

# ***Slc25a12*** Cas9-KO Strategy

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# Project Overview

**Project Name**

***Slc25a12***

**Project type**

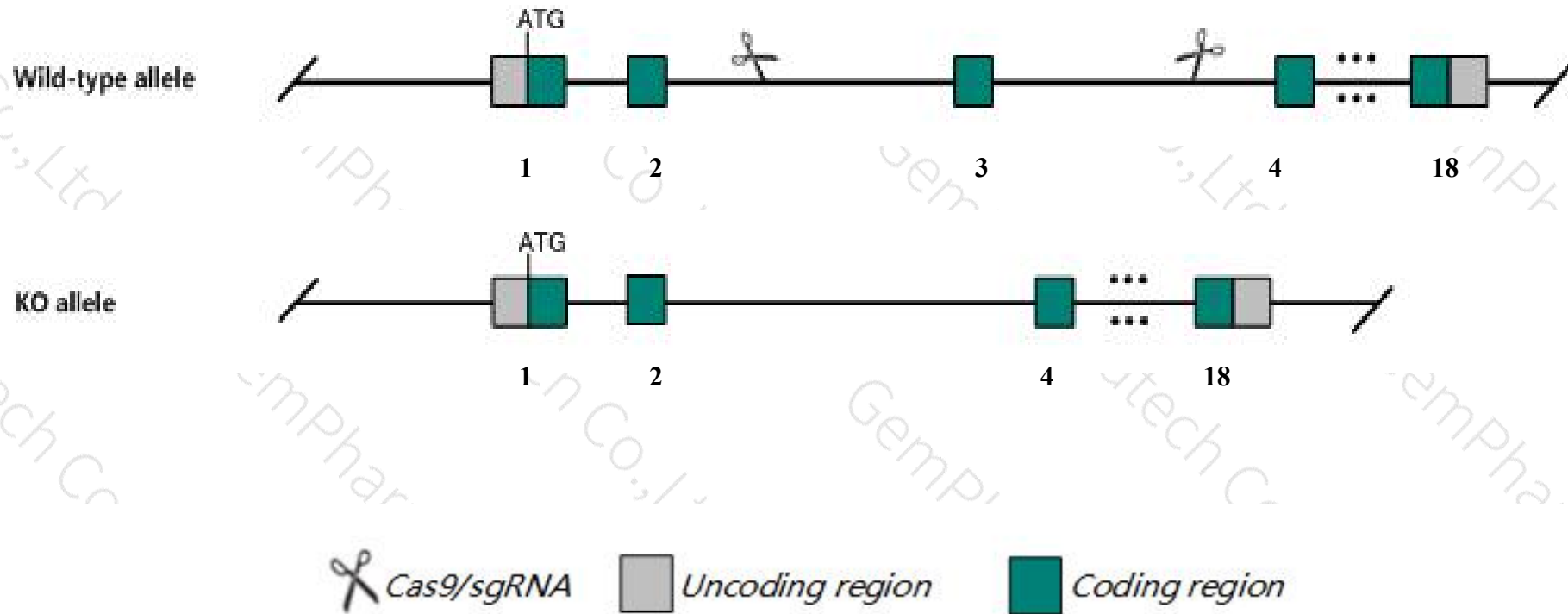
**Cas9-KO**

**Strain background**

**C57BL/6J**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slc25a12* gene. The schematic diagram is as follows:



- The *Slc25a12* gene has 7 transcripts. According to the structure of *Slc25a12* gene, exon3 of *Slc25a12-206* (ENSMUST00000151937.7) transcript is recommended as the knockout region. The region contains 143bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc25a12* 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.

- According to the existing MGI data, Mice homozygous for a null allele show severe growth defects, generalized tremors, postnatal lethality, impaired motor coordination, and CNS dysmyelination associated with decreased synthesis of myelin lipids and a striking reduction in brain aspartate and N-acetylaspartate levels.
- The *Slc25a12* 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 gene transcription and translation processes, all risks cannot be predicted under existing information.



# Gene information (NCBI)

## Slc25a12 solute carrier family 25 (mitochondrial carrier, Aralar), member 12 [Mus musculus (house mouse)]

Gene ID: 78830, updated on 7-Apr-2019

### Summary

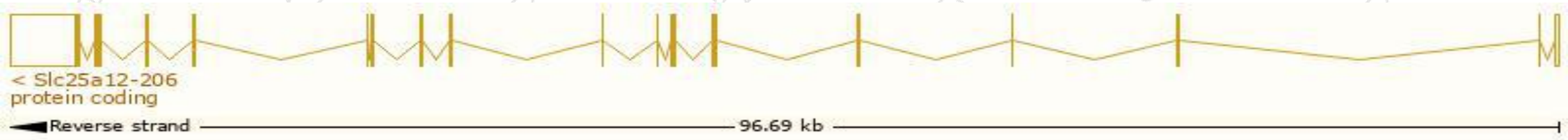
Official Symbol	Slc25a12 provided by <a href="#">MGI</a>
Official Full Name	solute carrier family 25 (mitochondrial carrier, Aralar), member 12 provided by <a href="#">MGI</a>
Primary source	<a href="#">MGI:MGI:1926080</a>
See related	<a href="#">Ensembl:ENSMUSG00000027010</a>
Gene type	protein coding
RefSeq status	VALIDATED
Organism	<a href="#">Mus musculus</a>
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	2610002D09Rik, A1839531, B230107K20Rik, BB129864
Expression	Ubiquitous expression in heart adult (RPKM 17.8), cerebellum adult (RPKM 11.2) and 27 other tissues <a href="#">See more</a>
Orthologs	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

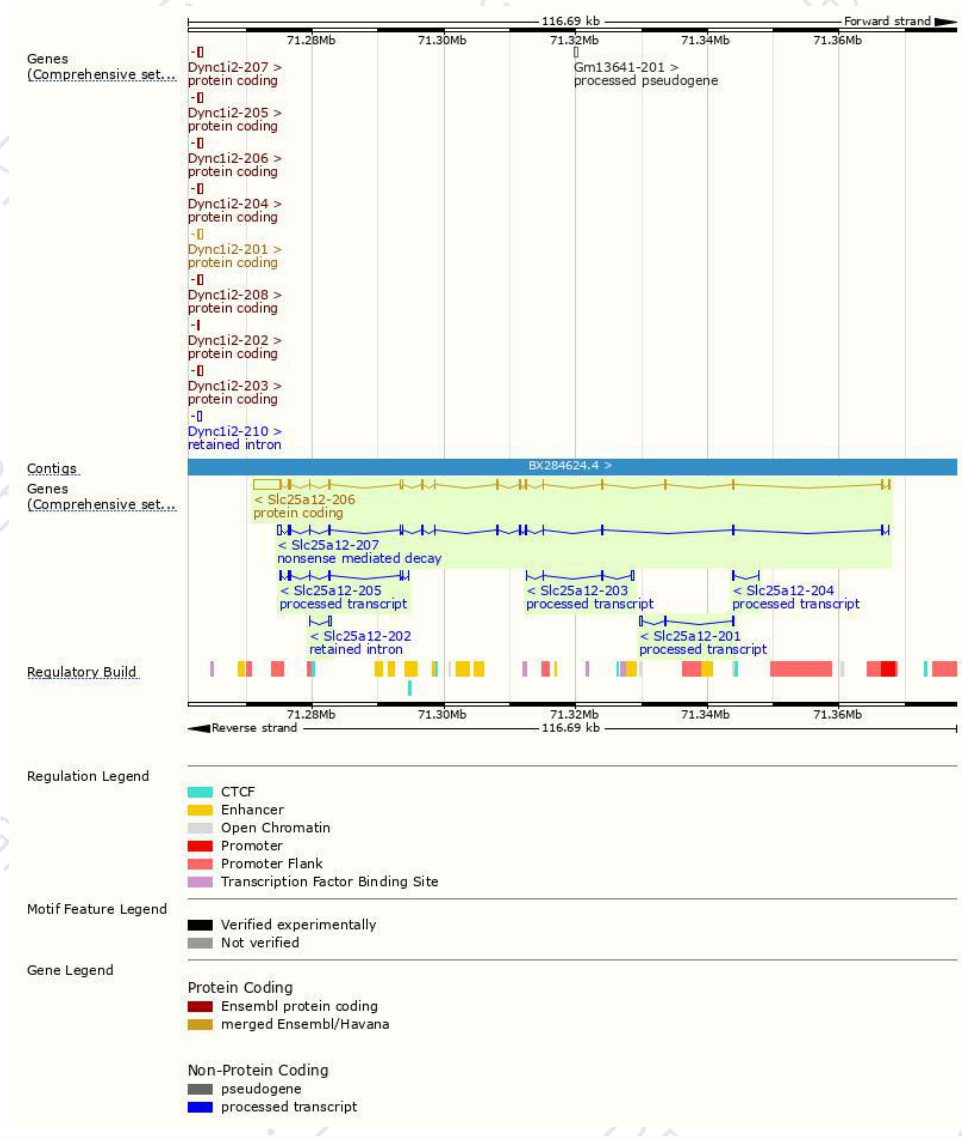
The gene has 7 transcript,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Slc25a12-206	<a href="#">ENSMUST00000151937.7</a>	6351	<a href="#">677aa</a>	Protein coding	<a href="#">CCDS16113</a>	<a href="#">Q8BH59</a>	TSL:1 GENCODE basic APPRIS P1
Slc25a12-207	<a href="#">ENSMUST00000184169.7</a>	2392	<a href="#">87aa</a>	Nonsense mediated decay	-	<a href="#">V9GXX9</a>	TSL:5
Slc25a12-205	<a href="#">ENSMUST00000147553.1</a>	994	No protein	Processed transcript	-	-	TSL:3
Slc25a12-203	<a href="#">ENSMUST00000137916.1</a>	790	No protein	Processed transcript	-	-	TSL:2
Slc25a12-201	<a href="#">ENSMUST00000126493.7</a>	550	No protein	Processed transcript	-	-	TSL:5
Slc25a12-204	<a href="#">ENSMUST00000146653.1</a>	204	No protein	Processed transcript	-	-	TSL:5
Slc25a12-202	<a href="#">ENSMUST00000130715.1</a>	414	No protein	Retained intron	-	-	TSL:1

The strategy is based on the design of *Slc25a12-206* transcript,The transcription is shown below

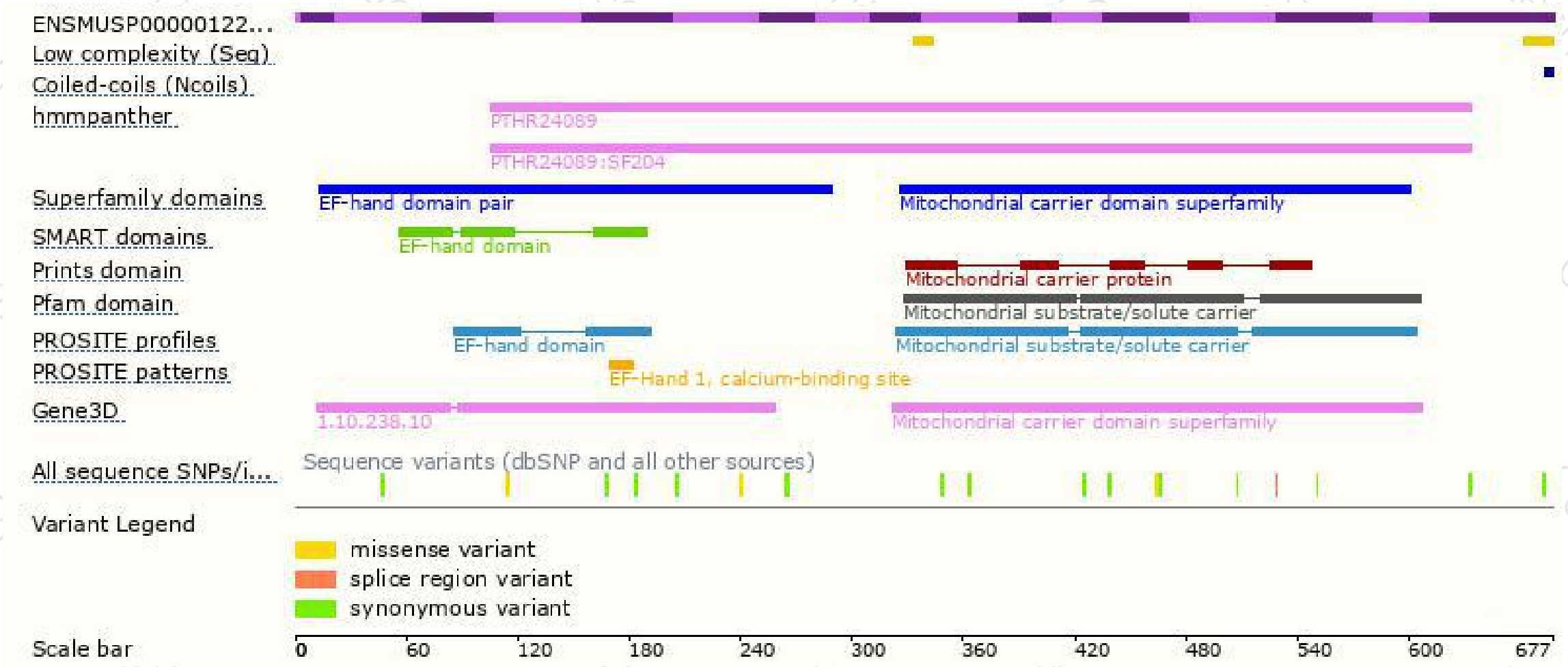


# Genomic location distribution

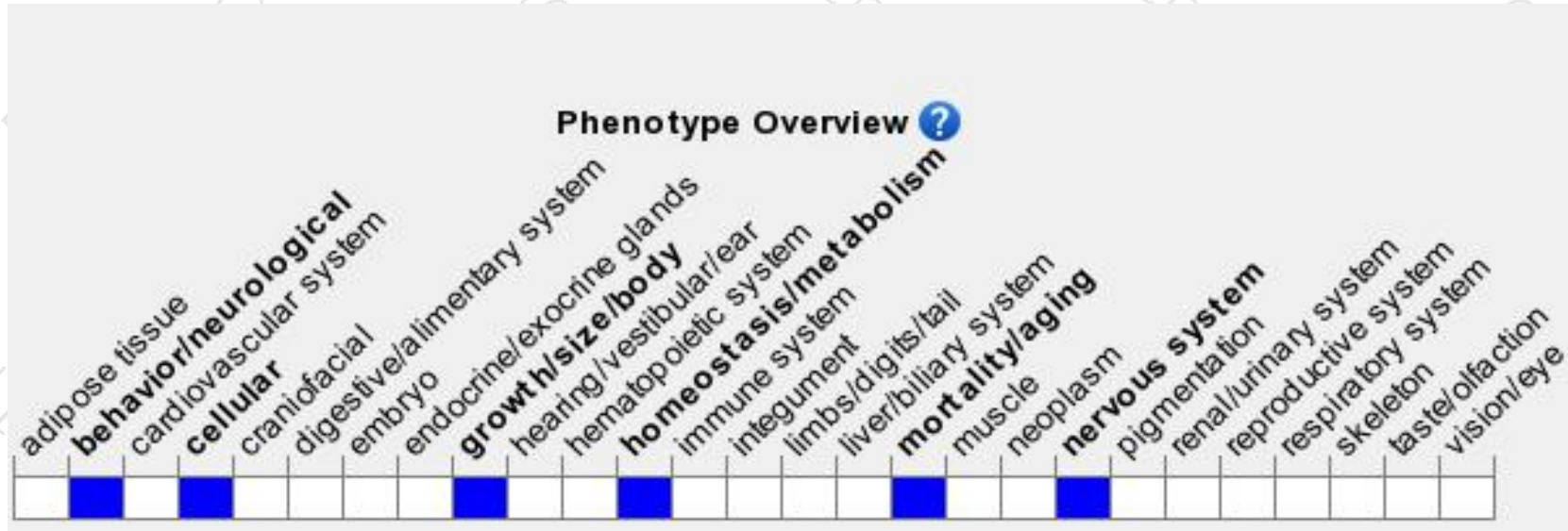




# Protein domain



# 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 show severe growth defects, generalized tremors, postnatal lethality, impaired motor coordination, and CNS dysmyelination associated with decreased synthesis of myelin lipids and a striking reduction in brain aspartate and N-acetylaspartate levels.

If you have any questions, you are welcome to inquire.

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