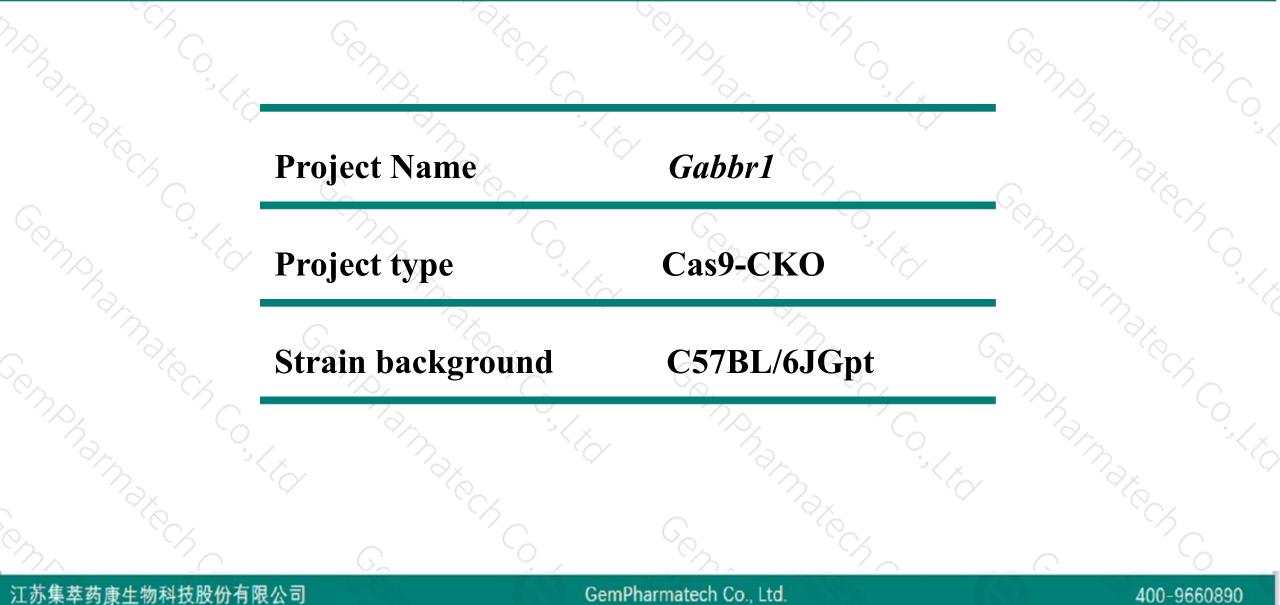


Gabbr1 Cas9-CKO Strategy

Designer: Xiaojing Li Design Date: 2019-9-19 Reviewer: JiaYu

Project Overview

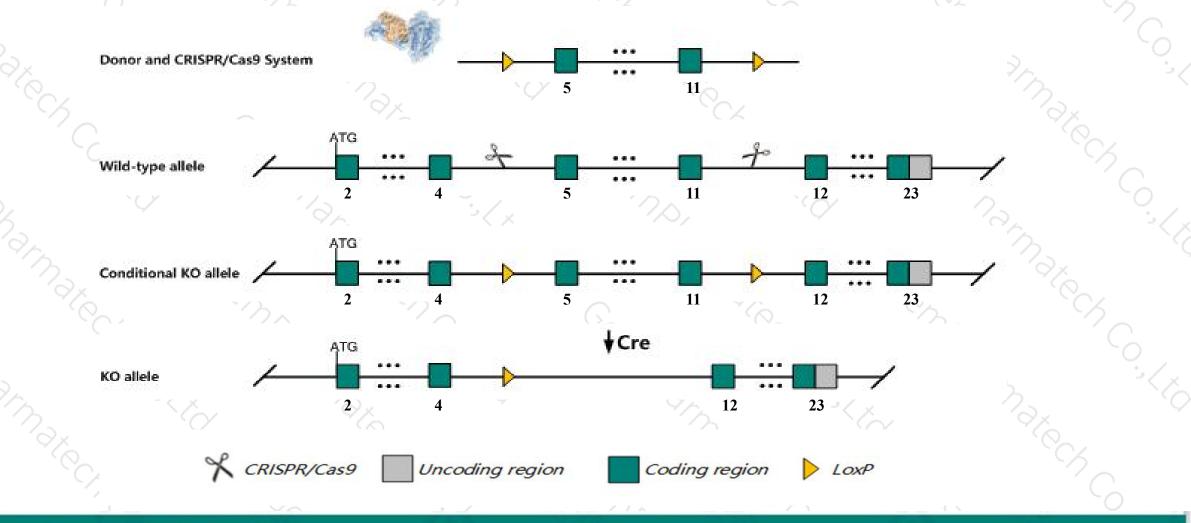




Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Gabbr1 gene. The schematic diagram is as follows:



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The Gabbr1 gene has 11 transcripts. According to the structure of Gabbr1 gene, exon5-exon11 of Gabbr1-201 (ENSMUST00000025338.15) transcript is recommended as the knockout region. The region contains 848bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Gabbr1* 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, Phenotypes of null mice vary depending on strain background and allele. Homozygous null mice may display seizures, premature death, and abnormal nervous system electrophysiology.
- The Gabbr1 gene is located on the Chr17. 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)



< ?

Gabbr1 gamma-aminobutyric acid (GABA) B receptor, 1 [Mus musculus (house mouse)]

Gene ID: 54393, updated on 19-Mar-2019

Summary

Official Symbol	Gabbr1 provided by MGI
Official Full Name	gamma-aminobutyric acid (GABA) B receptor, 1 provided by MGI
Primary source	MGI:MGI:1860139
See related	Ensembl:ENSMUSG0000024462
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	GABAB1, GABAbR1, bM573K1.1
Expression	Broad expression in cortex adult (RPKM 73.7), frontal lobe adult (RPKM 72.2) and 20 other tissues See more
Orthologs	human all

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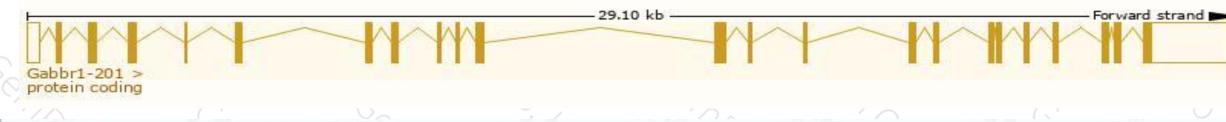
Transcript information (Ensembl)



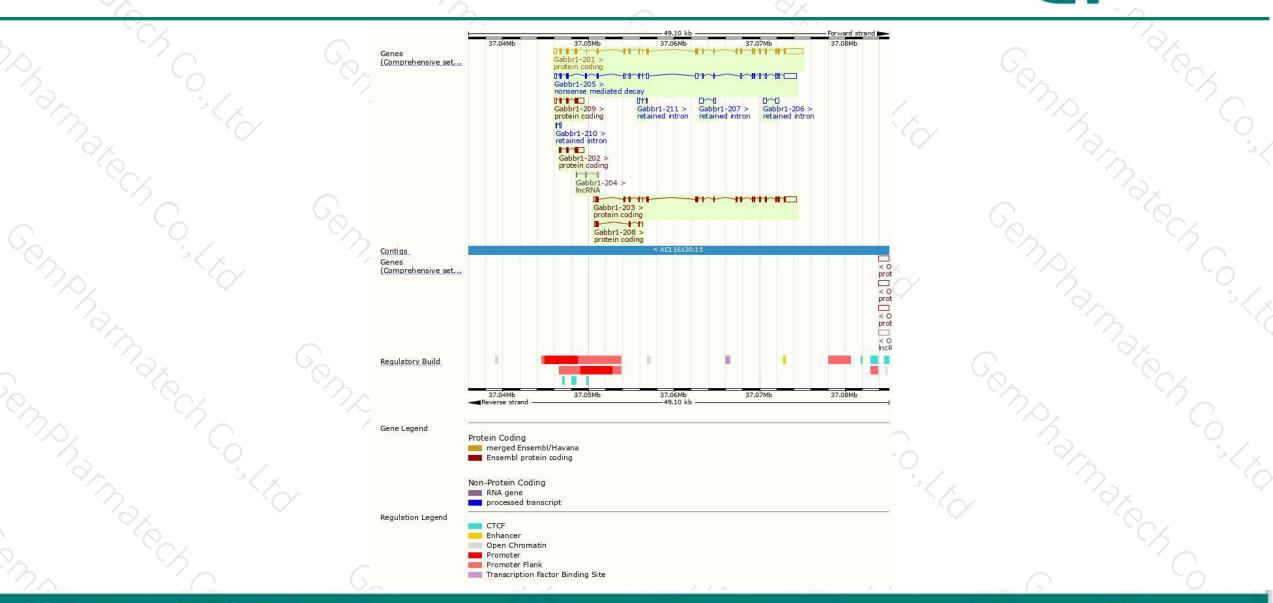
The gene has 11 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Gabbr1-201	ENSMUST00000025338.15	5248	<u>960aa</u>	Protein coding	CCDS37616	Q9WV18	TSL:1 GENCODE basic APPRIS P
Gabbr1-203	ENSMUST00000172792.7	4036	<u>844aa</u>	Protein coding	8.	Q9WV18	TSL:1 GENCODE basic
Gabbr1-209	ENSMUST00000174456.7	1595	<u>229aa</u>	Protein coding	3 4	Q3TWR2	TSL:1 GENCODE basic
Gabbr1-202	ENSMUST00000172789.1	1460	<u>229aa</u>	Protein coding	1 (2	Q3TWR2	TSL:1 GENCODE basic
Gabbr1-208	ENSMUST00000174347.1	821	<u>213aa</u>	Protein coding	17	G3UZ52	CDS 3' incomplete TSL:5
Gabbr1-205	ENSMUST00000173823.7	4204	<u>177aa</u>	Nonsense mediated decay	. 80	G3UXR6	TSL:1
Gabbr1-206	ENSMUST00000174071.1	716	No protein	Retained intron	8 4	-	TSL:3
Gabbr1-207	ENSMUST00000174181.1	608	No protein	Retained intron	62	22	TSL:2
Gabbr1-211	ENSMUST00000174866.1	378	No protein	Retained intron	65	7	TSL:5
Gabbr1-210	ENSMUST00000174826.1	237	No protein	Retained intron	19		TSL:2
Gabbr1-204	ENSMUST00000173564.1	303	No protein	IncRNA	84	-	TSL:5
			- / / N	/ >		N 31 - 24 mg	

The strategy is based on the design of Gabbr1-201 transcript, The transcription is shown below



Genomic location distribution



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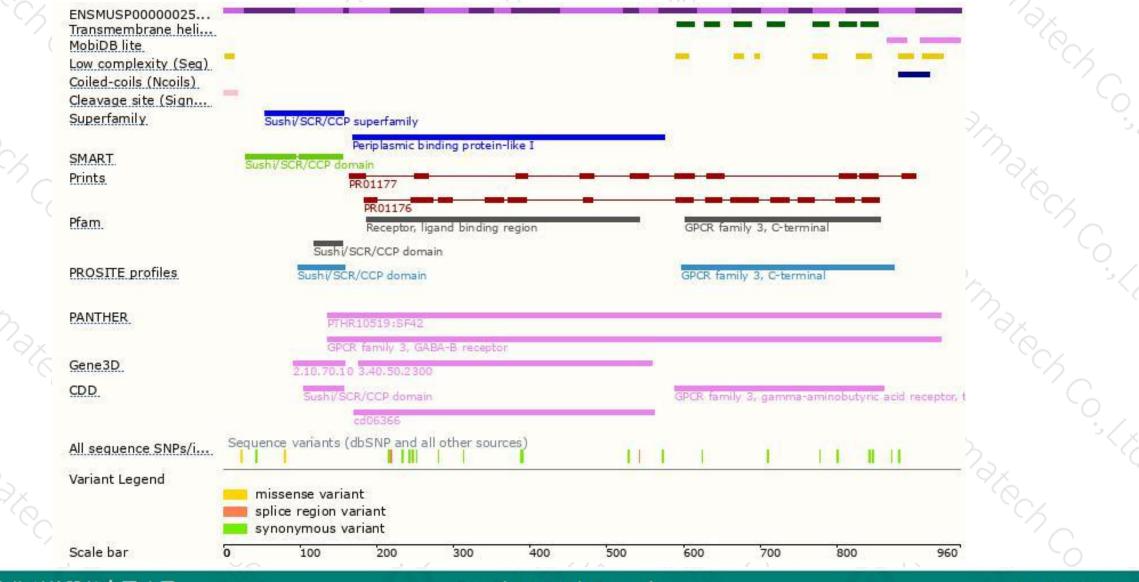
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Protein domain





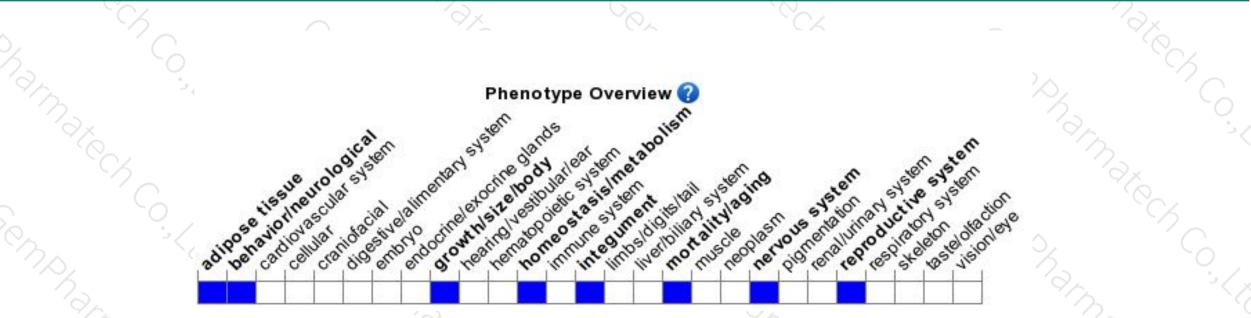
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Mouse phenotype description(MGI)





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, Phenotypes of null mice vary depending on strain background and allele. Homozygous null mice may display seizures, premature death, and abnormal nervous system electrophysiology.



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



