

Rab38 Cas9-CKO Strategy

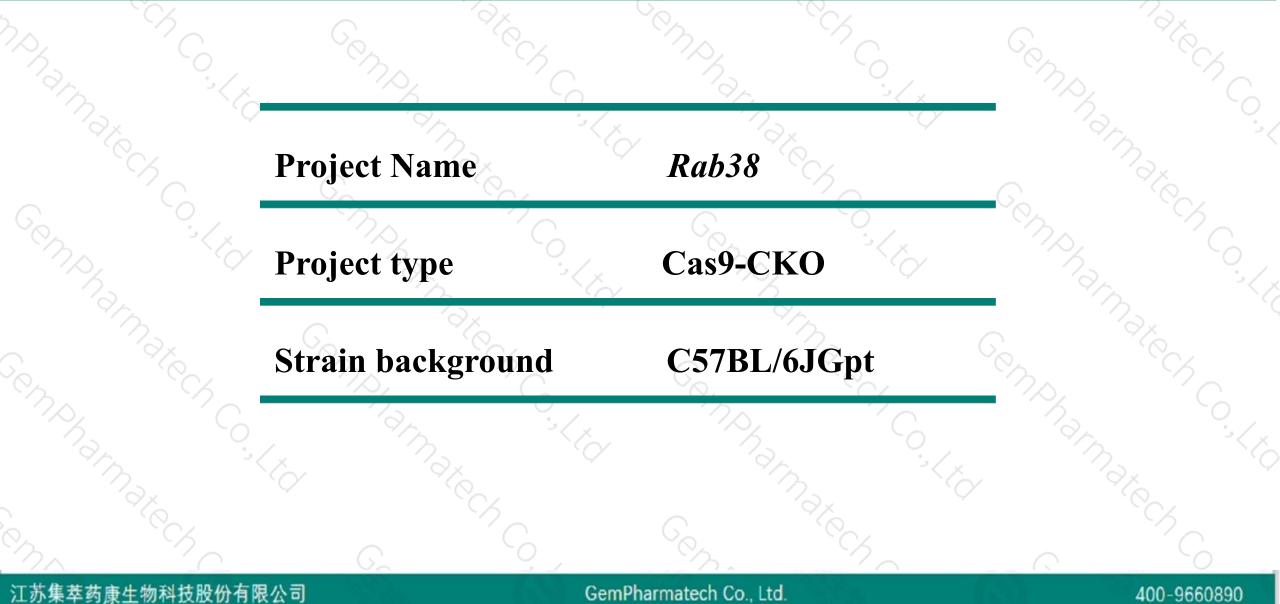
Designer: Reviewer:

Design Date:

Daohua Xu Huimin Su 2020-2-14

Project Overview

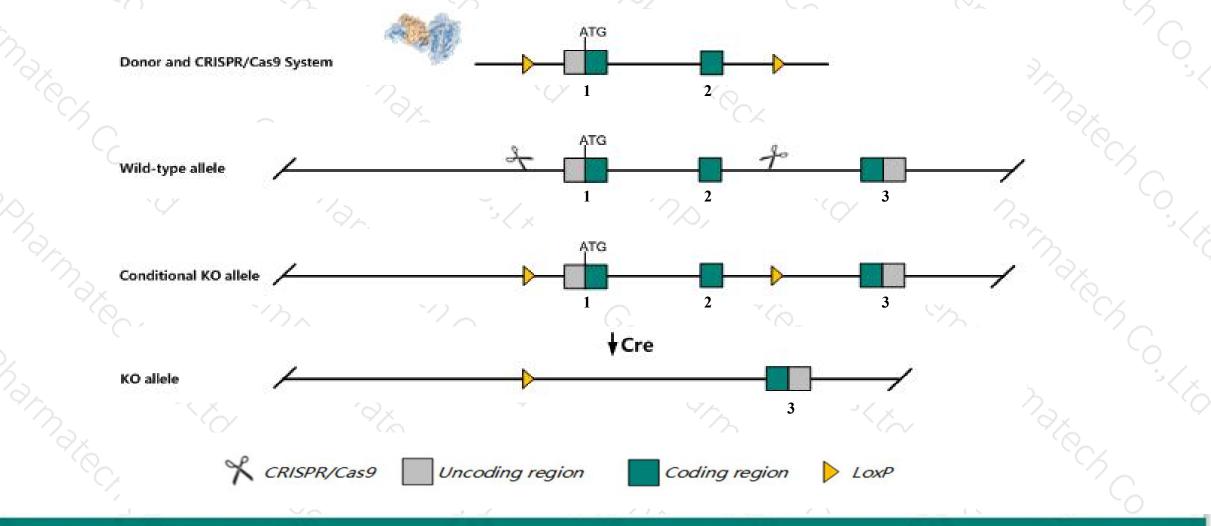




Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Rab38* gene. The schematic diagram is as follows:



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The Rab38 gene has 2 transcripts. According to the structure of Rab38 gene, exon1-exon2 of Rab38-201 (ENSMUST00000107256.3) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Rab38* 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 spontaneous mutation display oculocutaneous albinism, abnormal lung alveolar structure and aberrant pulmonary surfactant homeostasis.
 The KO region contains functional region of the *Gm15661* gene.Knockout the region may affect the function of *Gm15661* gene.
- The Rab38 gene is located on the Chr7. 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)



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Rab38 RAB38, member RAS oncogene family [Mus musculus (house mouse)]

Gene ID: 72433, updated on 31-Jan-2019

Summary

Official Symbol	Rab38 provided by MGI
Official Full Name	RAB38, member RAS oncogene family provided by MGI
Primary source	MGI:MGI:1919683
See related	Ensembl:ENSMUSG0000030559
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	2310011F14Rik, AU043391, cht
Expression	Broad expression in limb E14.5 (RPKM 5.1), kidney adult (RPKM 3.0) and 18 other tissuesSee more
Orthologs	human all

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Transcript information (Ensembl)



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

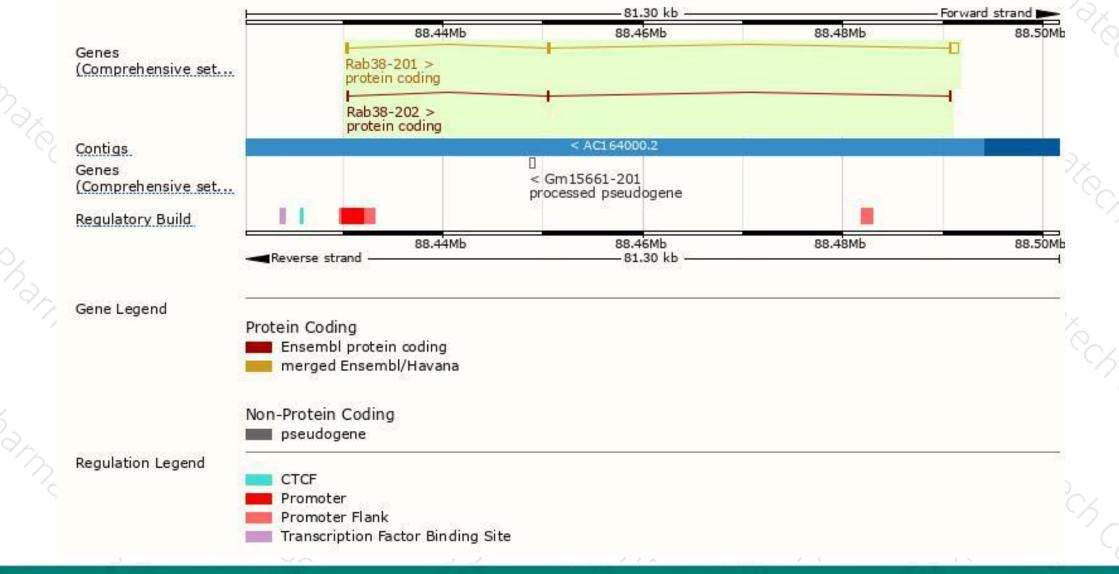
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Rab38-201	ENSMUST00000107256.3	1577	<u>211aa</u>	Protein coding	CCDS21439	Q5FW76 Q8QZZ8	TSL:1 GENCODE basic APPRIS P1
Rab38-202	ENSMUST00000208478.1	579	<u>121aa</u>	Protein coding	8	A0A140LHK2	TSL:3 GENCODE basic

The strategy is based on the design of Rab38-201 transcript, The transcription is shown below



Genomic location distribution





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Protein domain



ENSMUSP00000102... TIGRFAM

Superfamily SMART

Prints Pfam

PROSITE profiles PANTHER

Gene3D CDD

Scale bar

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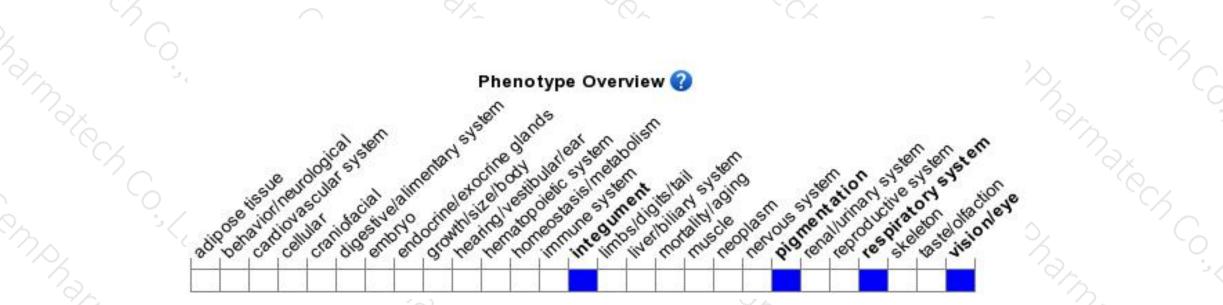
All sequence SNPs/i...

Variant Legend

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	las-related prot ence variants (0
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-	HR24073										
- 112.0	HR24073:SF839	9									
PS 51	1419										
	Small GTPase										
P	SM00176			<u>.</u>			-	<u></u>			
	SM00174									-	
S	M00175										
SM	00173										
P-lo	op containing r	nucleoside t	riphosphate	hydrolase							
S	mall GTP-bindi	ng protein o	omain								
						100					

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 spontaneous mutation display oculocutaneous albinism, abnorma lung alveolar structure and aberrant pulmonary surfactant homeostasis.



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



