

Nbeal2 Cas9-KO Strategy

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Design Date:

2020-2-20

Project Overview

Project Name

Nbeal2

Project type

Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Nbeal2* gene. The schematic diagram is as follows:



- The *Nbeal2* gene has 11 transcripts. According to the structure of *Nbeal2* gene, exon2-exon6 of *Nbeal2-210* (ENSMUST00000167320.7) transcript is recommended as the knockout region. The region contains 502bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Nbeal2* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygous null mice exhibit megakaryocyte and platelet abnormalities resulting in impaired arterial thrombus formation and protection from infarction following cerebral ischemia. Wound repair is impaired. These abnormalities result in a bleeding disorder similar to Gray Platelet Syndrome.
- The *Nbeal2* gene is located on the Chr9. 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)

Nbeal2 neurobeachin-like 2 [Mus musculus (house mouse)]

Gene ID: 235627, updated on 31-Jan-2019

Summary



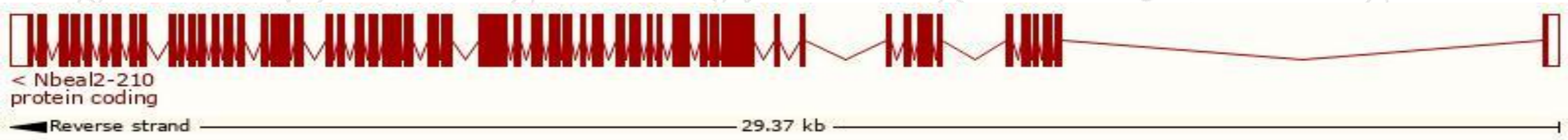
| | |
|---------------------------|---|
| Official Symbol | Nbeal2 provided by MGI |
| Official Full Name | neurobeachin-like 2 provided by MGI |
| Primary source | MGI:MGI:2448554 |
| See related | Ensembl:ENSMUSG00000056724 |
| Gene type | protein coding |
| RefSeq status | VALIDATED |
| Organism | Mus musculus |
| Lineage | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| Also known as | 1110014F23Rik, BC042396, mKIAA0540 |
| Expression | Broad expression in thymus adult (RPKM 58.1), colon adult (RPKM 27.1) and 15 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

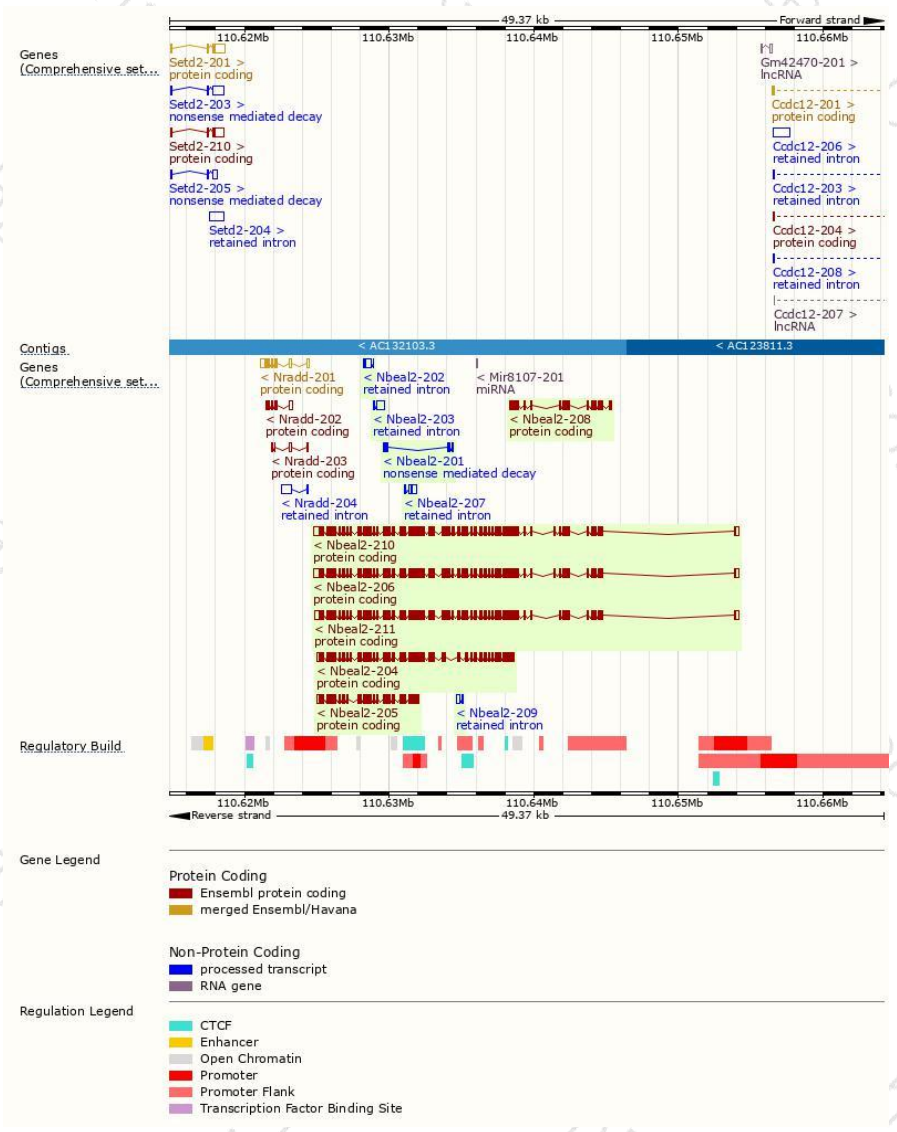
The gene has 11 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|------------|--------------------------------------|------|------------------------|-------------------------|---------------------------|----------------------------|---------------------------------|
| Nbeal2-210 | ENSMUST00000167320.7 | 8803 | 2750aa | Protein coding | CCDS52939 | E9Q9L6 | TSL:1 GENCODE basic APPRIS P2 |
| Nbeal2-206 | ENSMUST00000133191.7 | 8782 | 2743aa | Protein coding | - | Q6ZQA0 | TSL:5 GENCODE basic APPRIS ALT2 |
| Nbeal2-211 | ENSMUST00000196488.4 | 8599 | 2716aa | Protein coding | - | A0A0G2JFQ4 | TSL:5 GENCODE basic APPRIS ALT2 |
| Nbeal2-204 | ENSMUST00000130024.1 | 6180 | 2032aa | Protein coding | - | F6VTL9 | CDS 5' incomplete TSL:5 |
| Nbeal2-205 | ENSMUST00000131017.7 | 3303 | 1073aa | Protein coding | - | F6ZZB1 | CDS 5' incomplete TSL:5 |
| Nbeal2-208 | ENSMUST00000149089.1 | 1869 | 593aa | Protein coding | - | D3Z2K6 | CDS 3' incomplete TSL:5 |
| Nbeal2-201 | ENSMUST00000123996.1 | 457 | 96aa | Nonsense mediated decay | - | F7CGQ9 | CDS 5' incomplete TSL:5 |
| Nbeal2-203 | ENSMUST00000129095.1 | 664 | No protein | Retained intron | - | - | TSL:2 |
| Nbeal2-207 | ENSMUST00000138072.1 | 479 | No protein | Retained intron | - | - | TSL:2 |
| Nbeal2-202 | ENSMUST00000126088.3 | 432 | No protein | Retained intron | - | - | TSL:2 |
| Nbeal2-209 | ENSMUST00000153960.1 | 295 | No protein | Retained intron | - | - | TSL:5 |

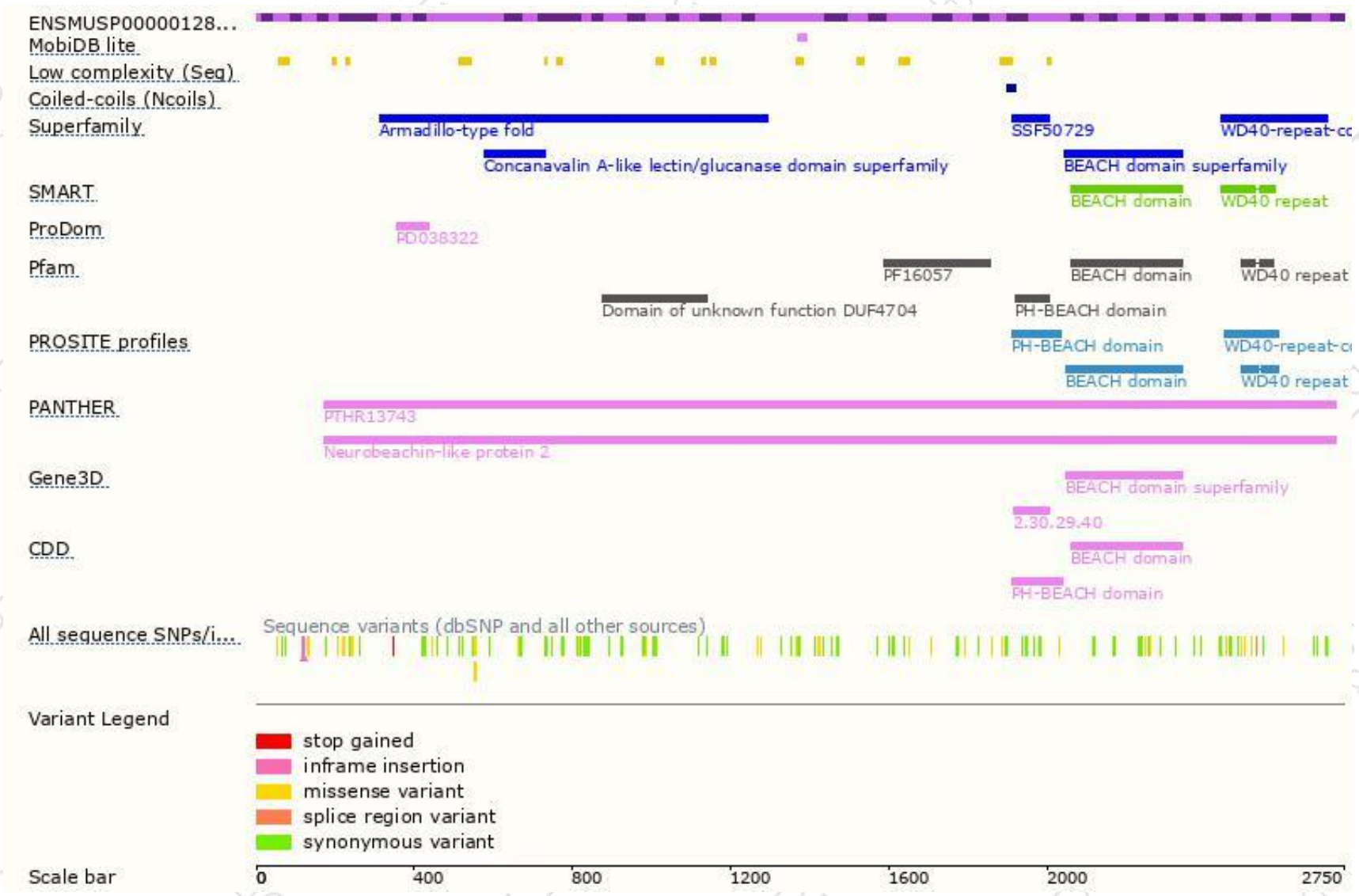
The strategy is based on the design of *Nbeal2-210* transcript,The transcription is shown below



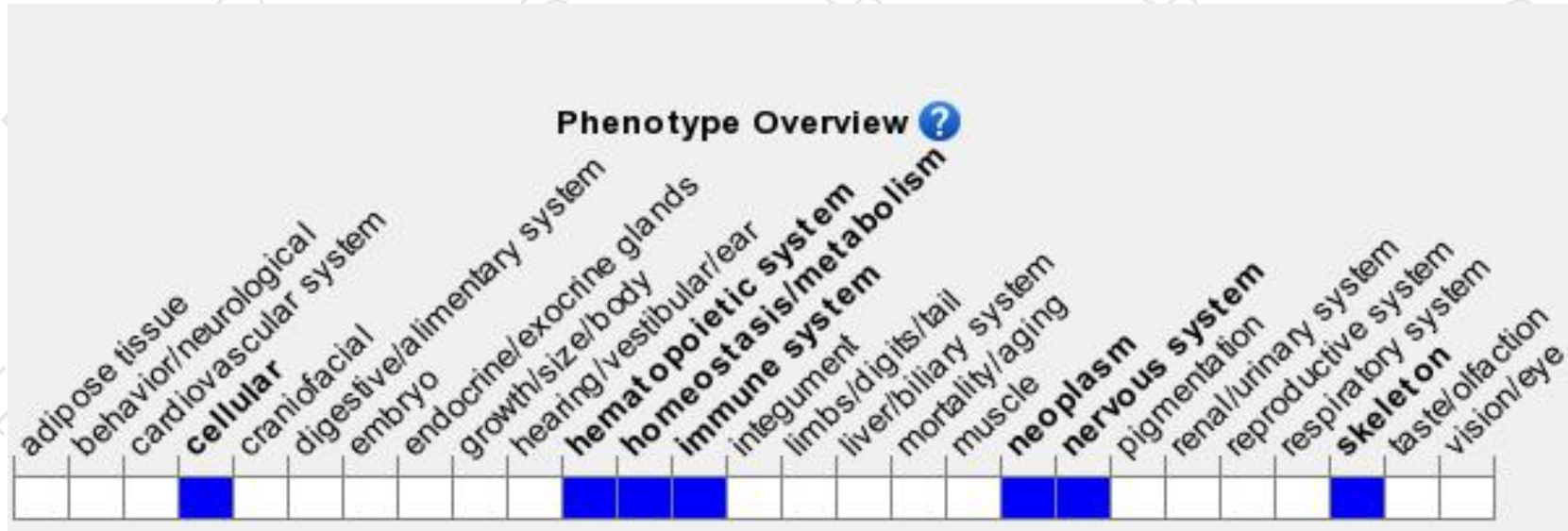
Genomic location distribution



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 exhibit megakaryocyte and platelet abnormalities resulting in impaired arterial thrombus formation and protection from infarction following cerebral ischemia. Wound repair is impaired. These abnormalities result in a bleeding disorder similar to Gray Platelet Syndrome.

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

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