

# *Slco1c1* Cas9-KO Strategy

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# Project Overview

**Project Name**

*Slco1c1*

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slco1c1* gene. The schematic diagram is as follows:



- The *Slco1c1* gene has 6 transcripts. According to the structure of *Slco1c1* gene, exon3-exon14 of *Slco1c1-201* (ENSMUST00000032362.11) transcript is recommended as the knockout region. The region contains 1796bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slco1c1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a knock-out allele exhibit decreased thyroxine and triiodothyronine levels in the forebrain, in the absence of overt growth, reproductive or neurological abnormalities.
- The *Slco1c1* gene is located on the Chr6. 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)

## Slco1c1 solute carrier organic anion transporter family, member 1c1 [ *Mus musculus* (house mouse) ]

Gene ID: 58807, updated on 5-Oct-2019

### Summary

**Official Symbol** Slco1c1 provided by [MGI](#)  
**Official Full Name** solute carrier organic anion transporter family, member 1c1 provided by [MGI](#)  
**Primary source** [MGI:MGI:1889679](#)  
**See related** [Ensembl:ENSMUSG00000030235](#)  
**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** Oatp2; Oatpf; OATP-F; OATP-14; Slc21a14  
**Expression** Biased expression in frontal lobe adult (RPKM 12.2), cortex adult (RPKM 11.8) and 4 other tissues [See more](#)  
**Orthologs** [human](#) [all](#)

### Genomic context

**Location:** 6; 6 G2

See Slco1c1 in [Genome Data Viewer](#)

**Exon count:** 15

Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCm38.p6 ( <a href="#">GCF_000001635.26</a> )	6	NC_000072.6 (141524354..141570177)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	6	NC_000072.5 (141472907..141518698)

# Transcript information (Ensembl)

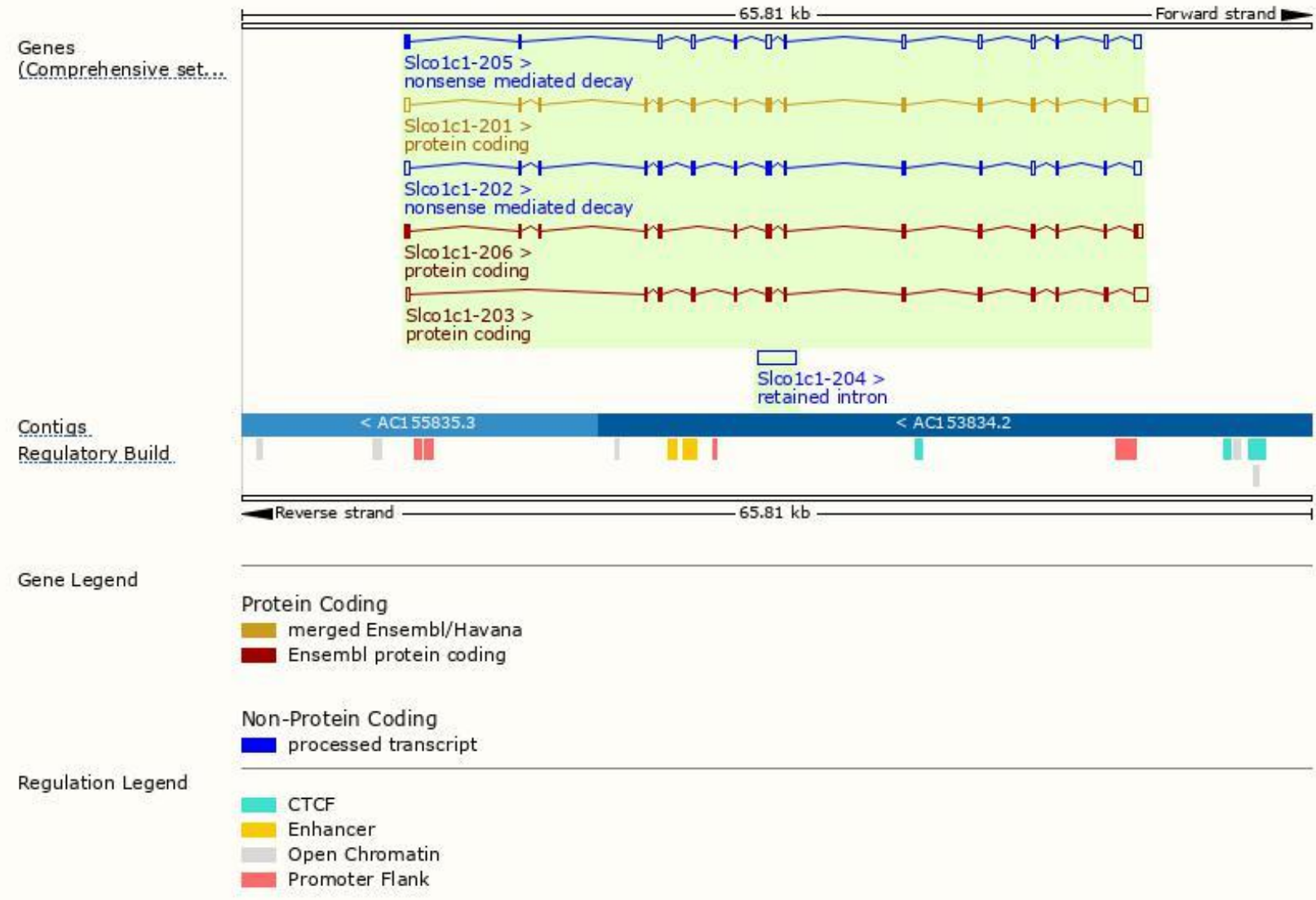
The gene has 6 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Slco1c1-201	<a href="#">ENSMUST00000032362.11</a>	3178	<a href="#">715aa</a>	Protein coding	<a href="#">CCDS20677</a>	<a href="#">Q9ERB5</a>	TSL:1 GENCODE basic APPRIS P2
Slco1c1-203	<a href="#">ENSMUST00000203140.1</a>	2771	<a href="#">544aa</a>	Protein coding	-	<a href="#">A0A0N4SVD7</a>	TSL:5 GENCODE basic APPRIS ALT2
Slco1c1-206	<a href="#">ENSMUST00000205214.2</a>	2515	<a href="#">666aa</a>	Protein coding	-	<a href="#">A0A0N4SUZ6</a>	TSL:5 GENCODE basic
Slco1c1-202	<a href="#">ENSMUST00000135562.7</a>	2680	<a href="#">469aa</a>	Nonsense mediated decay	-	<a href="#">Q66L38</a>	TSL:1
Slco1c1-205	<a href="#">ENSMUST00000204998.2</a>	2427	<a href="#">46aa</a>	Nonsense mediated decay	-	<a href="#">A0A0N4SW25</a>	TSL:5
Slco1c1-204	<a href="#">ENSMUST00000203755.1</a>	2300	No protein	Retained intron	-	-	TSL:NA

The strategy is based on the design of *Slco1c1-201* transcript,The transcription is shown below



# Genomic location distribution

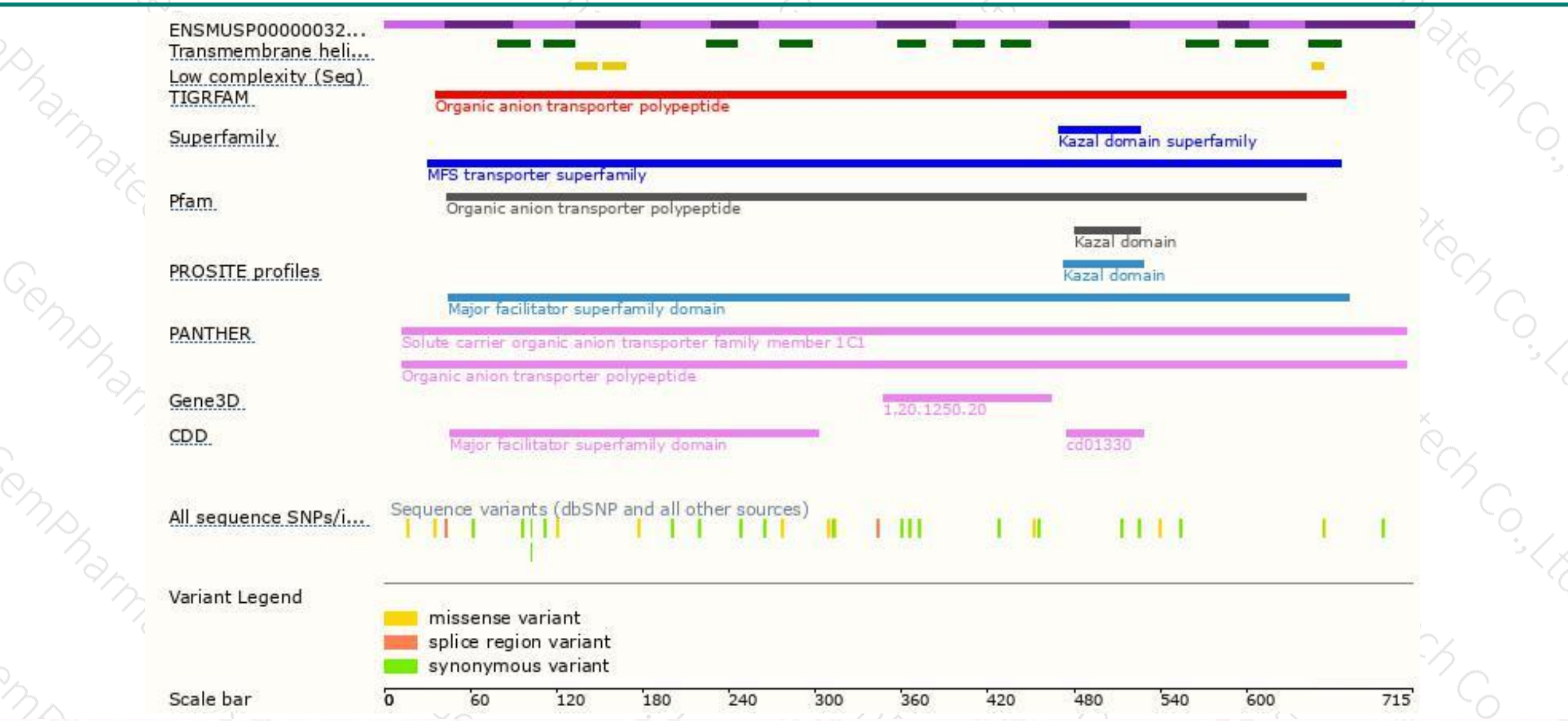




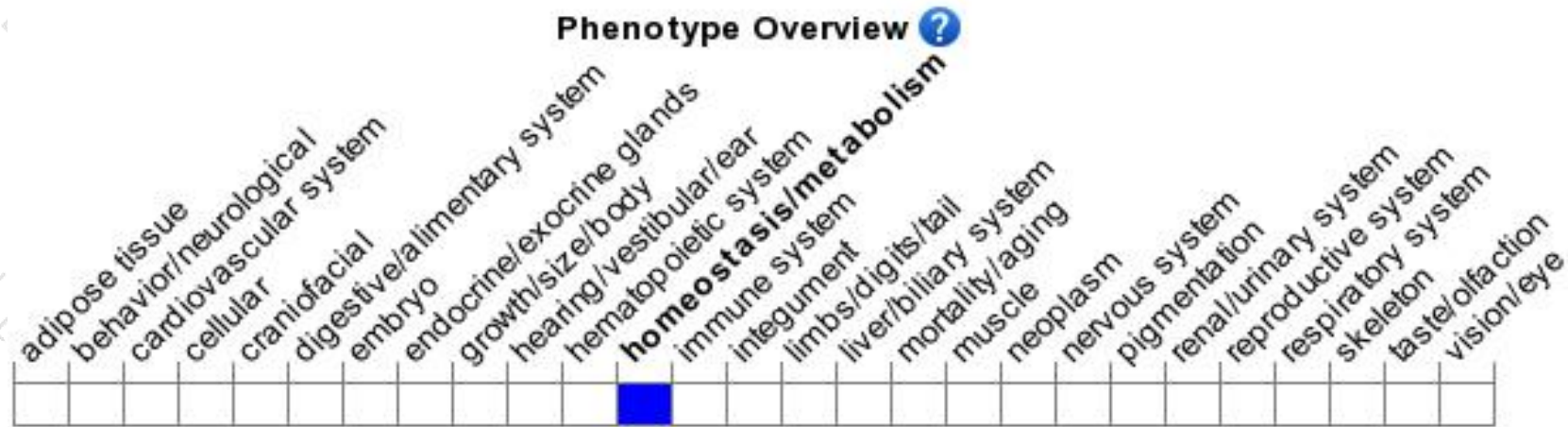
# Protein domain



集萃药康  
GemPharmatech



# 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 decreased thyroxine and triiodothyronine levels in the forebrain, in the absence of overt growth, reproductive or neurological abnormalities.

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

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