

Ap3s2 Cas9-CKO Strategy

Designer: Reviewer:

Design Date:

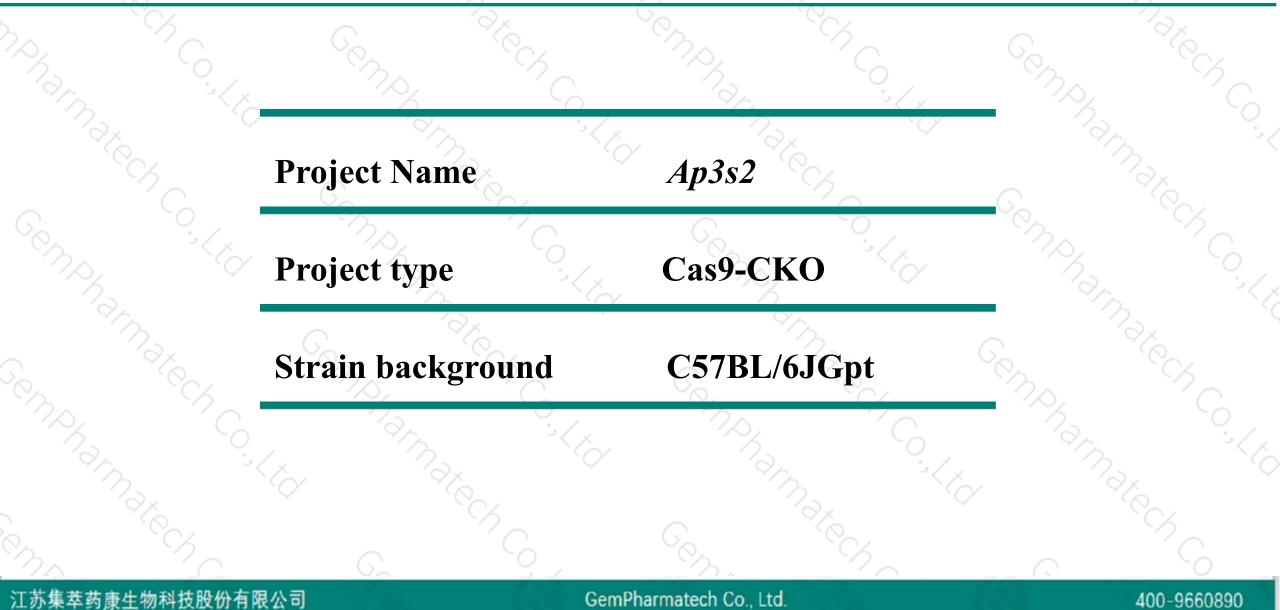
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2020-4-9

Project Overview

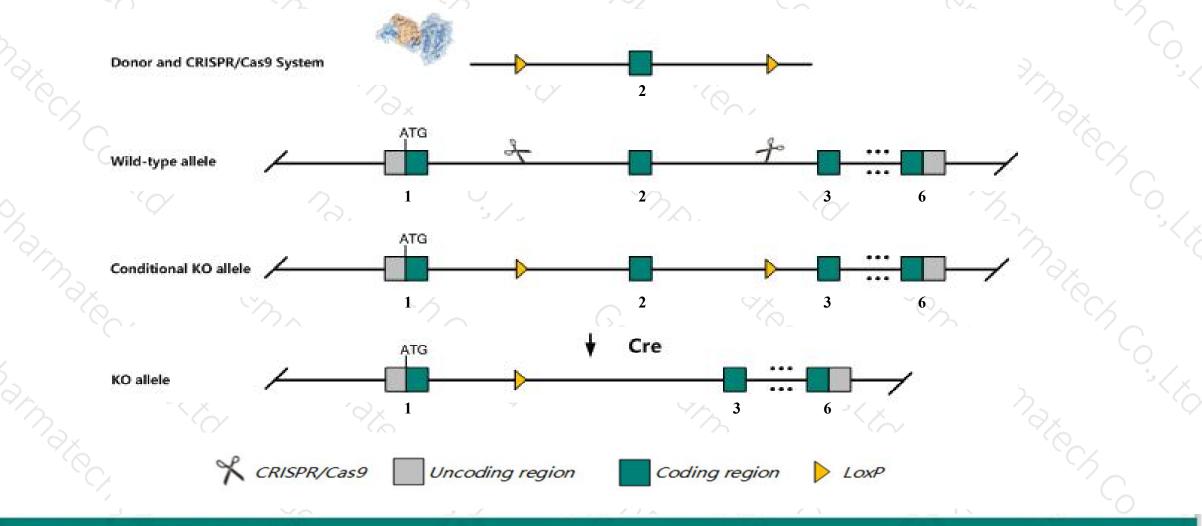




Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ap3s2* gene. The schematic diagram is as follows:



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400-9660890



The Ap3s2 gene has 2 transcripts. According to the structure of Ap3s2 gene, exon2 of Ap3s2-201 (ENSMUST00000075657.7) transcript is recommended as the knockout region. The region contains 92bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Ap3s2* 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



- ➤ The 5-terminal regulatin of 5430400D12Rik and Gm26646 may be affect.
- The Ap3s2 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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See Ap3s2 in Genome Data Viewer

Ap3s2 adaptor-related protein complex 3, sigma 2 subunit [Mus musculus (house mouse)]

Gene ID: 11778, updated on 13-Mar-2020

- Summary

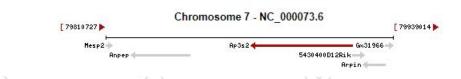
Official Symbol	Ap3s2 provided by MGI
Official Full Name	adaptor-related protein complex 3, sigma 2 subunit provided by MGI
Primary source	MGI:MGI:1337060
See related	Ensembl:ENSMUSG0000063801
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae;
	Murinae; Mus; Mus
Also known as	[s]3B
Expression	Ubiquitous expression in whole brain E14.5 (RPKM 17.2), CNS E14 (RPKM 15.6) and 28 other tissues See more
Orthologs	human all

Genomic context

Location: 7; 7 D2

Exon count: 6

Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	7	NC_000073.6 (7987532579920640, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	7	NC_000073.5 (8702021187065526, complement)



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Transcript information (Ensembl)



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

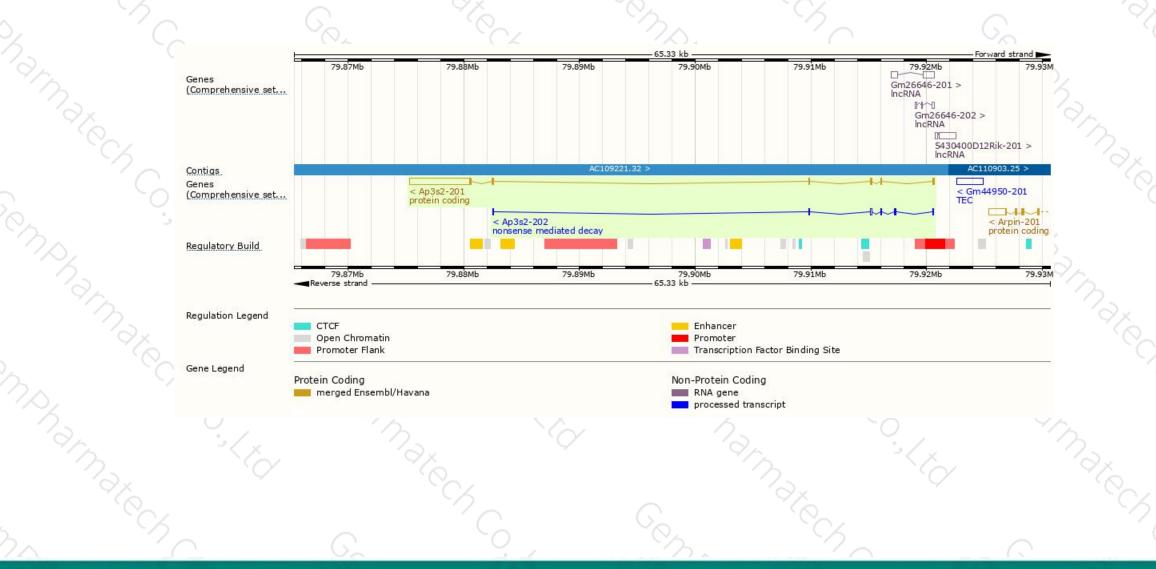
Name 🖕	Transcript ID	bp 🖕	Protein 🖕	Protein 🛊 Biotype 🔶		UniProt 💧	Flags 🍦		
Ap3s2-201	ENSMUST0000075657.7	5813	<u>193aa</u>	Protein coding	<u>CCDS21389</u> 교	Q8BSZ2 @	TSL:1	GENCODE basic	APPRIS P1
Ap3s2-202	ENSMUST00000206725.1	519	<u>42aa</u>	Nonsense mediated decay	-	<u>A0A0U1RQ64</u> ഗ്ര		TSL:3	

The strategy is based on the design of *Ap3s2-201* transcript, the transcription is shown below

< Ap3s2-201 protein coding

Genomic location distribution





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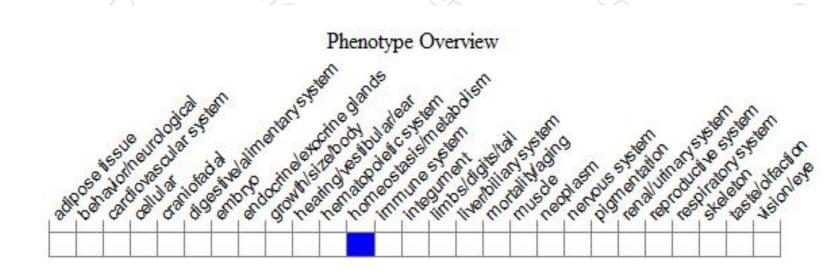
Protein domain



ENSMUSP00000075	Ç		24	X _n		'Ox			Γς.	
Superfamily Pfam	Longin-like domain superf									
PROSITE patterns	AP complex, mu/sigma subunit Clathrin adaptor complex, small chain									
PANTHER	PTHR11753:SF10									
Gene3D	Adaptor protein complex, 3.30.450.60	sigma subunit							-	
CDD	cd14834									
All sequence SNPs/i	Sequence variants (dbS	NP and all other sour	ces)						<u> </u>	
Variant Legend	splice region varia	nt			synonyn	nous variant				
Scale bar	0 20	40	60	80	100	120	140	160	193	
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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/).



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



