

Slc38a3 Cas9-KO Strategy

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Design Date: 2020-4-28

Project Overview



Project Name

Slc38a3

Project type

Cas9-KO

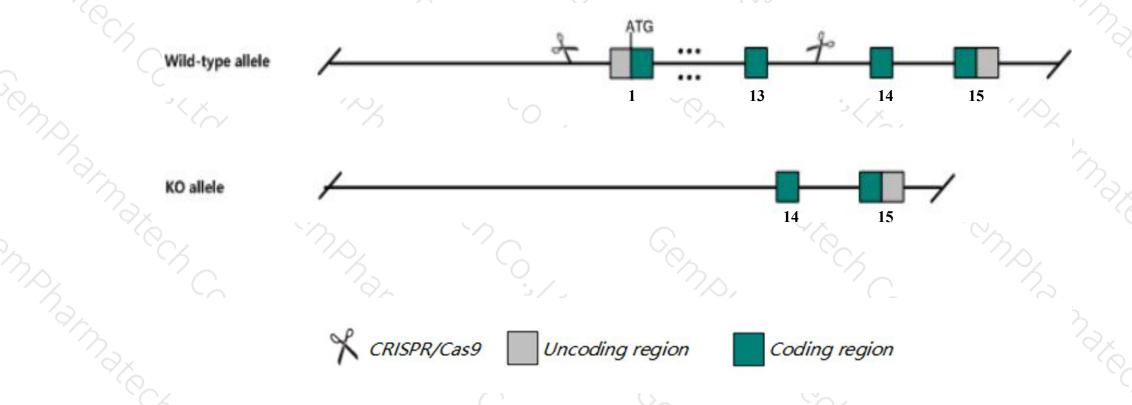
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc38a3 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Slc38a3* gene has 14 transcripts. According to the structure of *Slc38a3* gene, exon1-exon13 of *Slc38a3-202* (ENSMUST00000167868.7) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify Slc38a3 gene. The brief process is as follows: CRISPR/Cas9 syste

Notice



- ➤ According to the existing MGI data, mice homozygous for an enu allele exhibit complete postnatal lethality between p18 and p20, altered amino acid levels in the serum, liver and brain, and decreased ammonia excretion.
- The *Slc38a3* gene is located on the Chr9. 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)



Slc38a3 solute carrier family 38, member 3 [Mus musculus (house mouse)]

Gene ID: 76257, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Slc38a3 provided by MGI

Official Full Name solute carrier family 38, member 3 provided by MGI

Primary source MGI:MGI:1923507

See related Ensembl: ENSMUSG00000010064

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 Sn1; Nat1; mNAT; Snat3; D9Ucla2; Slc38-3; 0610012J02Rik

Expression Biased expression in liver adult (RPKM 1073.0), liver E18 (RPKM 123.2) and 1 other tissue See more

Orthologs human all

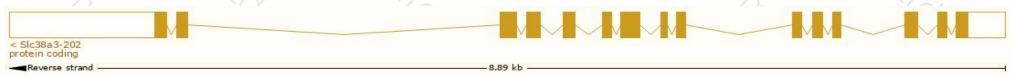
Transcript information (Ensembl)



The gene has 14 transcripts, all transcripts are shown below:

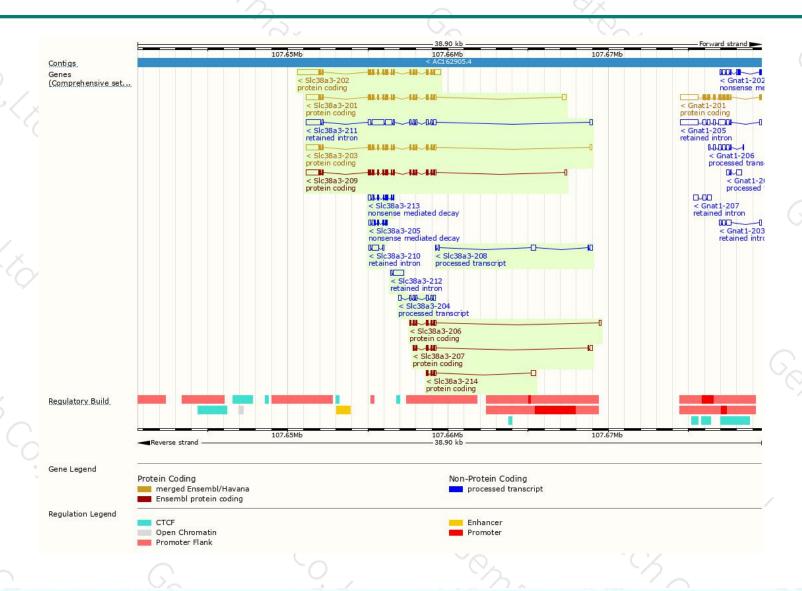
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ragio leads	Name 🍦	Transcript ID	bp 🍦	Protein 🍦	Biotype	CCDS	UniProt	Flags	
	SIc38a3-202	ENSMUST00000167868.7	3151	<u>505aa</u>	Protein coding	CCDS23503 ₺	Q9DCP2₽	TSL:1 GENCODE basic APPRIS P1	
	SIc38a3-201	ENSMUST00000010208.13	2585	<u>505aa</u>	Protein coding	CCDS23503₽	Q9DCP2₽	TSL:1 GENCODE basic APPRIS P1	
	SIc38a3-209	ENSMUST00000193932.5	2463	<u>505aa</u>	Protein coding	CCDS23503 ₽	Q9DCP2₽	TSL:1 GENCODE basic APPRIS P1	
	SIc38a3-203	ENSMUST00000177567.7	2456	<u>505aa</u>	Protein coding	CCDS23503₽	Q9DCP2₽	TSL:1 GENCODE basic APPRIS P1	Ī
	SIc38a3-206	ENSMUST00000192323.5	696	<u>181aa</u>	Protein coding	\$ 5 72	<u>A0A0A6YX60</u> ₽	CDS 3' incomplete TSL:3	
	SIc38a3-207	ENSMUST00000192990.5	691	<u>154aa</u>	Protein coding		A0A0A6YWF5₺	CDS 3' incomplete TSL:3	
	SIc38a3-214	ENSMUST00000195843.1	638	<u>99aa</u>	Protein coding	(-)	A0A0A6YWH4₽	CDS 3' incomplete TSL:3	
	SIc38a3-213	ENSMUST00000195739.1	703	<u>160aa</u>	Nonsense mediated decay	7 - 7/	A0A0A6YWC0@	CDS 5' incomplete TSL:3	
	SIc38a3-205	ENSMUST00000192211.1	641	<u>125aa</u>	Nonsense mediated decay	(44)	A0A0A6YWI8@	CDS 5' incomplete TSL:3	
	SIc38a3-204	ENSMUST00000191923.1	731	No protein	Processed transcript	8200	2	TSL:3	
	SIc38a3-208	ENSMUST00000193495.1	624	No protein	Processed transcript	120	2	TSL:3	
	SIc38a3-211	ENSMUST00000194895.5	3133	No protein	Retained intron		6	TSL:2	
	SIc38a3-212	ENSMUST00000195033.1	780	No protein	Retained intron	232		TSL:2	
	SIc38a3-210	ENSMUST00000194230.1	614	No protein	Retained intron		-	TSL:5	
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The strategy is based on the design of Slc38a3-202 transcript, the transcription is shown below



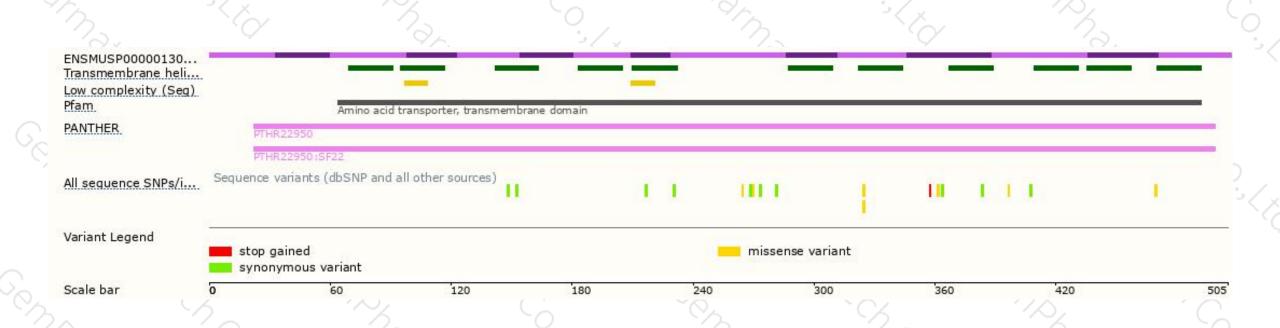
Genomic location distribution





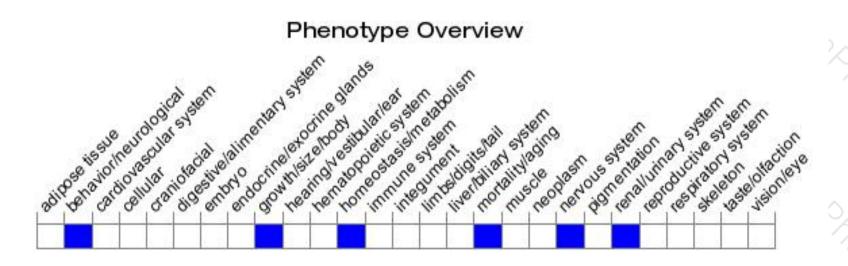
Protein domain





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 an ENU allele exhibit complete postnatal lethality between P18 and P20, altered amino acid levels in the serum, liver and brain, and decreased ammonia excretion.



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





