

# Eef2 Cas9-KO Strategy

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# Overview

## Target Gene Name

- Eef2

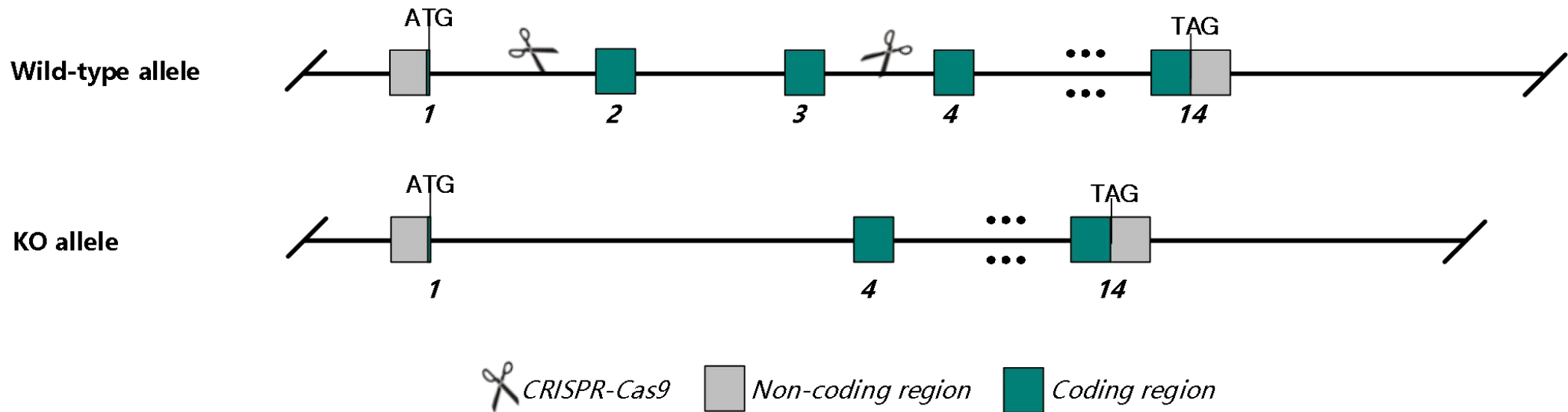
## Project Type

- Cas9-KO

## Genetic Background

- C57BL/6JGpt

# Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the *Eef2* gene.

# Technical Information

- The *Eef2* gene has 3 transcripts. According to the structure of *Eef2* gene, exon2-3 of *Eef2*-201 (ENSMUST00000047864.11) transcript is recommended as the knockout region. The region contains 397 bp of coding sequences. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Eef2* gene. The brief process is as follows: gRNAs were transcribed in vitro. Cas9 and gRNAs 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 on-target amplicon sequencing. A stable F1-generation mouse strain was obtained by mating positive F0-generation mice with C57BL/6JGpt mice and confirmation of the desired mutant allele was carried out by PCR and on-target amplicon sequencing.

# Gene Information

## Eef2 eukaryotic translation elongation factor 2 [ *Mus musculus* (house mouse) ]

Gene ID: 13629, updated on 12-Apr-2023

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### Summary

<b>Official Symbol</b>	Eef2 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	eukaryotic translation elongation factor 2 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:95288</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000034994</a> <a href="#">AllianceGenome:MGI:95288</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	Ef-2
<b>Summary</b>	Enables GTPase activity. Acts upstream of or within hematopoietic progenitor cell differentiation and translational elongation. Located in cytoplasm. Part of polysome. Is active in synapse. Is expressed in brain. Human ortholog(s) of this gene implicated in glaucoma and spinocerebellar ataxia type 26. Orthologous to human EEF2 (eukaryotic translation elongation factor 2). [provided by Alliance of Genome Resources, Apr 2022]
<b>Expression</b>	Ubiquitous expression in ovary adult (RPKM 1248.0), colon adult (RPKM 810.0) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>
<b>NEW</b>	Try the new <a href="#">Gene table</a> Try the new <a href="#">Transcript table</a>

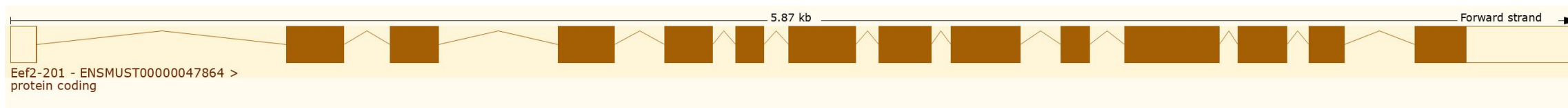
Source: <https://www.ncbi.nlm.nih.gov/>

# Transcript Information

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

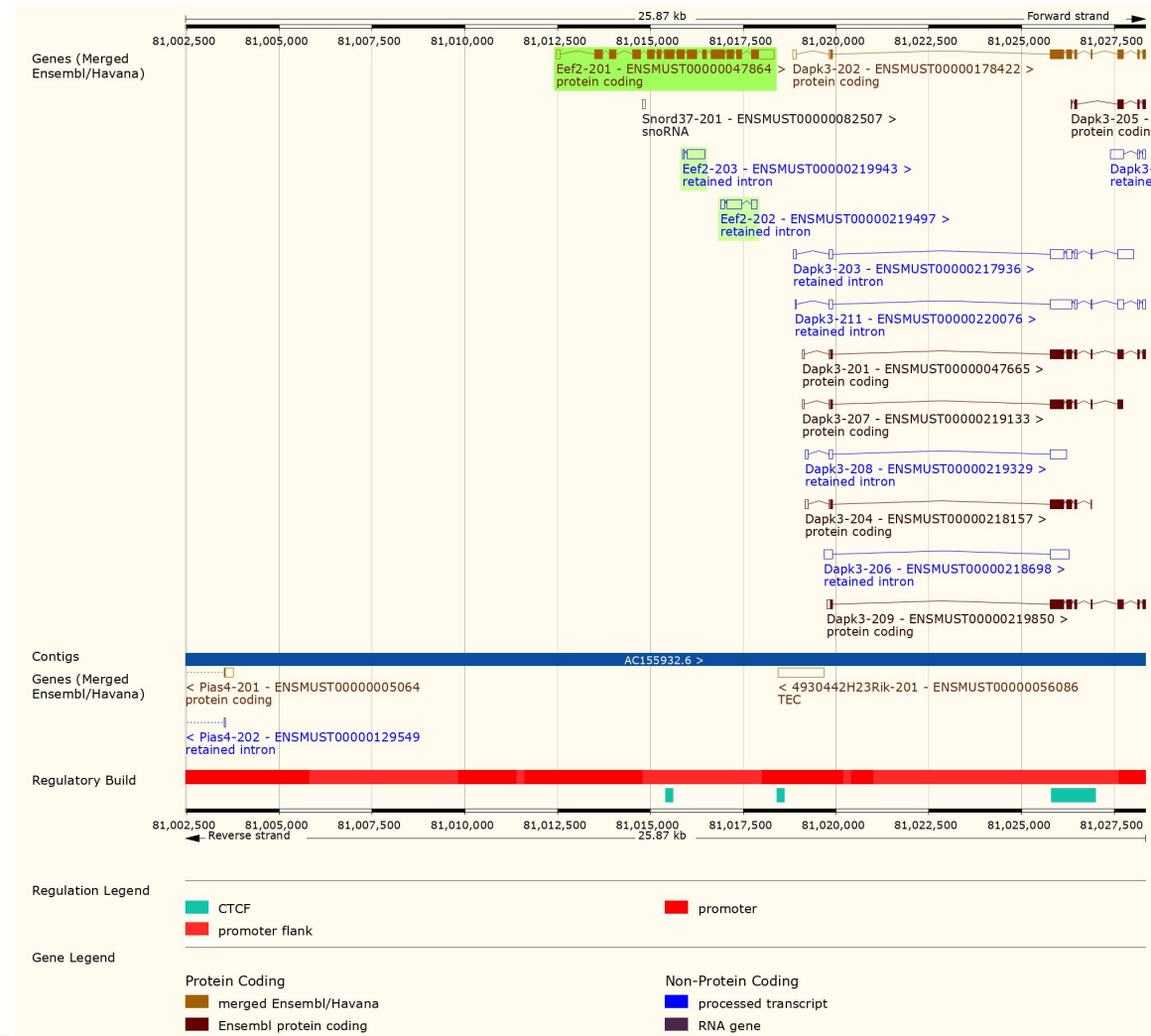
Show/hide columns (1 hidden)							Filter	
Transcript ID	Name	bp	Protein	Biotype	CCDS	UniProt Match	Flags	
<a href="#">ENSMUST00000047864.11</a>	Eef2-201	3089	<a href="#">858aa</a>	Protein coding	<a href="#">CCDS35993</a>	<a href="#">P58252</a>	Ensembl Canonical	GENCODE basic APPRIS P1 TSL:1
<a href="#">ENSMUST00000219497.2</a>	Eef2-202	651	No protein	Retained intron		-		TSL:3
<a href="#">ENSMUST00000219943.2</a>	Eef2-203	519	No protein	Retained intron		-		TSL:1

The strategy is based on the design of *Eef2*-201 transcript, the transcription is shown below:

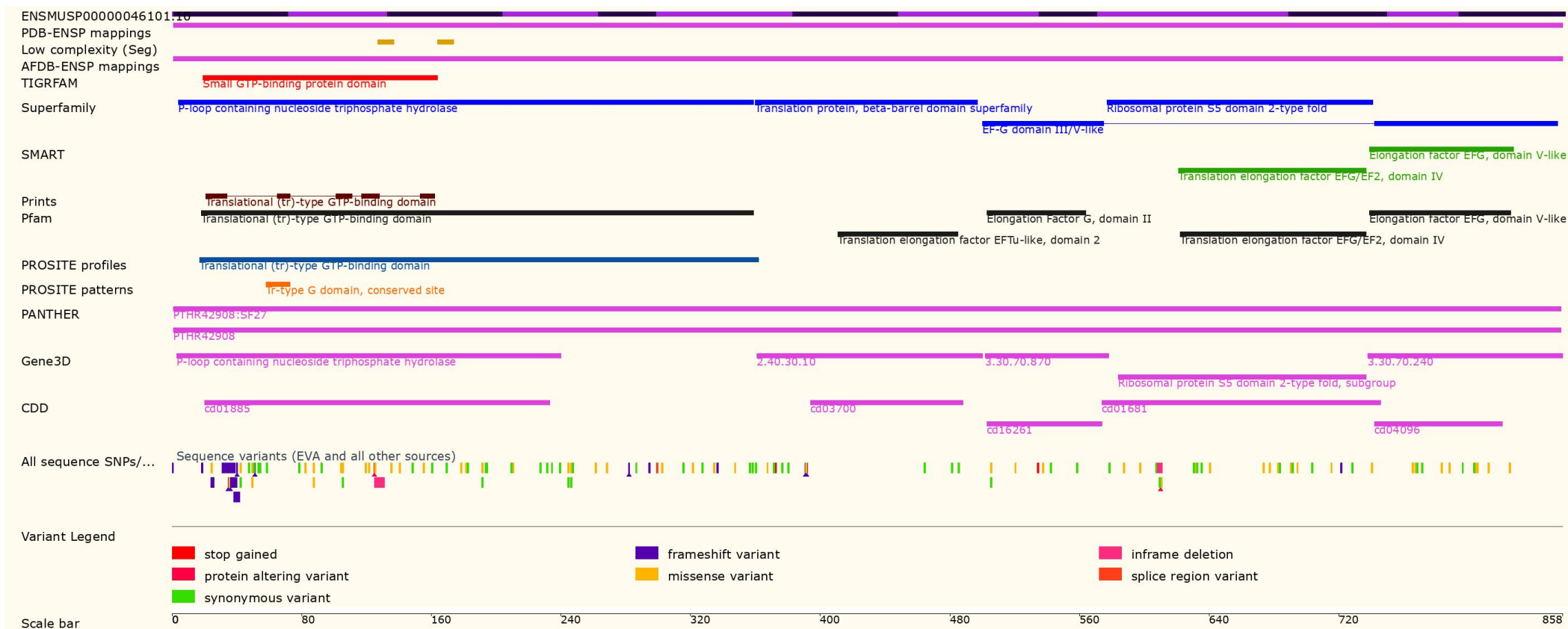


Source: <https://www.ensembl.org>

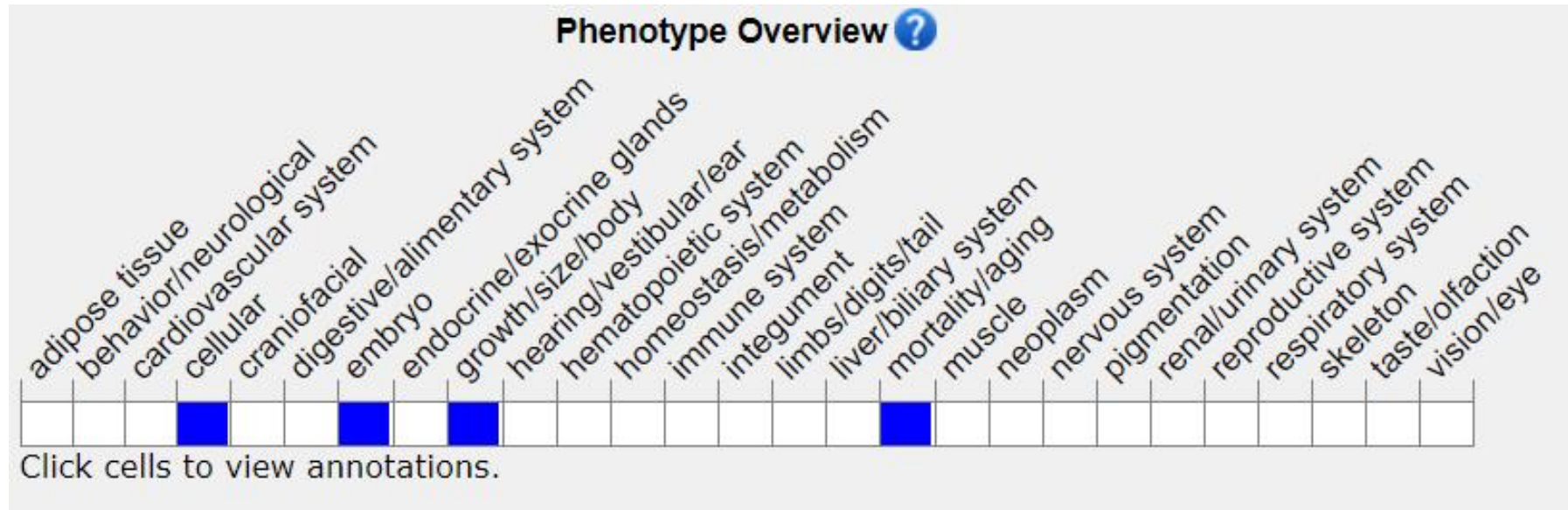
# Genomic Information



# Protein Information



# Mouse Phenotype Information (MGI)



- Mice homozygous for a mutation removing the diphthamide modification display partial neonatal lethality, fetal growth retardation and abnormal cell physiology.

# Important Information

- The effect of *Eef2*-202, *Eef2*-203 is unknown.
- This Strategy may effect the N-terminal regulation of *Snord37* and *Dapk3*.
- *Eef2* is located on Chr10. If the knockout mice are crossed with other mouse strains to obtain double homozygous mutant offspring, please avoid the situation that the second gene is on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risks of the mutation on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.