Bcat2 Cas9-KO Strategy Ronnohamakech Co.

Designer: Condata de Ch

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Project Overview



Project Name

Bcat2

Project type

Cas9-KO

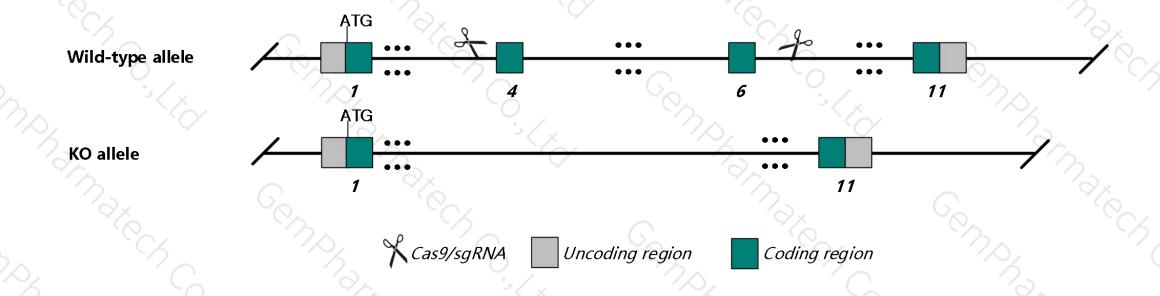
Animal background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Bcat2 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Bcat2* gene has 8 transcripts. According to the structure of *Bcat2* gene, exon4-exon6 of *Bcat2*-201 transcript is recommended as the knockout region. The region contains 395bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Bcat2* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- According to the existing MGI data, The metabolism of branched chain amino acid is impaired in homozygous null mice, resulting in a phenotype similar to human maple syrup urine disease. Mutants exhibit a failure to thrive and die prematurely, though the severity of the symptoms can be ameliorated with a restricted diet.
- ➤ The *Bcat2* 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)



Bcat2 branched chain aminotransferase 2, mitochondrial [Mus musculus (house mouse)]

Gene ID: 12036, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Bcat2 provided by MGI

Official Full Name branched chain aminotransferase 2, mitochondrial provided by MGI

Primary source MGI:MGI:1276534

See related Ensembl: ENSMUSG00000030826

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 Eca40; Bcat-2; Bcat(m)

Expression Ubiquitous expression in adrenal adult (RPKM 36.2), subcutaneous fat pad adult (RPKM 34.2) and 28 other tissues See more

Orthologs human all

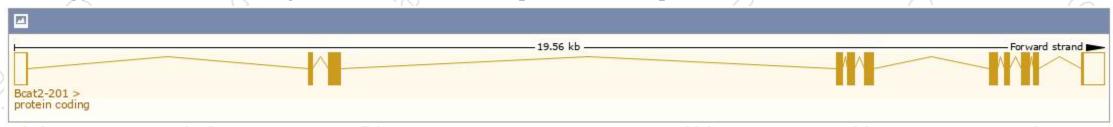
Transcript information (Ensembl)



The gene has 8 transcripts, and all transcripts are shown below:

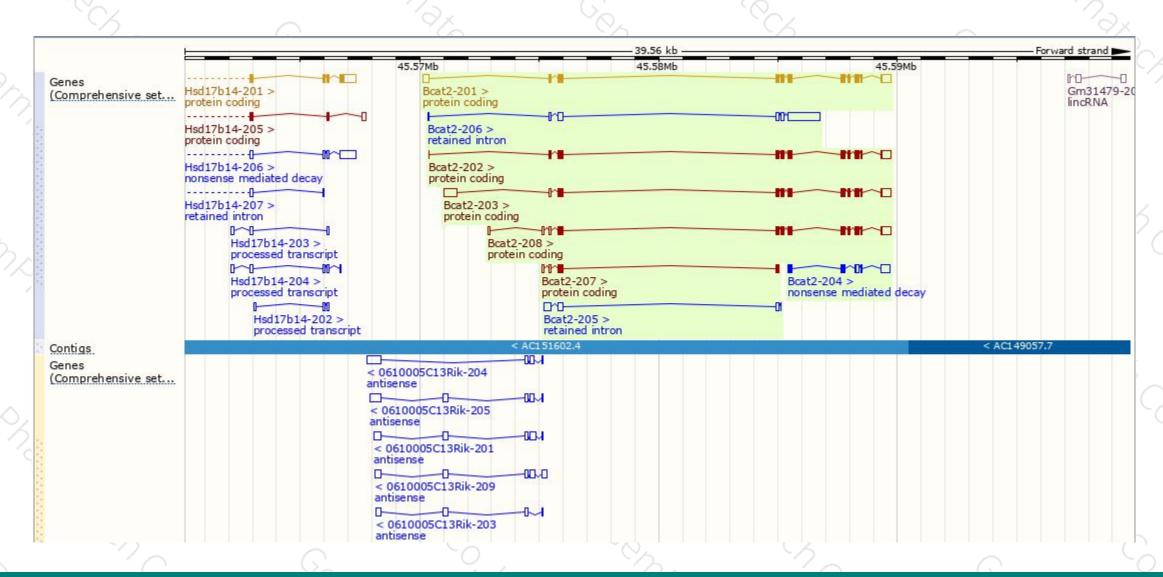
Name 4	Transcript ID	bp 🌢	Protein &	Biotype	CCDS	UniProt 4	Flags
Bcat2-203	ENSMUST00000209204.1	2081	353aa	Protein coding	CCDS57549 ₽	<u>088374</u> ₽	TSL:1 GENCODE basic
3cat2-201	ENSMUST00000033098.15	1772	<u>393aa</u>	Protein coding	CCDS21252₽	<u>035855</u> 윤 <u>Q3ULU3</u> 윤	TSL:1 GENCODE basic APPRIS P
Bcat2-208	ENSMUST00000211173.1	1684	<u>353aa</u>	Protein coding	CCDS57549₽	<u>088374</u> ₽	TSL:1 GENCODE basic
3cat2-202	ENSMUST00000120864.9	1536	388aa	Protein coding	9.7	A0A1B0GX27₺	TSL:5 GENCODE basic
3cat2-207	ENSMUST00000210811.1	486	<u>97aa</u>	Protein coding	5.00	A0A1B0GST1 ₽	CDS 3' incomplete TSL:3
Bcat2-204	ENSMUST00000209410.1	917	<u>101aa</u>	Nonsense mediated decay	(*)	A0A1B0GQY4₽	CDS 5' incomplete TSL:5
Bcat2-206	ENSMUST00000209569.1	1905	No protein	Retained intron	-	-	TSL:1
Bcat2-205	ENSMUST00000209543.1	632	No protein	Retained intron	-	-	TSL:2

The strategy is based on the design of *Bcat2-201* transcript, The transcription is shown below



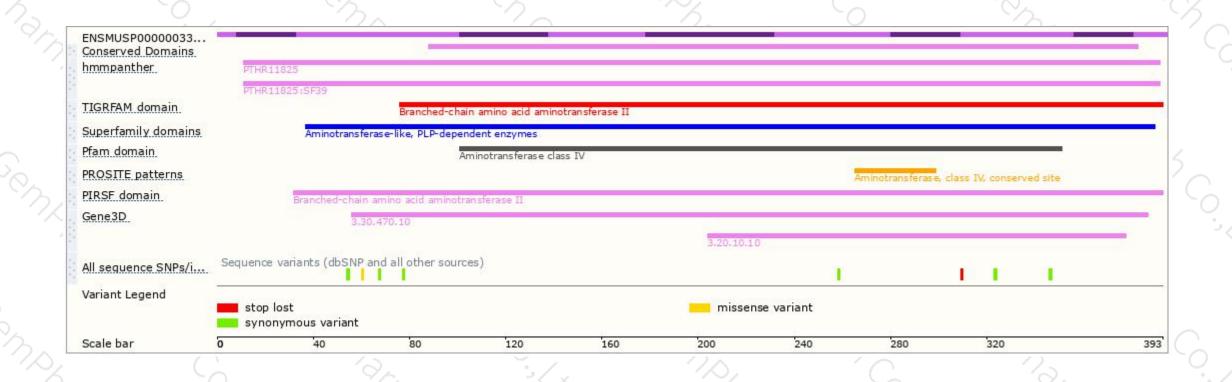
Genomic location distribution





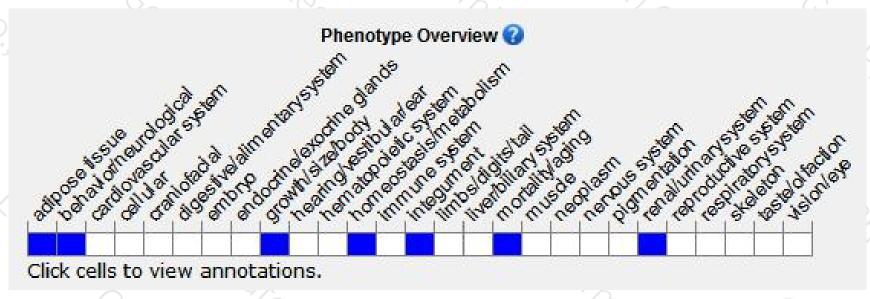
Protein domain





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, The metabolism of branched chain amino acid is impaired in homozygous null mice, resulting in a phenotype similar to human maple syrup urine disease. Mutants exhibit a failure to thrive and die prematurely, though the severity of the symptoms can be ameliorated with a restricted diet.

If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





