

# *Slc34a2* Cas9-KO Strategy

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# Project Overview



**Project Name**

***Slc34a2***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slc34a2* gene. The schematic diagram is as follows:



- The *Slc34a2* gene has 3 transcripts. According to the structure of *Slc34a2* gene, exon2-exon13 of *Slc34a2-201* (ENSMUST00000094787.7) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc34a2* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygous null mice display embryonic lethality, embryonic growth arrest, failure of embryo turning and somitogenesis, impaired placental development and impaired yolk sac vascular remodeling.
- The *Slc34a2* gene is located on the Chr5. 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)

## Slc34a2 solute carrier family 34 (sodium phosphate), member 2 [ *Mus musculus* (house mouse) ]

Gene ID: 20531, updated on 12-Aug-2019

### Summary

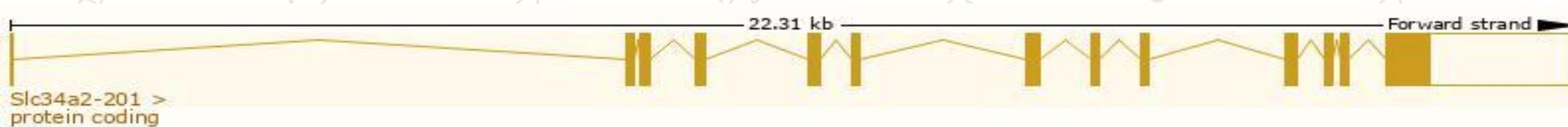
<b>Official Symbol</b>	Slc34a2 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	solute carrier family 34 (sodium phosphate), member 2 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1342284</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000029188</a>
<b>Gene type</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>	Npt2b; NaPi-2b; AA536683; D5Erd227e
<b>Expression</b>	Biased expression in lung adult (RPKM 498.2), large intestine adult (RPKM 142.1) and 1 other tissue <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

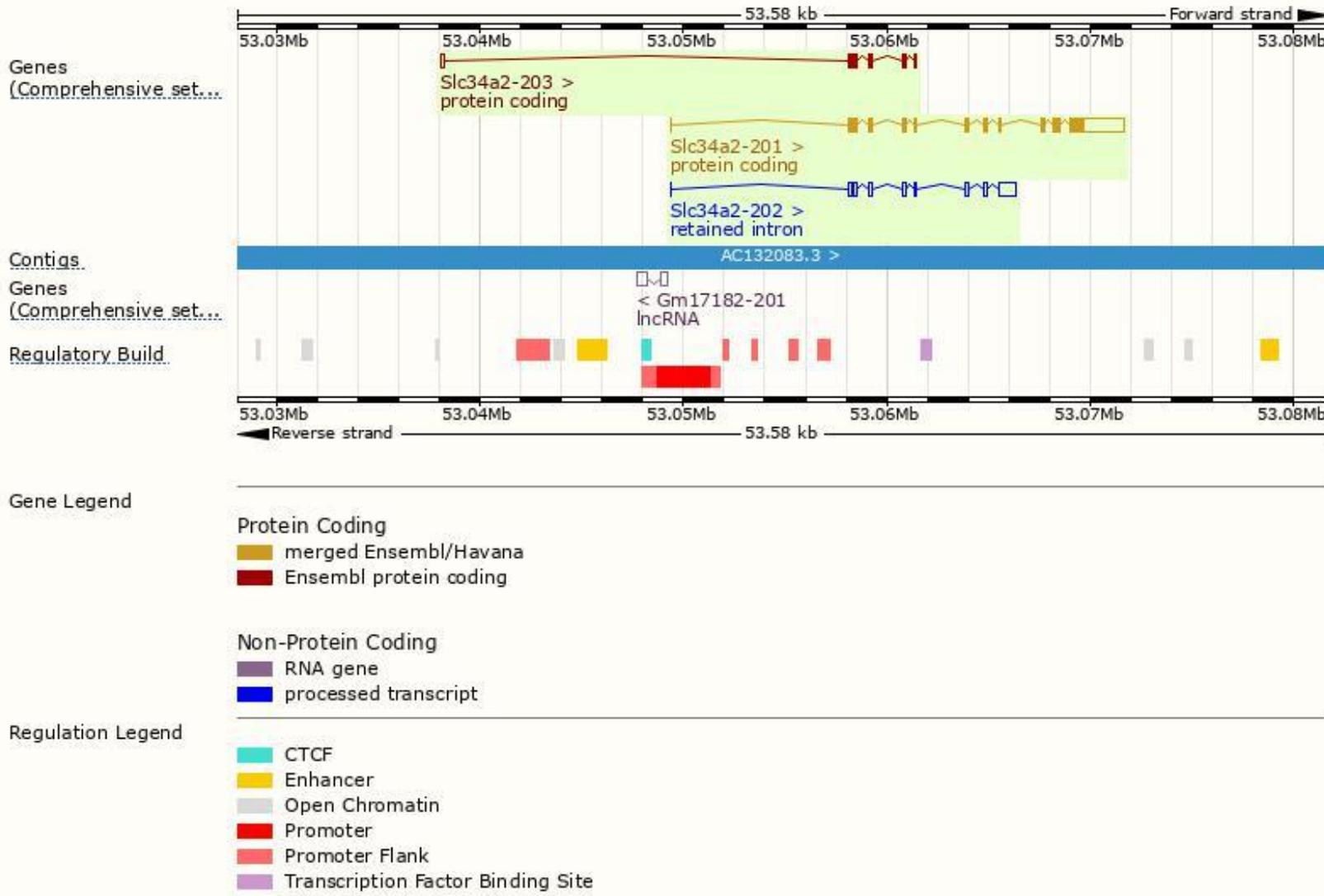
The gene has 3 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
<b>Slc34a2-201</b>	<a href="#">ENSMUST00000094787.7</a>	4176	<a href="#">697aa</a>	Protein coding	<a href="#">CCDS19291</a>	<a href="#">Q9DBP0</a>	TSL:1 GENCODE basic APPRIS P1
<b>Slc34a2-203</b>	<a href="#">ENSMUST00000170523.7</a>	717	<a href="#">186aa</a>	Protein coding	-	<a href="#">E9QAX5</a>	CDS 3' incomplete TSL:3
<b>Slc34a2-202</b>	<a href="#">ENSMUST00000147243.3</a>	1778	No protein	Retained intron	-	-	TSL:1

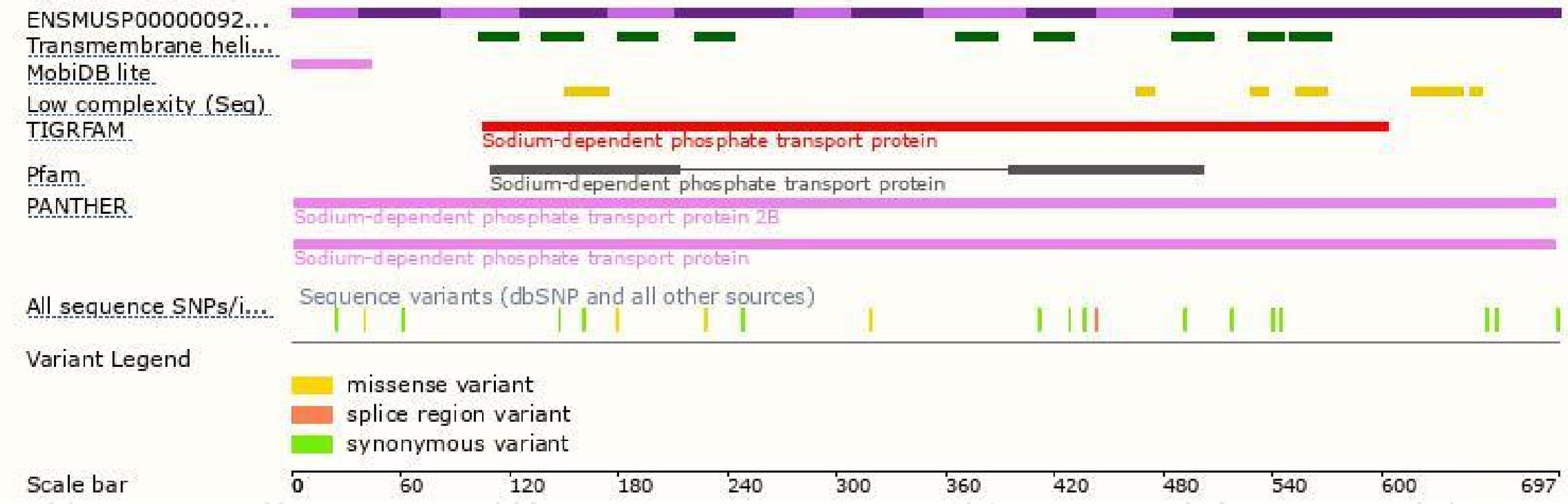
The strategy is based on the design of *Slc34a2-201* transcript, The transcription is shown below



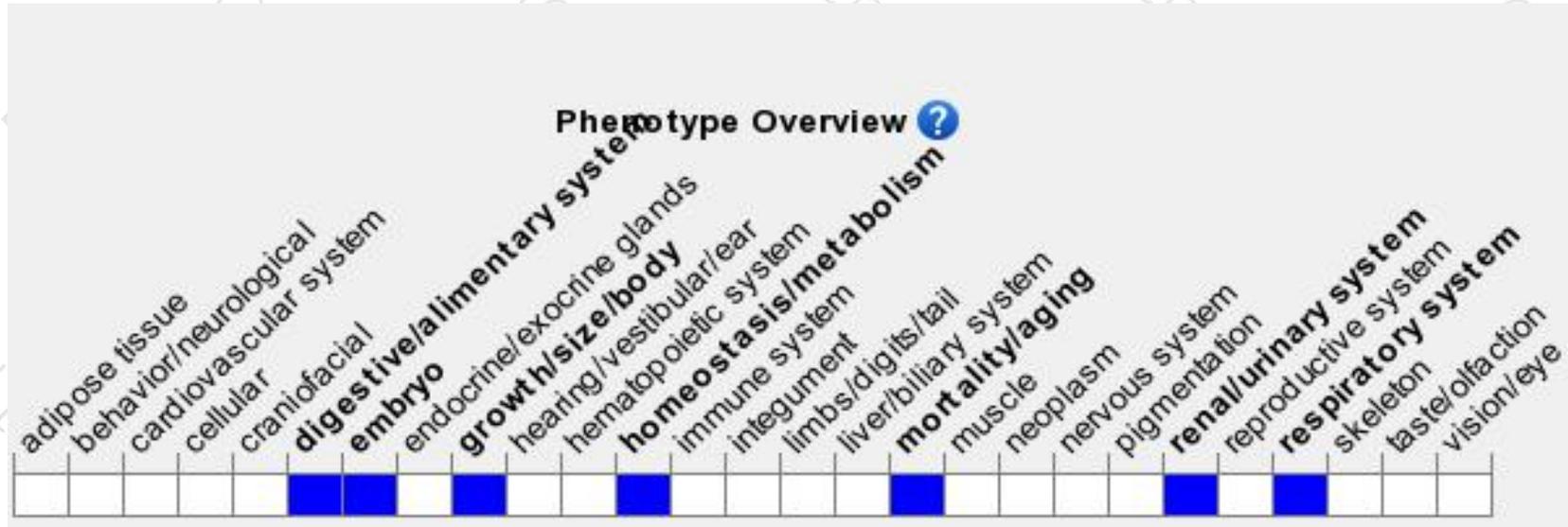
# Genomic location distribution



# 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, Homozygous null mice display embryonic lethality, embryonic growth arrest, failure of embryo turning and somitogenesis, impaired placental development and impaired yolk sac vascular remodeling.

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

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