]

Gene ID: 12702, updated on 2-Apr-2019

Summary

Official SymbolSocs3 provided by MGIOfficial Full Namesuppressor of cytokine signaling 3 provided by MGIPrimary sourceMGI:MGI:1201791See relatedEnsembl:ENSMUSG0000053113Gene typeprotein codingRefSeq statusVALIDATEDOrganismMus musculusLineageEukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;
Muroidea; Murinae; Mus; MusAlso knownaGis3, Cish3, EF-10, Ef10, SSI-3, Ssi3ExpressionBroad expression in large intestine adult (RPKM 19.8), duodenum adult (RPKM 16.3) and 25 other tissues
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The gene has 3 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags		
Socs3-201	ENSMUST0000054002.3	2552	<u>225aa</u>	Protein coding	CCDS25697	<u>035718</u>	TSL:1 GENCODE basic APPRIS P1		
Socs3-202	ENSMUST00000123255.1	490	No protein	IncRNA	-	1.0	TSL:3		
Socs3-203	ENSMUST00000132295.1	246	No protein	IncRNA	-	(22)	TSL:5		

The strategy is based on the design of Socs3-201 transcript, The transcription is shown below

< Socs3-201 protein coding

Reverse strand

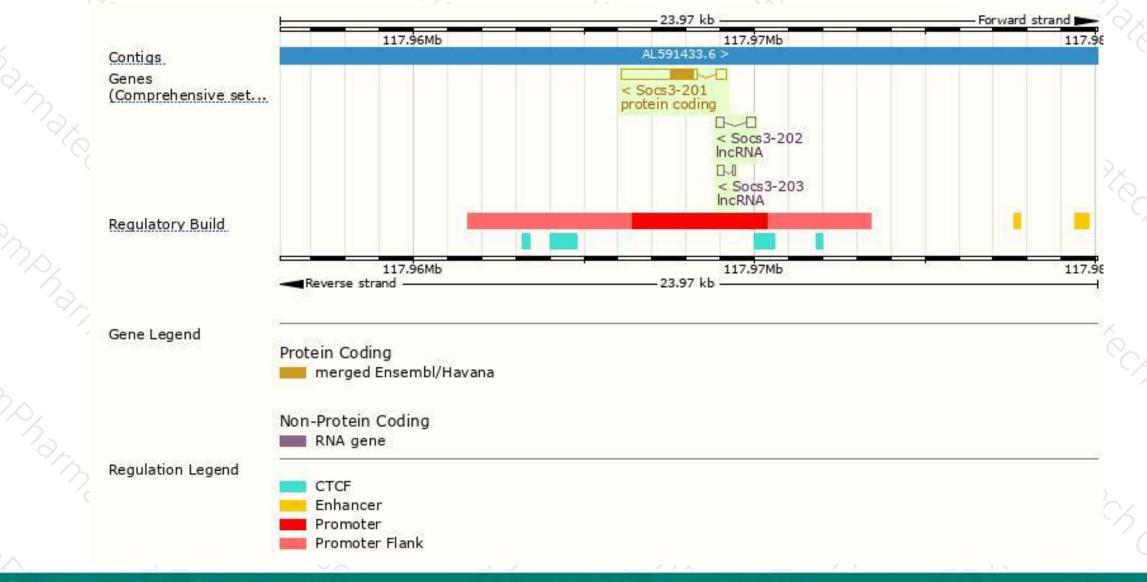
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Genomic location distribution





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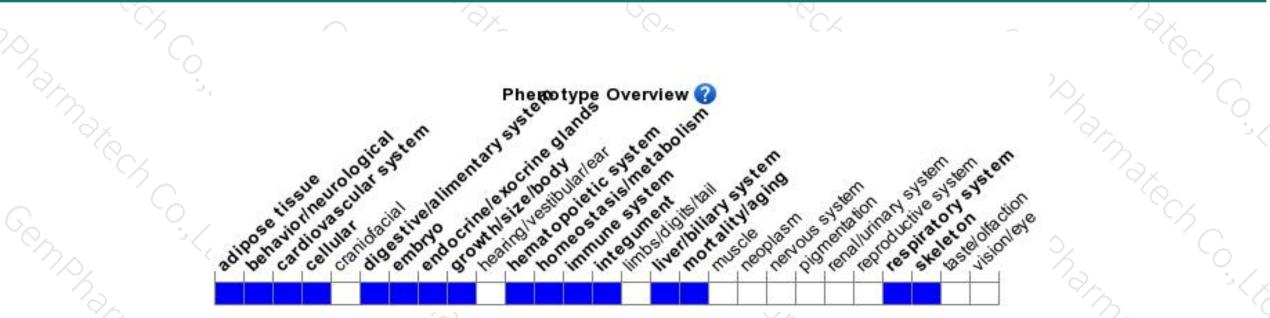
Protein domain



Scale bar	0 20	40	60	80	100	120	140	160	180	200 2	25
Variant Legend	synonymo	ous variant									5
All sequence SNPs/i	Sequence varia	ants (dbSNP	and all othe	er sources)	1						
<u>CDD</u>		SOCS3, S	H2 domain						cd03737		
Gene3D	saled 2 of an and a find the co	omain superfa	100.00						-		
PANTHER	PTHR10155 Suppressor of cy	rtokine signali	ing 3								
PROSITE profiles		SH	2 domain						SOCS box do	omain	-
Pfam		SH:	2 domain							oox domain	
SMART		SH2	domain			_			SOCS BOX	oox domain	
MobiDB lite Low complexity (Seg) Superfamily		SH2 domair	n superfamily			_			SOCS box-lik	e domain sup	erfa

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 mutations cause impairment if gene expression show immunologic and hematopoetic abnormalities. Complete gene disruption causes lethality.



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



