

Barhl2 Cas9-CKO Strategy

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Project Overview

Project Name

Barhl2

Project type

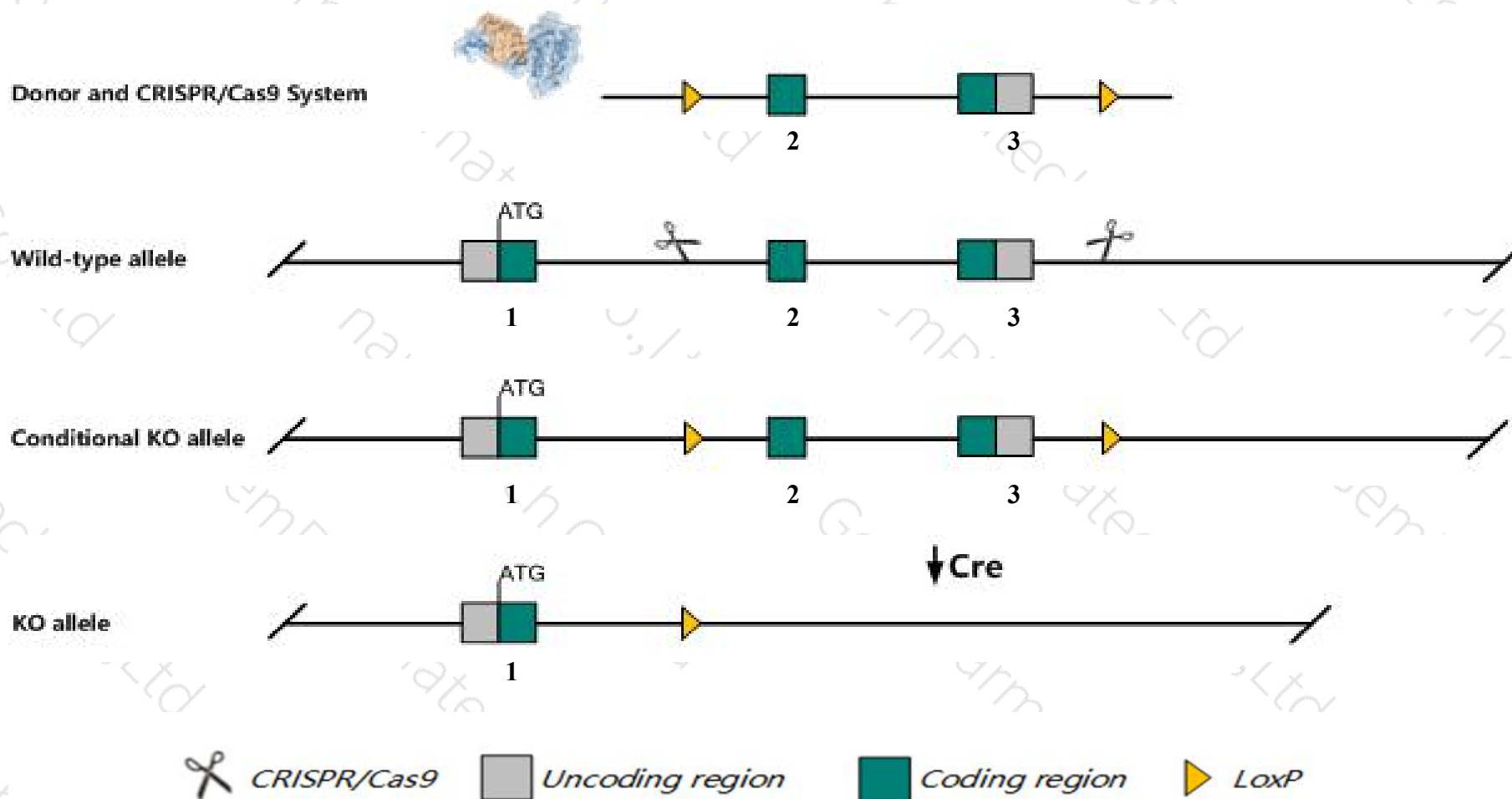
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Barhl2* gene. The schematic diagram is as follows:



- The *Barhl2* gene has 1 transcript. According to the structure of *Barhl2* gene, exon2-exon3 of *Barhl2-201* (ENSMUST00000086795.7) transcript is recommended as the knockout region. The region contains the stop codon. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Barhl2* gene. The brief process is as follows: gRNA was transcribed in vitro, donor was constructed. Cas9, gRNA and Donor were microinjected into the fertilized eggs of C57BL/6JGpt 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/6JGpt mice.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

- According to the existing MGI data, mice homozygous for a null allele display postnatal lethality with slow weight gain, impaired coordination, decreased numbers of retinal ganglion cells and retinal amacrine cells, and abnormal eye electrophysiology.
- The *Barhl2* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)

Barhl2 BarH like homeobox 2 [Mus musculus (house mouse)]

Gene ID: 104382, updated on 13-Mar-2020

Summary



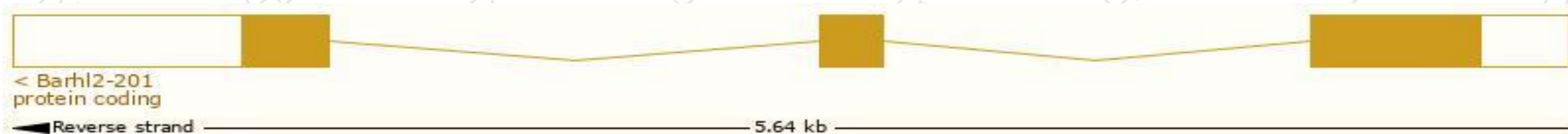
Official Symbol	Barhl2 provided by MGI
Official Full Name	BarH like homeobox 2 provided by MGI
Primary source	MGI:MGI:1859314
See related	Ensembl:ENSMUSG00000034384
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	E130309B19Rik, MBH1
Expression	Biased expression in cerebellum adult (RPKM 10.0), CNS E11.5 (RPKM 8.8) and 4 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

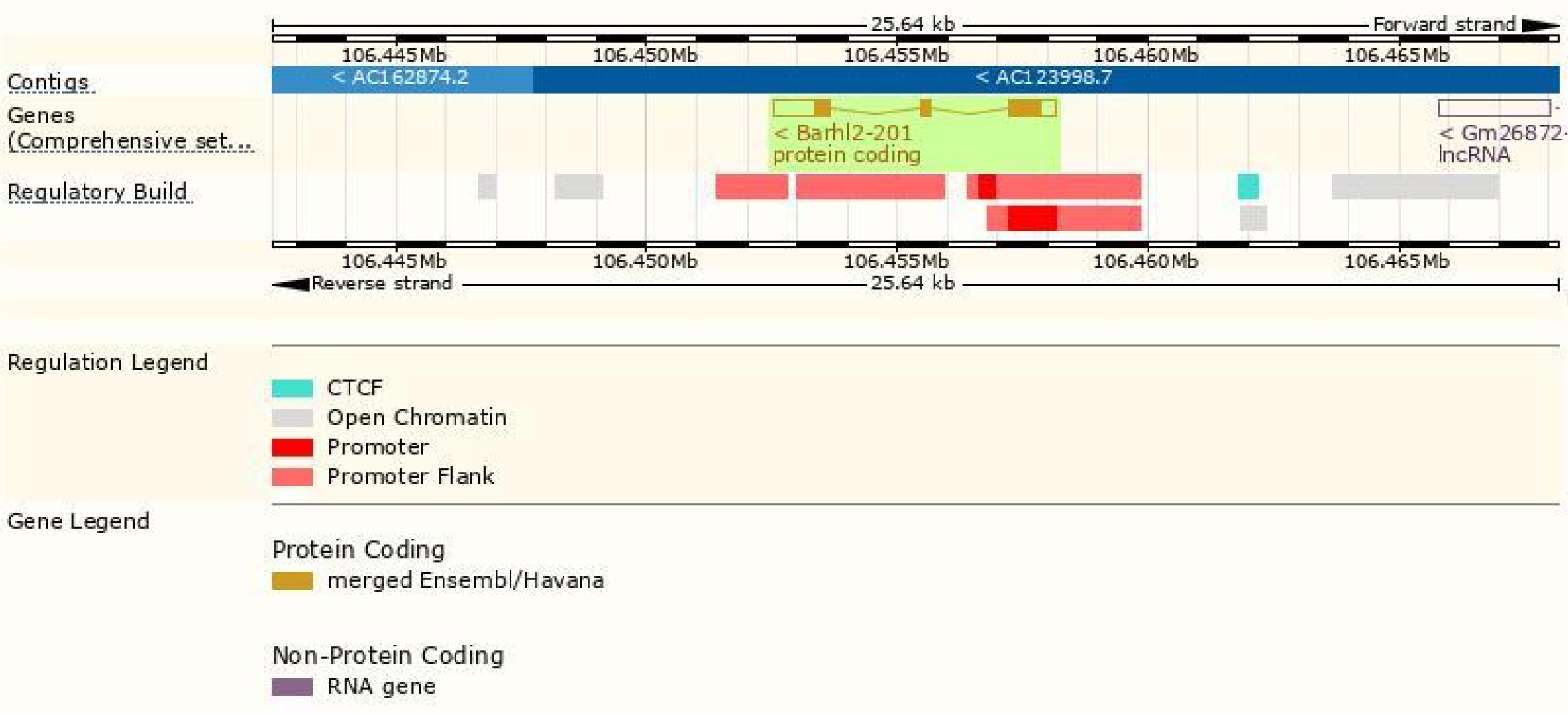
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Barhl2-201	ENSMUST00000086795.7	2314	384aa	Protein coding	CCDS19497	Q8VIB5	TSL:1 Gencode basic APPRIS P1

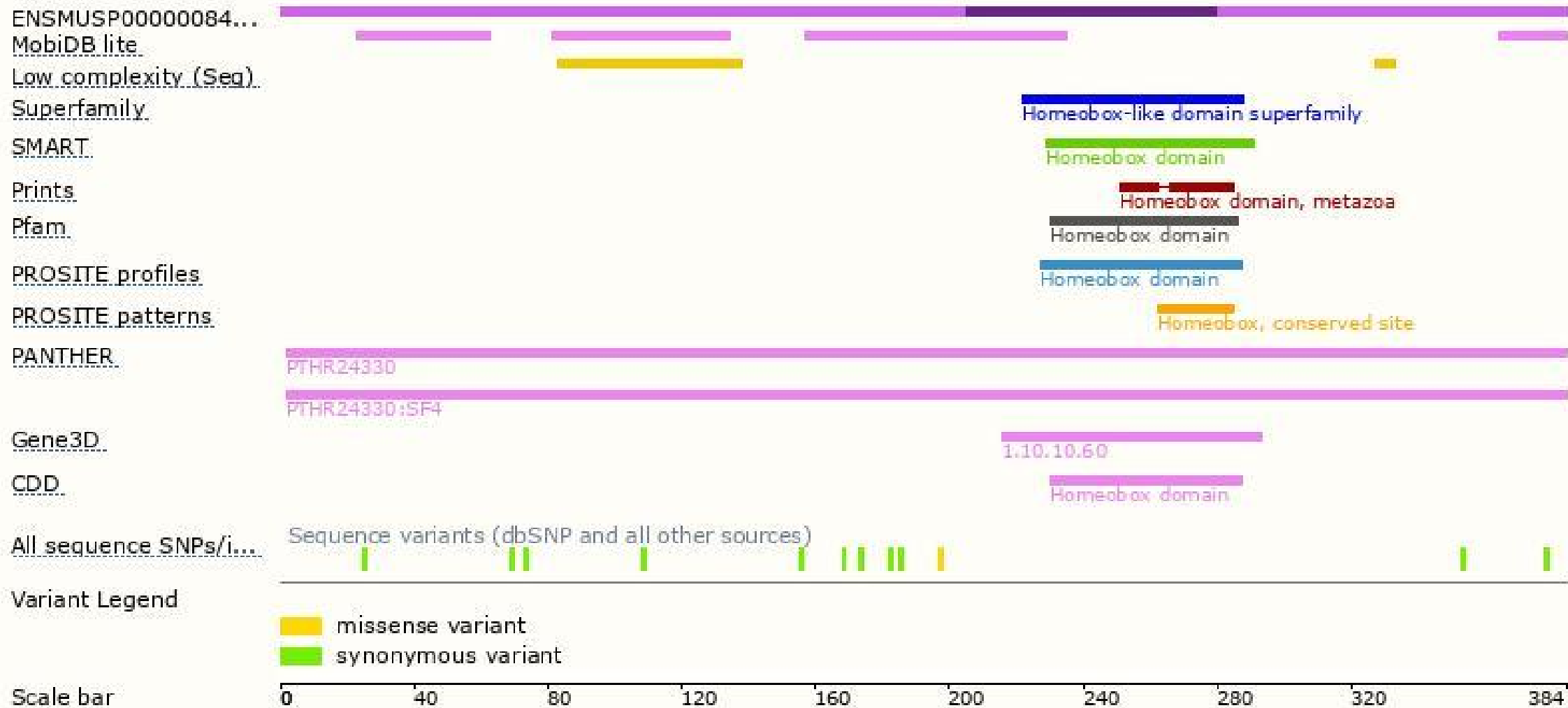
The strategy is based on the design of *Barhl2-201* transcript, the transcription is shown below:



Genomic location distribution

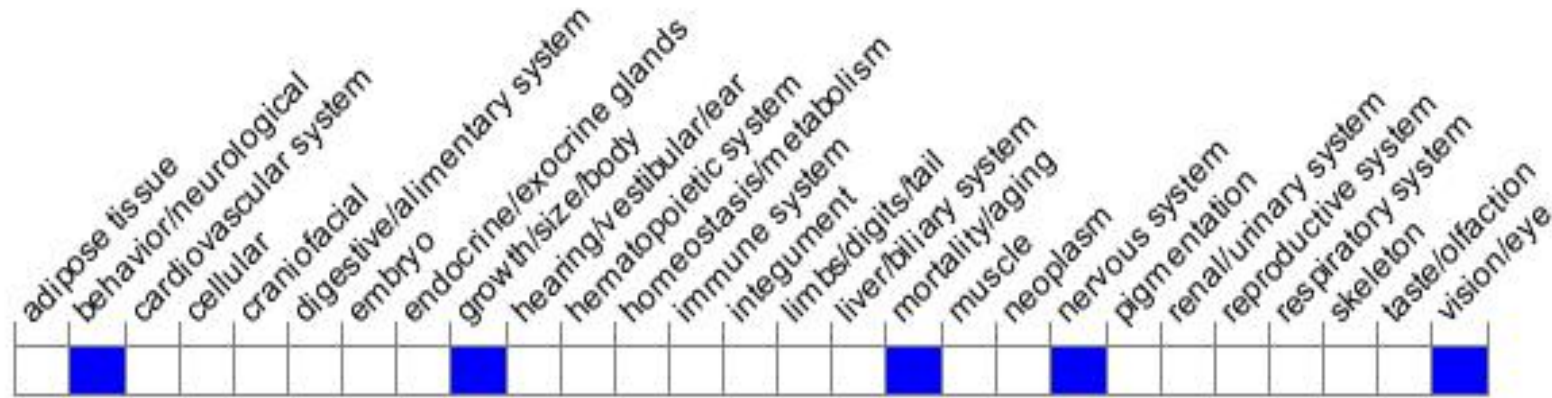


Protein domain



Mouse phenotype description(MGI)

Phenotype Overview



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 display postnatal lethality with slow weight gain, impaired coordination, decreased numbers of retinal ganglion cells and retinal amacrine cells, and abnormal eye electrophysiology.

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

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