

# **Dnah5** Cas9-CKO Strategy

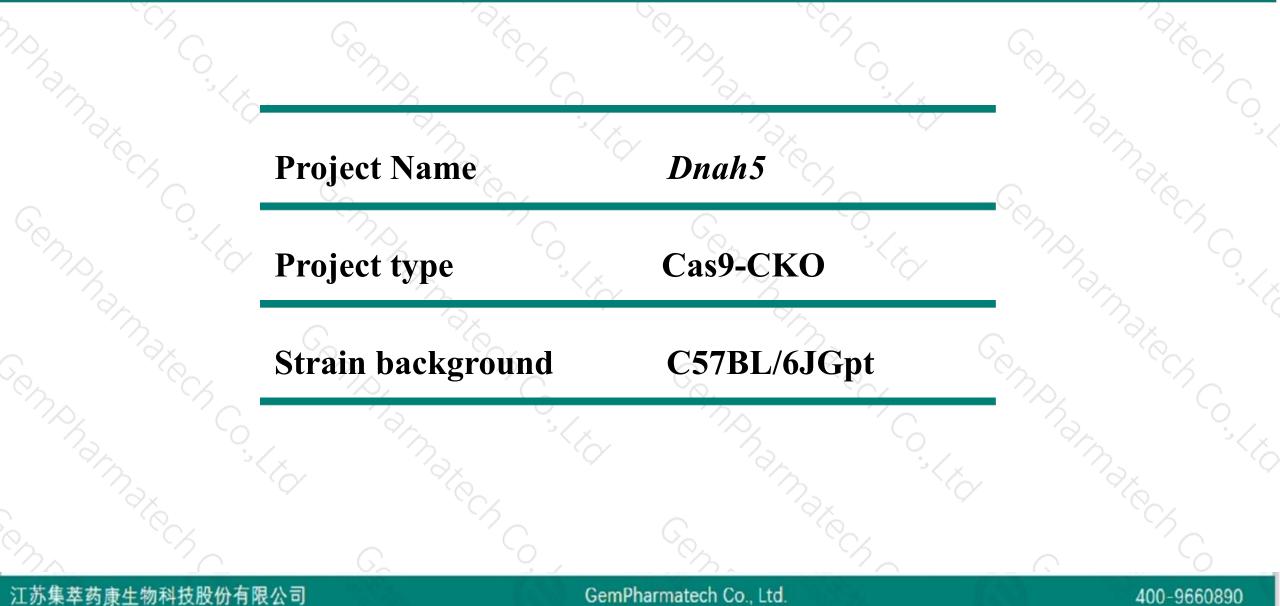
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**Reviewer: JiaYu** 

Design Date: 2020-6-9

# **Project Overview**

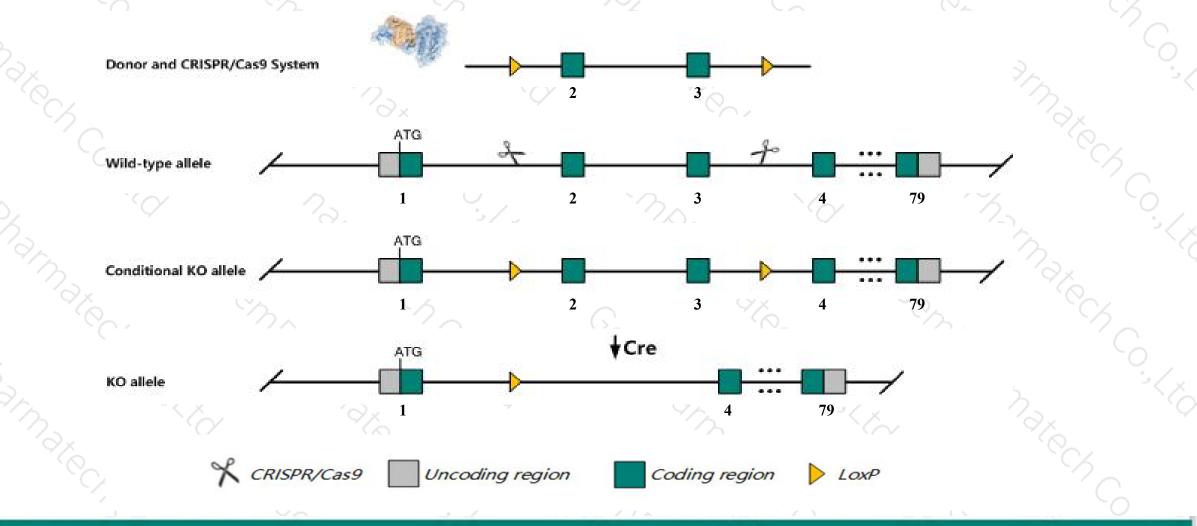




# **Conditional Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Dnah5* gene. The schematic diagram is as follows:



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The Dnah5 gene has 1 transcript. According to the structure of Dnah5 gene, exon2-exon3 of Dnah5-201 (ENSMUST00000067048.7) transcript is recommended as the knockout region. The region contains 220bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Dnah5* gene. The brief process is as follows:CRISPR/Cas9 system 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 disruption in this gene display postnatal lethality, hydrocephalus, respiratory infections, situs inversus and ciliary immotility.
- The *Dnah5* gene is located on the Chr15. 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)**



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### Dnah5 dynein, axonemal, heavy chain 5 [Mus musculus (house mouse)]

Gene ID: 110082, updated on 13-Mar-2020

### Summary

| Official Symbol           | Dnah5 provided by MGI   |
|---------------------------|---|
| <b>Official Full Name</b> | dynein, axonemal, heavy chain 5 provided by <u>MGI</u>  |
| <b>Primary source</b>     | MGI:MGI:107718  |
| See related               | Ensembl:ENSMUSG0000022262   |
| 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             | AU022615, Dnahc5, Mdnah5, b2b1134Clo, b2b1154Clo, b2b1537Clo, b2b1565Clo, b2b3491Clo, b2b601Clo, mKIAA1603  |
| Expression                | Low expression observed in reference datasetSee more  |
| Orthologs                 | human all   |

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

| Name      | Transcript ID       | bp    | Protein       | Biotype        | CCDS      | UniProt | Flags                         |
|-----------|---------------------|-------|---------------|----------------|-----------|---------|-------------------------------|
| Dnah5-201 | ENSMUST0000067048.7 | 15637 | <u>4621aa</u> | Protein coding | CCDS27404 | Q8VHE6  | TSL:5 GENCODE basic APPRIS P1 |

The strategy is based on the design of *Dnah5-201* transcript, the transcription is shown below:

Dnah5-201 > protein coding

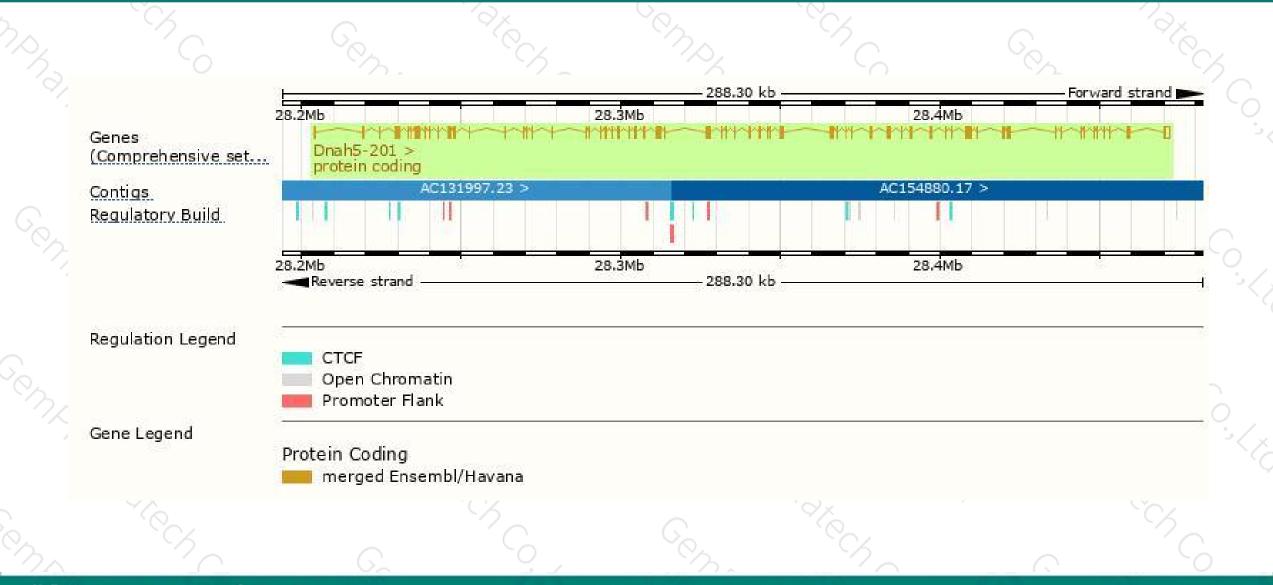
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268.30 kb

Forward strand

## **Genomic location distribution**



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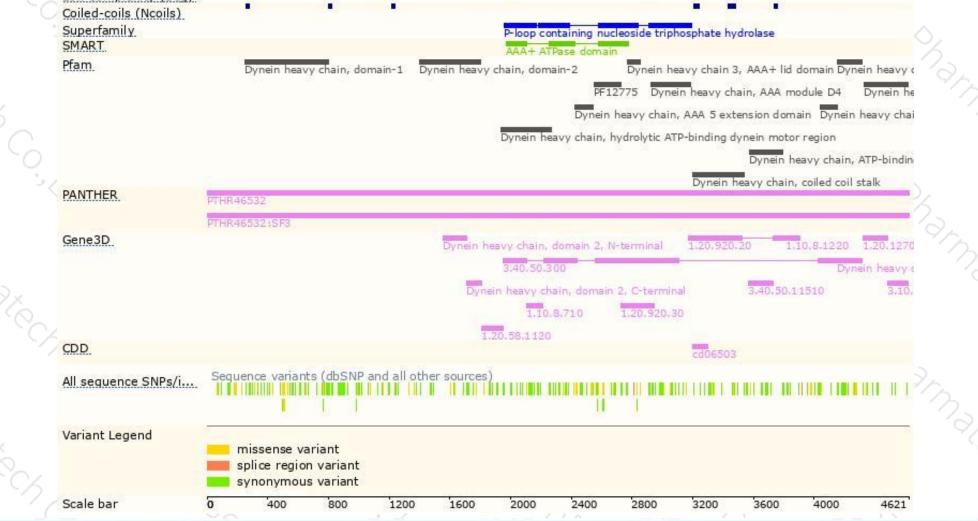
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#### ENSMUSP00000069... MobiDB lite Low complexity (Seg) Coiled-coils (Neoils)

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## **Protein domain**

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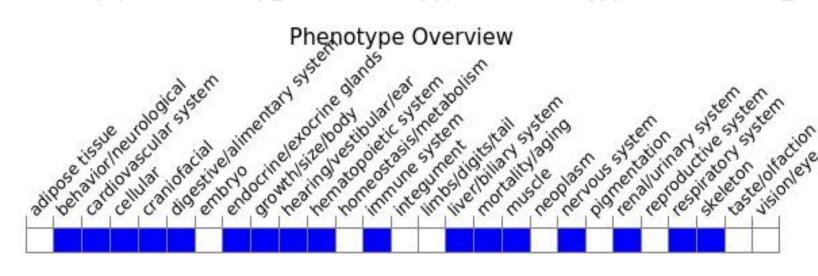


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# Mouse phenotype description(MGI)





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 disruption in this gene display postnatal lethality, hydrocephalus, respiratory infections, situs inversus and ciliary immotility.



If you have any questions, you are welcome to inquire. Tel: 400-9660890



