

# *Slc23a1* Cas9-KO Strategy

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# Project Overview

**Project Name**

***Slc23a1***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slc23a1* gene. The schematic diagram is as follows:



- The *Slc23a1* gene has 4 transcripts. According to the structure of *Slc23a1* gene, exon2-exon13 of *Slc23a1-201* (ENSMUST00000025212.7) transcript is recommended as the knockout region. The region contains 1534bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc23a1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a knock-out allele exhibit abnormal ascorbate homeostasis and early postnatal lethality associated with lethargy and lack of gastric milk. Heterozygous mice of homozygous dams exhibit a similar phenotype.
- The *Slc23a1* gene is located on the Chr18. 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)

## Slc23a1 solute carrier family 23 (nucleobase transporters), member 1 [ *Mus musculus* (house mouse) ]

Gene ID: 20522, updated on 10-Oct-2019

### Summary

<b>Official Symbol</b>	Slc23a1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	solute carrier family 23 (nucleobase transporters), member 1 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1341903</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000024354</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	SVCT1; YSPL3; Slc23a2; D18Ucla2
<b>Expression</b>	Biased expression in kidney adult (RPKM 78.2), liver adult (RPKM 28.3) and 6 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

### Genomic context

**Location:** 18 B2; 18 19.17 cM

See Slc23a1 in [Genome Data Viewer](#)

**Exon count:** 19

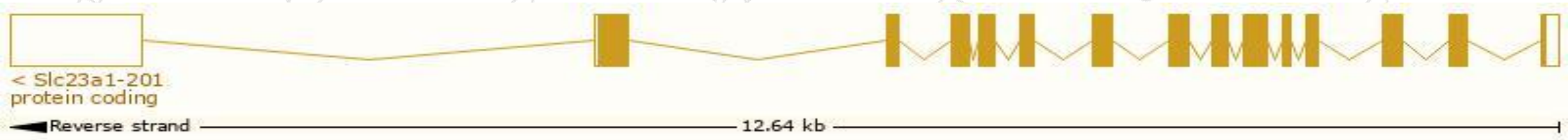
Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCm38.p6 ( <a href="#">GCF_000001635.26</a> )	18	NC_000084.6 (35604224..35629845, complement)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	18	NC_000084.5 (35774258..35786881, complement)

# Transcript information (Ensembl)

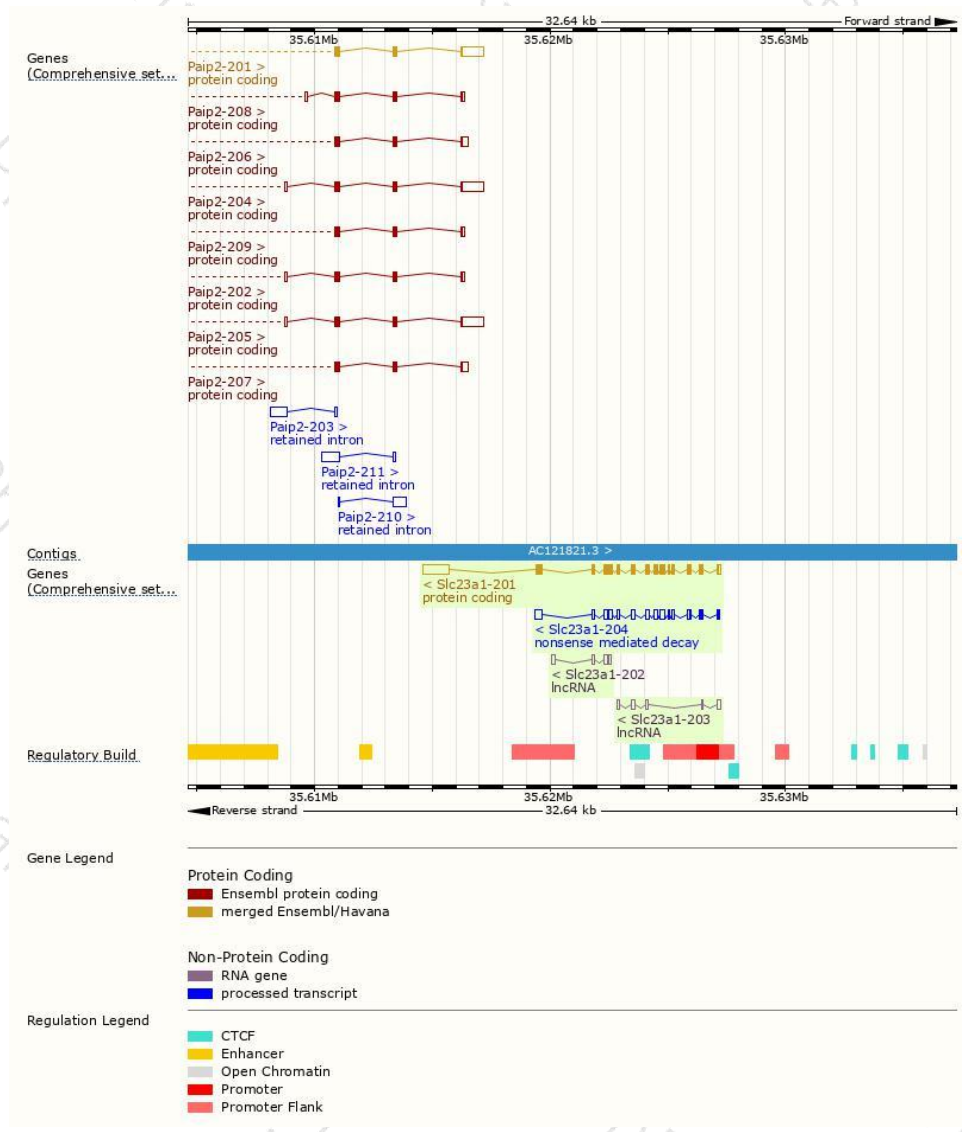
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Slc23a1-201	<a href="#">ENSMUST00000025212.7</a>	3026	<a href="#">605aa</a>	Protein coding	<a href="#">CCDS29144</a>	<a href="#">Q9Z2J0</a>	TSL:1 GENCODE basic APPRIS P1
Slc23a1-204	<a href="#">ENSMUST00000237305.1</a>	1889	<a href="#">53aa</a>	Nonsense mediated decay	-	<a href="#">D6RDS7</a>	
Slc23a1-203	<a href="#">ENSMUST00000236196.1</a>	565	No protein	lncRNA	-	-	
Slc23a1-202	<a href="#">ENSMUST00000235744.1</a>	498	No protein	lncRNA	-	-	

The strategy is based on the design of *Slc23a1-201* transcript,The transcription is shown below

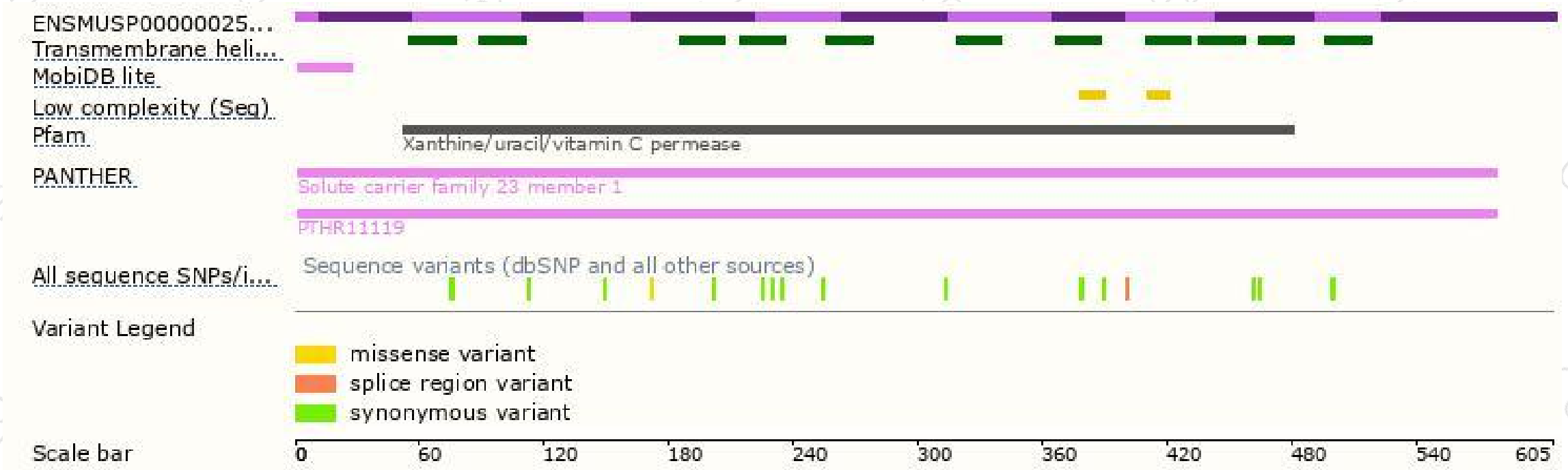


# Genomic location distribution





# 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, Mice homozygous for a knock-out allele exhibit abnormal ascorbate homeostasis and early postnatal lethality associated with lethargy and lack of gastric milk. Heterozygous mice of homozygous dams exhibit a similar phenotype.

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

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