Sox11 Cas9-KO Strategy

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Project Overview

Project Name	*Sox11*
Project type	**Cas9-KO**
Strain background	**C57BL/6JGpt**

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Sox11* gene. The schematic diagram is as follows:

Technical routes

➢ The *Sox11* gene has 1 transcript. According to the structure of *Sox11* gene, exon1 of *Sox11-201* (ENSMUST00000079063.6) 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 *Sox11* gene. The brief process is as follows: CRISPR/Cas9 system

➤ According to the existing MGI data,Homozygous null mice display neonatal lethality with impaired ossification and impaired development of the heart, lung, spleen, stomach, skeleton and pancreas. Mice homozygous for a different knock-out allele exhibit abnormal nervous system development and complete neonatal lethality.

➤ The *Sox11* gene is located on the Chr12. 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.

Sox11 SRY (sex determining region Y)-box 11 [Mus musculus (house mouse)]

Gene ID: 20666, updated on 12-Mar-2019

▲Summary

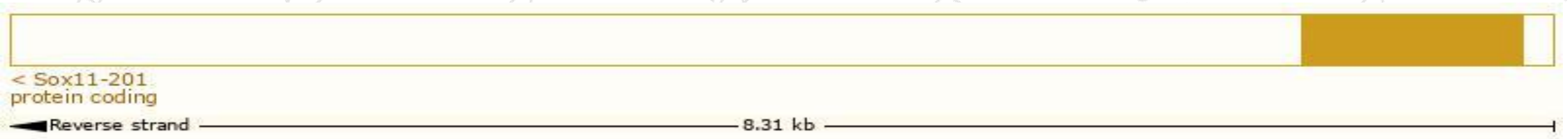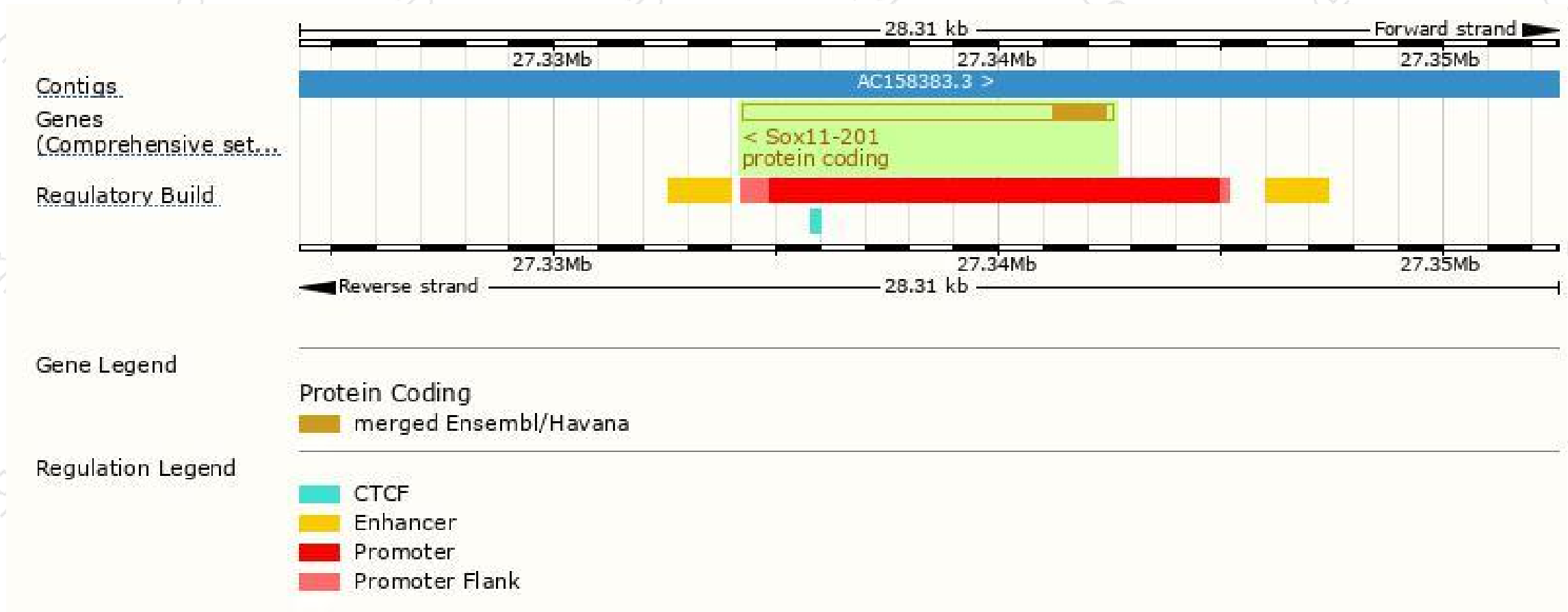|	
Official Symbol	Sox11 provided by MGI
Official Full Name	SRY (sex determining region Y)-box 11 provided by MGI
Primary source	MGI:MGI:98359
See related	Ensembl:ENSMUSG00000063632
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	1110038H03Rik, 6230403H02Rik, AI836553, end1
Orthologs	human all

Transcript information（Ensembl）

The gene has 1 transcript,and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Sox11-201	ENSMUST00000079063.6	8311	395aa	Protein coding	CCDS25849	Q7M6Y2	TSL:NA GENCODE basic APPRIS P1

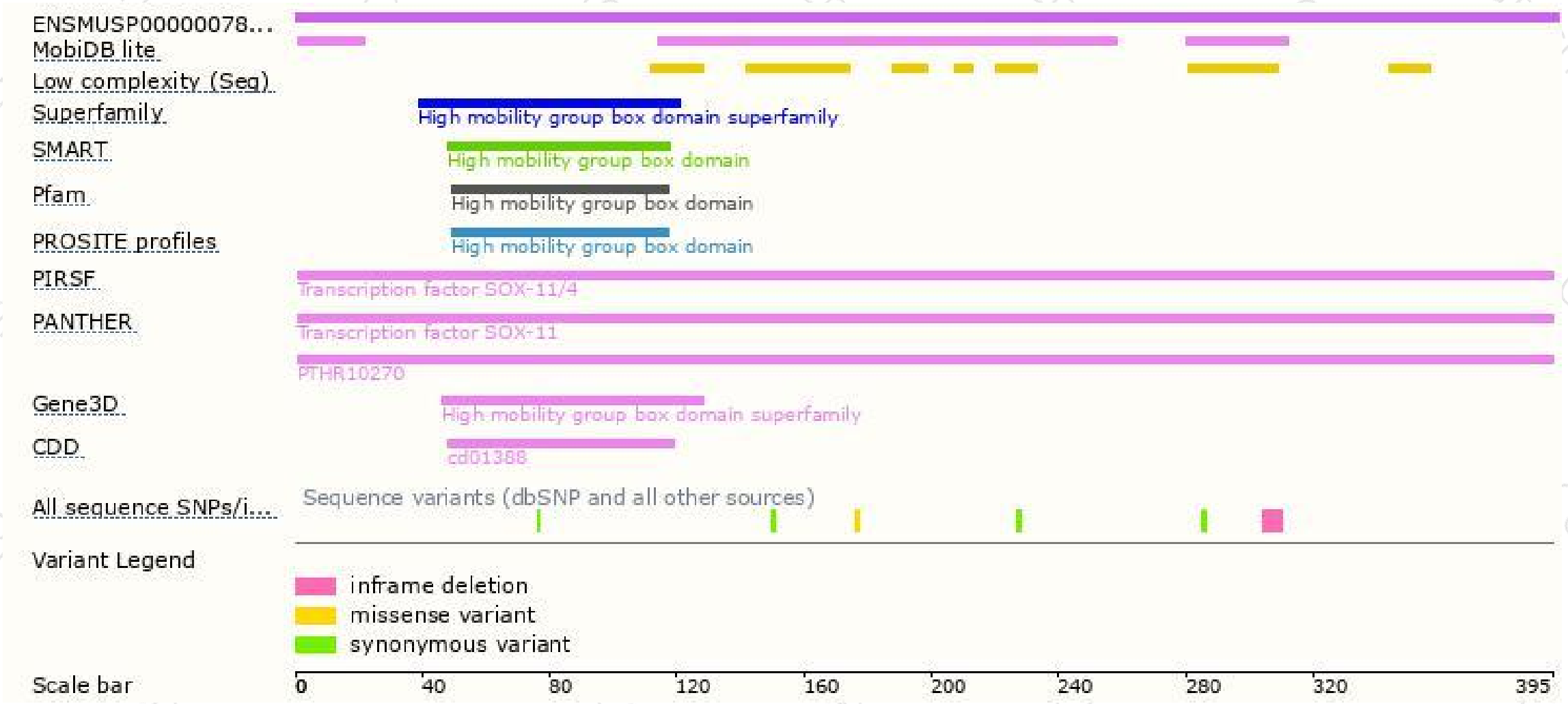The strategy is based on the design of *Sox11-201* transcript,The transcription is shown below

< Sox11-201
protein coding

◄ Reverse strand ——————————————— 8.31 kb ——————————————

Genomic location distribution

Protein domain

ENSMUSP00000078...	
MobiDB lite	
Low complexity (Seg)	
Superfamily	High mobility group box domain superfamily
SMART	High mobility group box domain
Pfam	High mobility group box domain
PROSITE profiles	High mobility group box domain
PIRSF	Transcription factor SOX-11/4
PANTHER	Transcription factor SOX-11
	PTHR10270
Gene3D	High mobility group box domain superfamily
CDD	cd01388
All sequence SNPs/i...	Sequence variants (dbSNP and all other sources)
Variant Legend	

Variant Legend
- ■ inframe deletion
- ■ missense variant
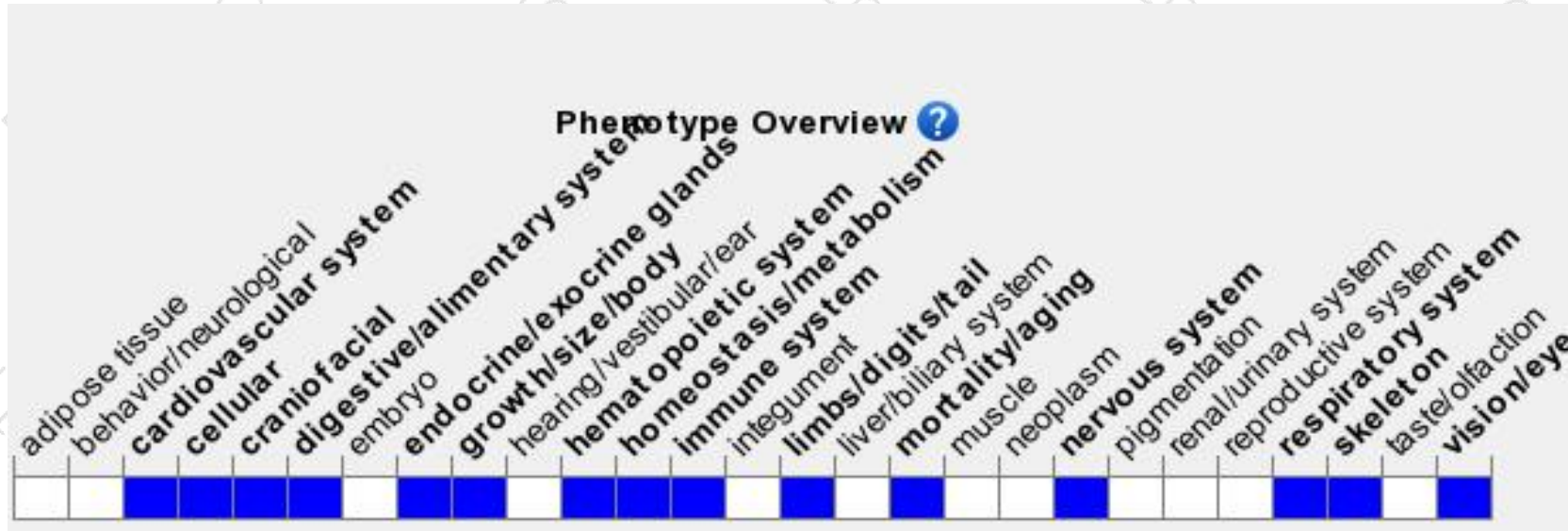- ■ synonymous variant

Scale bar

0   40   80   120   160   200   240   280   320   395

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 neonatal lethality with impaired ossification and impaired development of the heart, lung, spleen, stomach, skeleton and pancreas. Mice homozygous for a different knock-out allele exhibit abnormal nervous system development and complete neonatal lethality.

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