

Hydin Cas9-CKO Strategy

Designer:

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Project Overview



Project Name

Hydin

Project type

Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

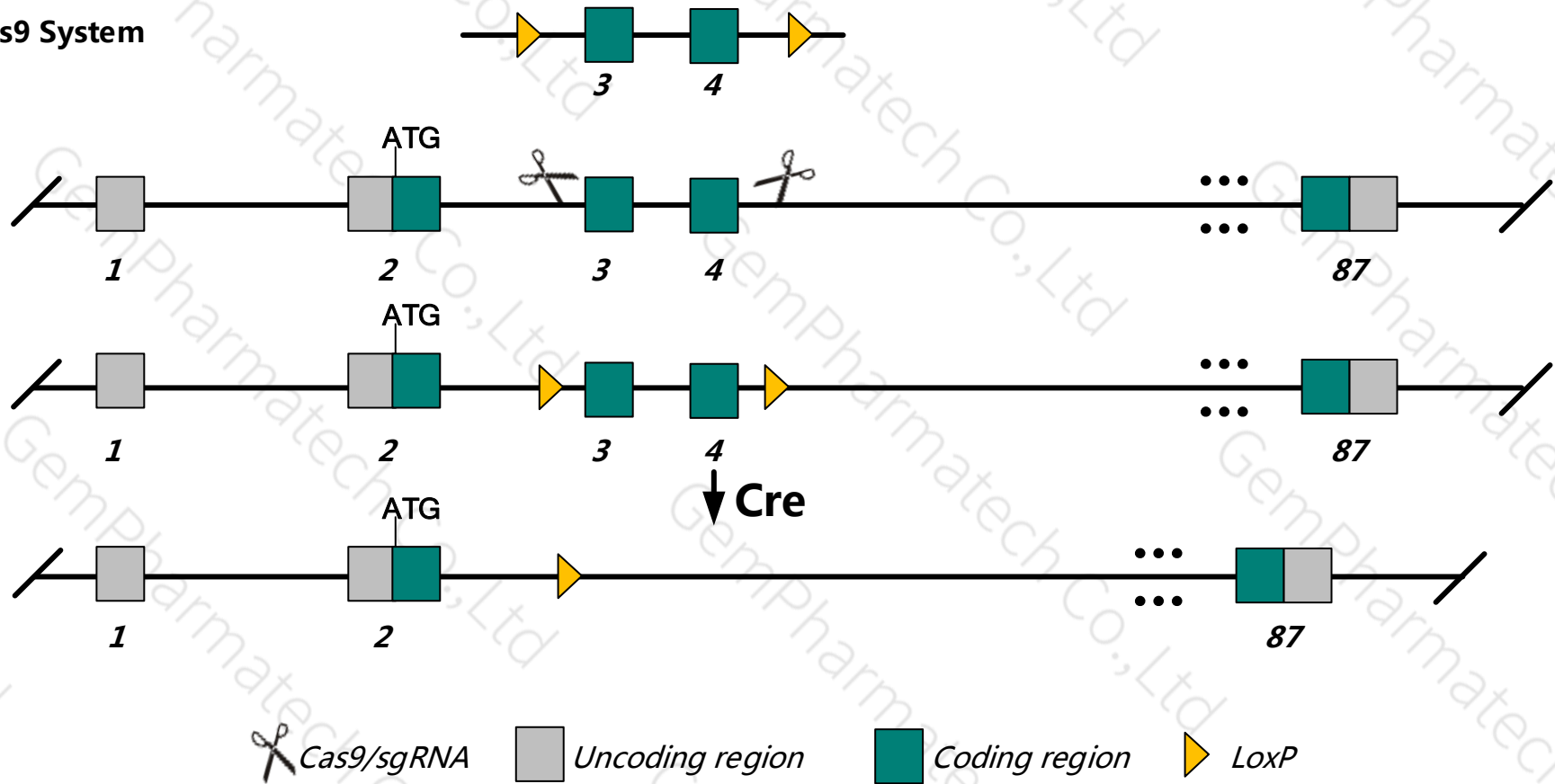
This model will use CRISPR/Cas9 technology to edit the *Hydin* gene. The schematic diagram is as follows:

Donor and CRISPR/Cas9 System

Wild-type allele

Conditional KO allele

KO allele



- The *Hydin* gene has 3 transcripts. According to the structure of *Hydin* gene, exon3-exon4 of *Hydin*-201 (ENSMUST00000043141.6) transcript is recommended as the knockout region. The region contains 284bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Hydin* gene. The brief process is as follows: sgRNA was transcribed in vitro, donor vector was constructed. Cas9, sgRNA 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 was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues or cell types.

- According to the existing MGI data , Mice homozygous for a mutation in this gene develop hydrocephaly after birth. Symptoms develop after 3-5 days. Affected animals usually die before 2 months of age.
- The *Hydin* gene is located on the Chr8. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)

Hydin HYDIN, axonemal central pair apparatus protein [*Mus musculus* (house mouse)]

Gene ID: 244653, updated on 2-Apr-2019

Summary

Official Symbol Hydin provided by [MGI](#)

Official Full Name HYDIN, axonemal central pair apparatus protein provided by [MGI](#)

Primary source [MGI:MGI:2389007](#)

See related [Ensembl:ENSMUSG00000059854](#)

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 hy3; hy-3; hyrh; 672398; Gm9558; 4932703P14; A830061H17; 1700034M11Rik; 4930545D19Rik; AC069308.21gm4

Expression Biased expression in testis adult (RPKM 10.5), lung adult (RPKM 1.1) and 1 other tissue [See more](#)

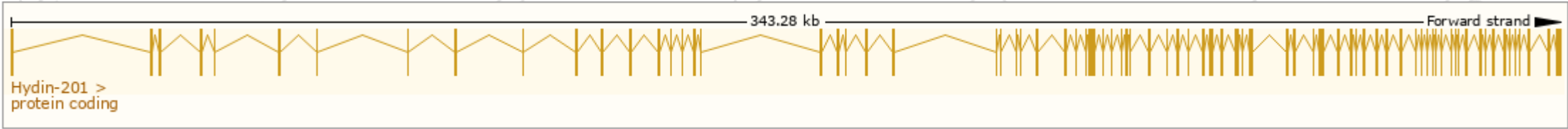
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

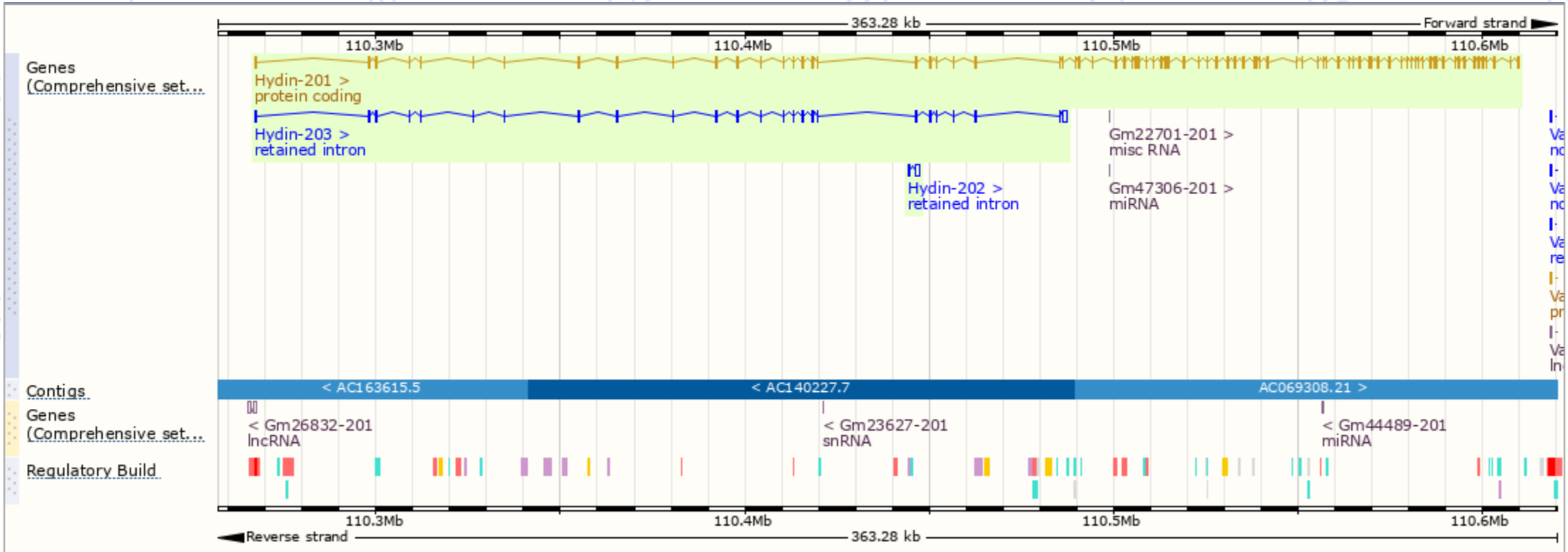
The gene has 3 transcripts, and all transcripts are shown below:

Show/hide columns (1 hidden)							Filter	
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Hydin-201	ENSMUST00000043141.6	15783	5154aa	Protein coding	CCDS40476	Q80W93	TSL:1	GENCODE basic APPRIS P1
Hydin-203	ENSMUST00000212218.1	5330	No protein	Retained intron	-	-	TSL:1	
Hydin-202	ENSMUST00000212034.1	1267	No protein	Retained intron	-	-	TSL:1	

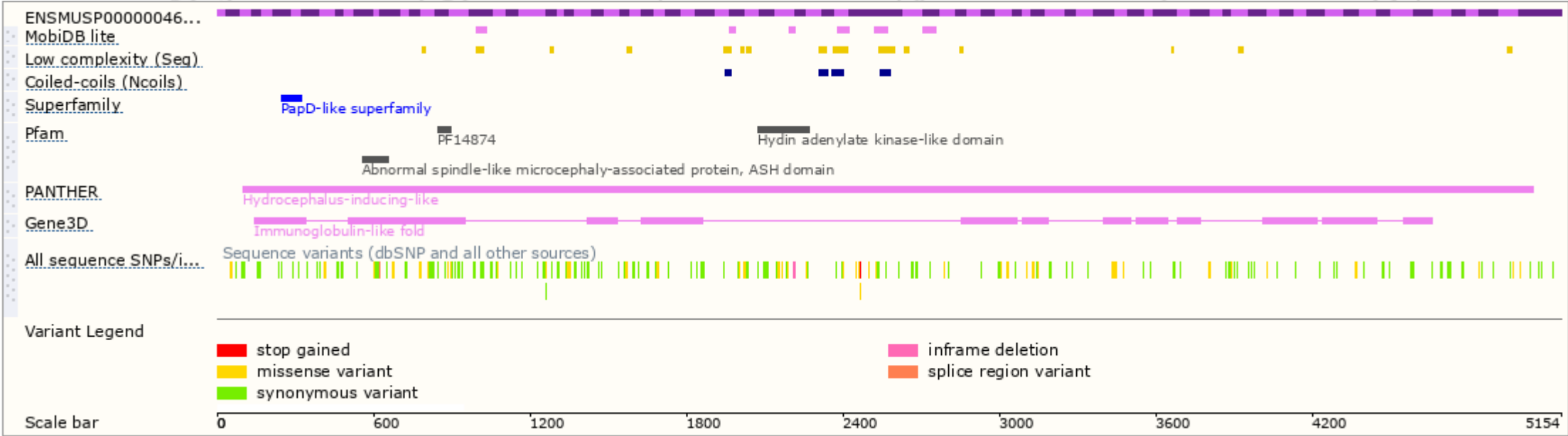
The strategy is based on the design of *Hydin*-201 transcript, The transcription is shown below



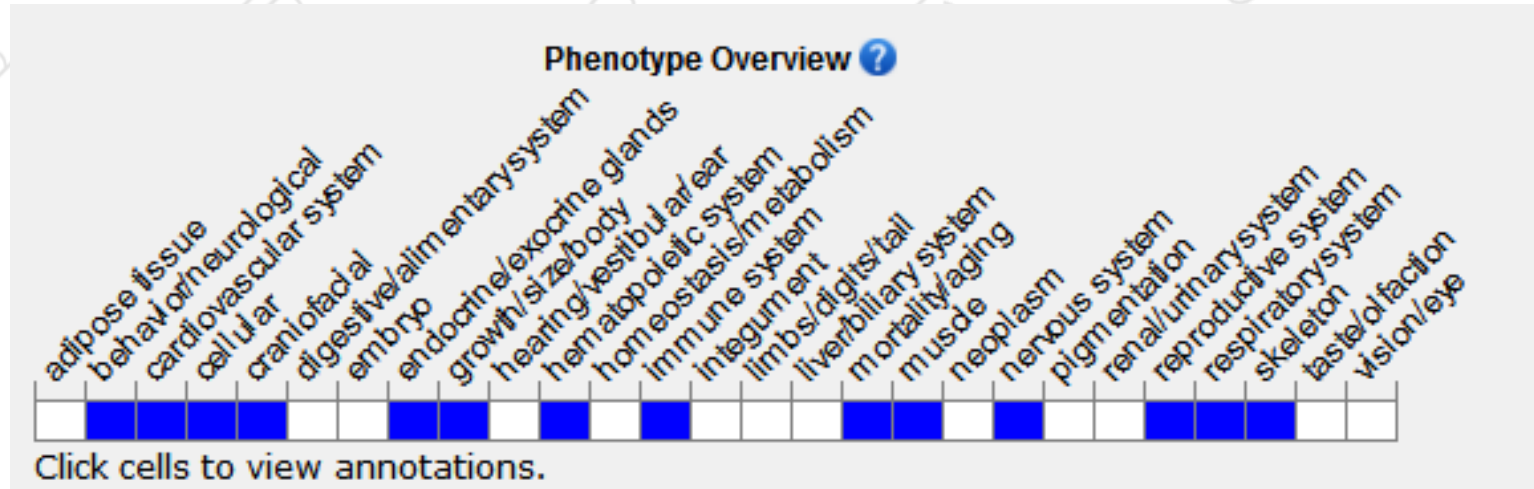
Genomic location distribution



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 mutation in this gene develop hydrocephaly after birth.

Symptoms develop after 3-5 days. Affected animals usually die before 2 months of age.

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