

Cep290 Cas9-KO Strategy

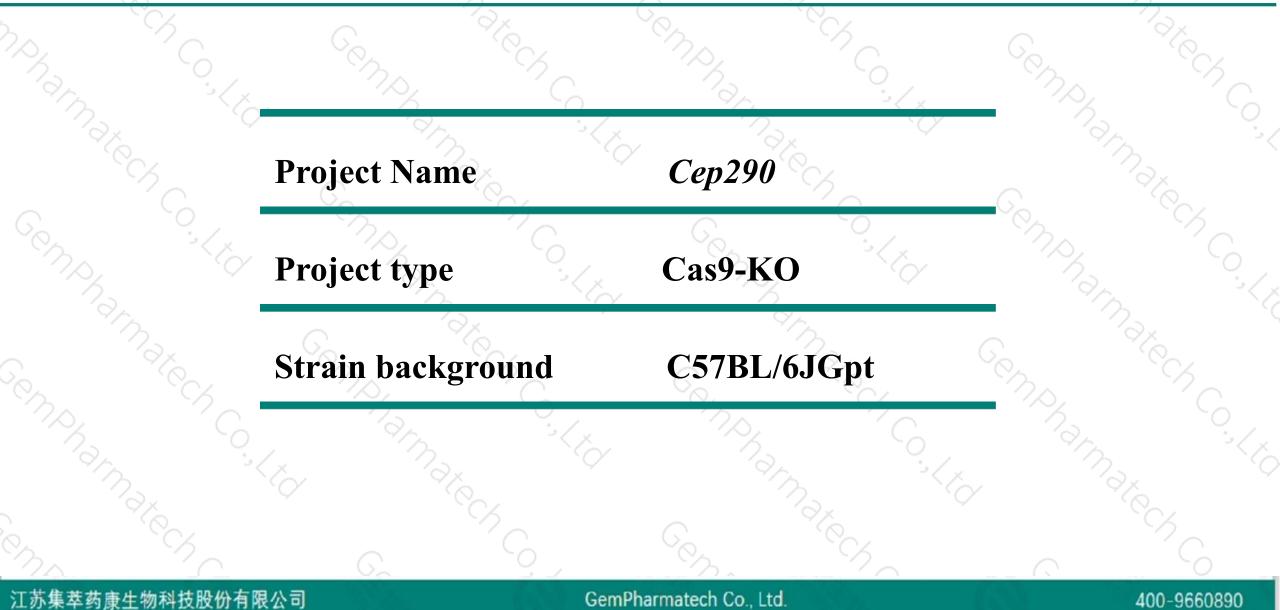
Designer: Reviewer:

Design Date:

Daohua Xu Huimin Su 2019-10-28

Project Overview

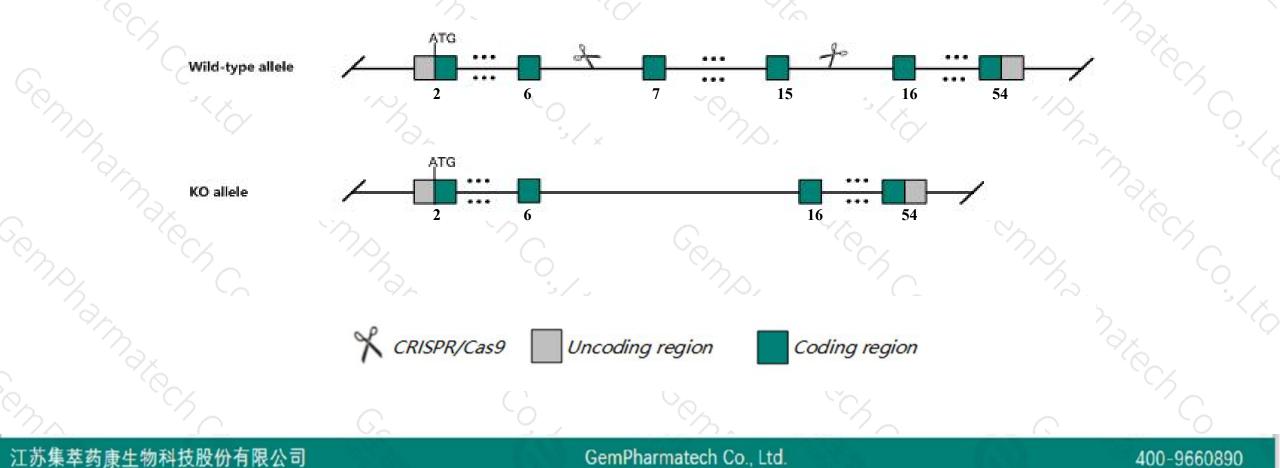




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Cep290 gene. The schematic diagram is as follows:





- The Cep290 gene has 10 transcripts. According to the structure of Cep290 gene, exon7-exon15 of Cep290-210 (ENSMUST00000220346.1) transcript is recommended as the knockout region. The region contains 1084bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Cep290 gene. The brief process is as follows: CRISPR/Cas9 system



- According to the existing MGI data, Mutant mice display mislocalization of ciliary and phototransduction proteins resulting in early-onset retinal degeneration. Heterotaxy with transposition of the great arteries (TGA), atrioventricular septal defect (AVSD), left bronchial isomerism, and hypoplastic spleen is also seen.
- The Cep290 gene is located on the Chr10. 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)



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Cep290 centrosomal protein 290 [Mus musculus (house mouse)]

Gene ID: 216274, updated on 5-Mar-2019

Summary

Official Symbol	Cep290 provided by MGI
Official Full Name	centrosomal protein 290 provided byMGI
Primary source	MGI:MGI:2384917
See related	Ensembl:ENSMUSG00000019971
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	BC004690, Nphp6, b2b1454Clo, b2b1752Clo
Expression	Biased expression in CNS E11.5 (RPKM 1.5), testis adult (RPKM 1.4) and 14 other tissues See more
Orthologs	human all

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Transcript information (Ensembl)



The gene has 10 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Cep290-210	ENSMUST00000220346.1	8211	<u>2479aa</u>	Protein coding	CCDS48685	<u>E9Q9M0</u>	TSL:5 GENCODE basic APPRIS P2	
Cep290-201	ENSMUST00000164751.1	8006	<u>2479aa</u>	Protein coding	CCDS48685	<u>E9Q9M0</u>	TSL:5 GENCODE basic APPRIS P2	
Cep290-206	ENSMUST00000219765.1	7985	<u>2472aa</u>	Protein coding	(4)	Q6A078	TSL:5 GENCODE basic APPRIS ALT2	
Cep290-204	ENSMUST00000219408.1	1177	<u>237aa</u>	Protein coding	1.27	A0A1W2P6U7	CDS 5' incomplete TSL:2	
Cep290-207	ENSMUST00000219889.1	2170	No protein	Retained intron	-	₹.6	TSL:1	
Cep290-202	ENSMUST00000218000.1	1567	No protein	Retained intron	6.84	, 4	TSL:5	
Cep290-203	ENSMUST00000218703.1	1406	No protein	Retained intron	(4)	49	TSL:5	
Cep290-208	ENSMUST00000220231.1	903	No protein	Retained intron	1 <u>17</u> 1	<u>1</u> 0	TSL:5	
Cep290-209	ENSMUST00000220331.1	423	No protein	Retained intron	173	₹.6	TSL:3	
Cep290-205	ENSMUST00000219643.1	2738	No protein	IncRNA	(.	- 3	TSL:1	

The strategy is based on the design of Cep290-210 transcript, The transcription is shown below

Cep290-210 > protein coding

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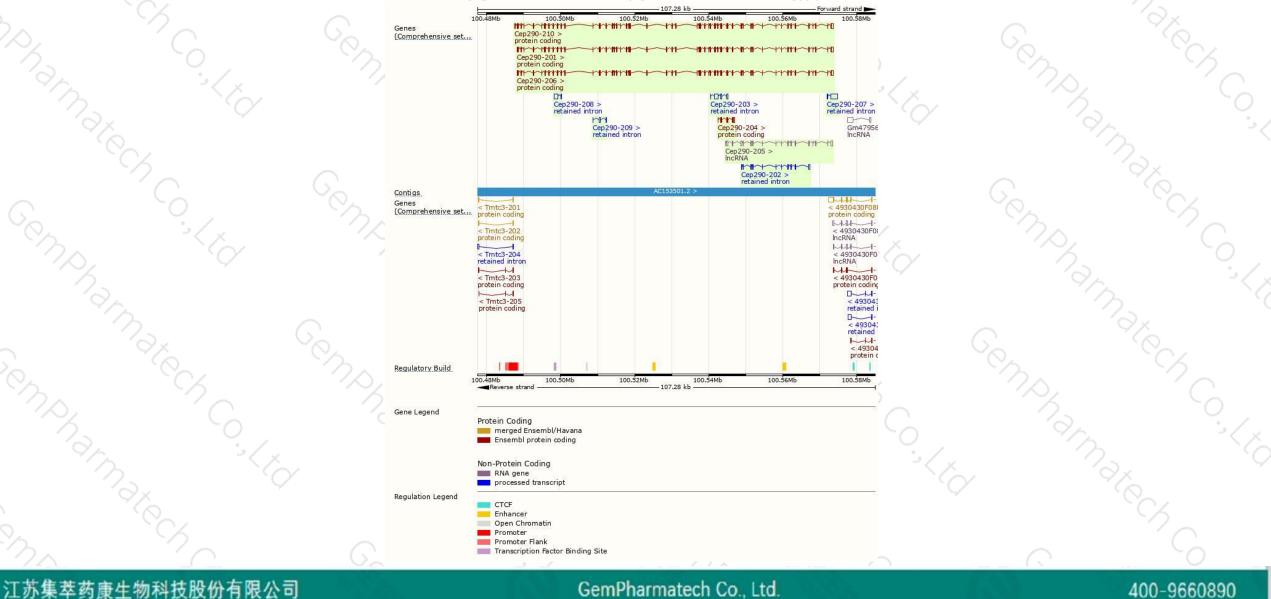
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Forward strand

Genomic location distribution





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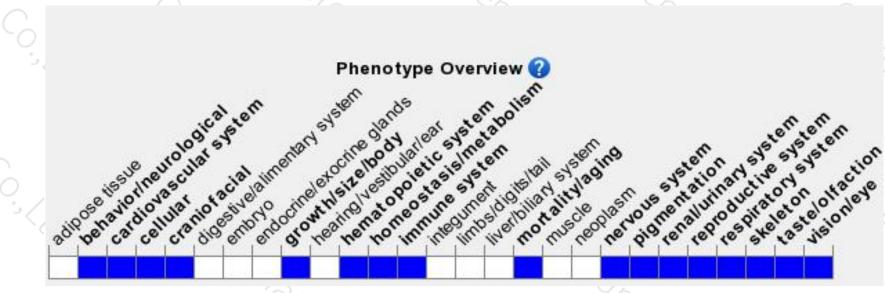
Protein domain



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47	ENSMUSP00000151 MobiDB lite Low complexity (Seg) Coiled-coils (Ncoils) Pfam				osomal protein of 290k	Da. coiled-coil region	
Con .	PANTHER	PTHR18879:SF20 Centrosomal protein of	290kDa	- Juenti	eleventi en eleventi en eleventi	en, carea con region	
19	All sequence SNPs/i	Sequence variants (db)		urces)	0.0100		
Sent	Variant Legend	stop gained frameshift variant missense variant splice region varia	ant				
	Scale bar	<b>o</b> 400	800	1200	1600	2000	2479
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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, Mutant mice display mislocalization of ciliary and phototransduction proteins resulting in early-onset retinal degeneration. Heterotaxy with transposition of the great arteries (TGA), atrioventricular septal defect (AVSD), left bronchial isomerism, and hypoplastic spleen is also seen.

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



