

Gabpb1 Cas9-KO Strategy

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Project Overview



Project Name

Gabpb1

Project type

Cas9-KO

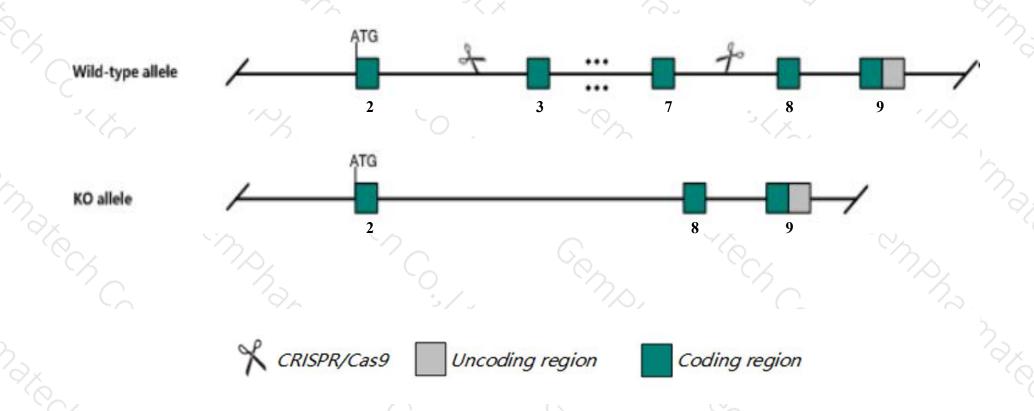
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Gabpb1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Gabpb1* gene has 9 transcripts. According to the structure of *Gabpb1* gene, exon3-exon7 of *Gabpb1*-206(ENSMUST00000110425.8) transcript is recommended as the knockout region. The region contains 775bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Gabpb1* gene. The brief process is as follows: CRISPR/Cas9 system 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.

Notice



- > According to the existing MGI data, mice homozygous for a null allele exhibit embryonic lethality by E12.5.
- > The *Gabpb1* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Gabpb1 GA repeat binding protein, beta 1 [Mus musculus (house mouse)]

Gene ID: 14391, updated on 13-Mar-2020





Official Symbol Gabpb1 provided by MGI

Official Full Name GA repeat binding protein, beta 1 provided by MGI

Primary source MGI:MGI:95611

See related Ensembl: ENSMUSG00000027361

Gene type protein coding

RefSeq status VALIDATED

Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as BABPB2, E4TF1, E4TF1-47, E4TF1-53, E4TF1B, GABPB, GABPB-1, GABPB-2, GABPB1-1, GABPB1-2, NRF2B1, NRF2B2

Expression Ubiquitous expression in CNS E11.5 (RPKM 6.0), limb E14.5 (RPKM 4.8) and 28 other tissuesSee more

Orthologs <u>human</u> <u>all</u>

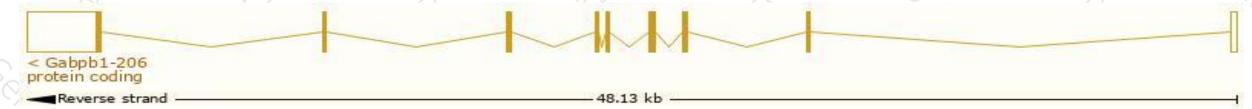
Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

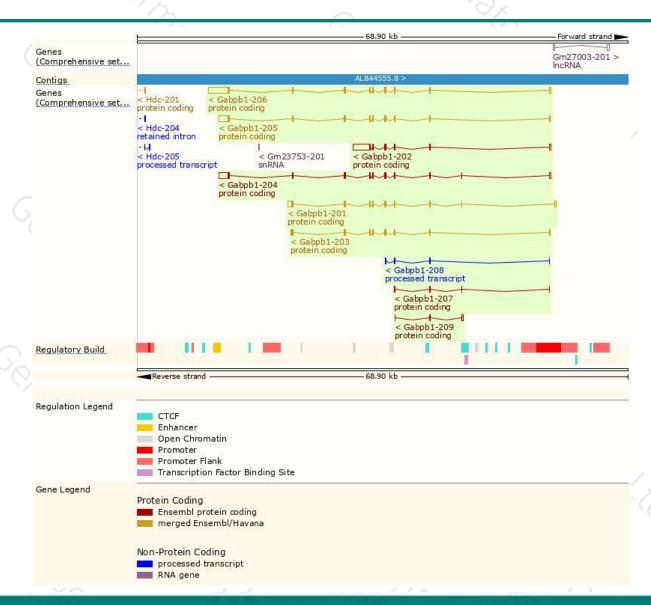
							1
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Gabpb1-206	ENSMUST00000110425.8	4153	383aa	Protein coding	CCDS71137	Q00420	TSL:1 GENCODE basic APPRIS ALT
Gabpb1-202	ENSMUST00000089745.10	3254	233aa	Protein coding	CCDS71135	Q00420	TSL:1 GENCODE basic
Gabpb1-204	ENSMUST00000103227.7	2751	382aa	Protein coding	CCDS16685	Q00420 Q3US22	TSL:1 GENCODE basic APPRIS P3
Gabpb1-205	ENSMUST00000110424.8	2613	382aa	Protein coding	CCDS16685	Q00420 Q3US22	TSL:1 GENCODE basic APPRIS P3
Gabpb1-201	ENSMUST00000039978.12	1991	347aa	Protein coding	CCDS71136	<u>Q3UT69</u>	TSL:1 GENCODE basic
Gabpb1-203	ENSMUST00000103226.9	1338	348aa	Protein coding	CCDS16686	Q00420	TSL:1 GENCODE basic
Gabpb1-209	ENSMUST00000137335.1	439	<u>46aa</u>	Protein coding	-:	A2AQ72	CDS 3' incomplete TSL:3
Gabpb1-207	ENSMUST00000124972.7	372	<u>72aa</u>	Protein coding	29	A2AQ71	CDS 3' incomplete TSL:3
Gabpb1-208	ENSMUST00000130263.1	423	No protein	Processed transcript	-	859	TSL:3
	1111	/ / \			1. 2" 200	A. V. aug.	7 ; ;

The strategy is based on the design of *Gabpb1-206* transcript, the transcription is shown below:



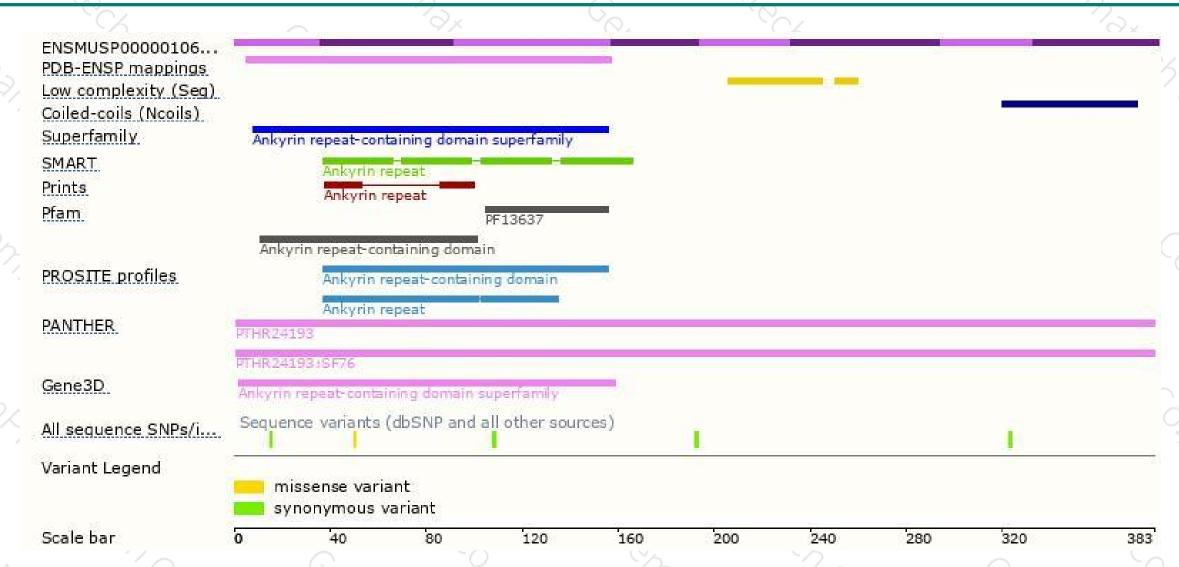
Genomic location distribution





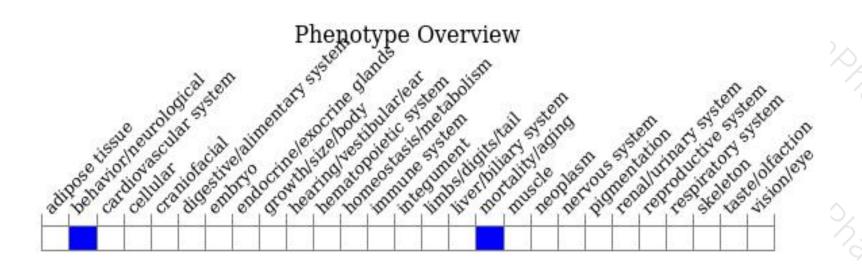
Protein domain





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, mice homozygous for a null allele exhibit embryonic lethality by E12.5.



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





