

Chp1 Cas9-KO Strategy

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Overview

Target Gene Name

- Chp1

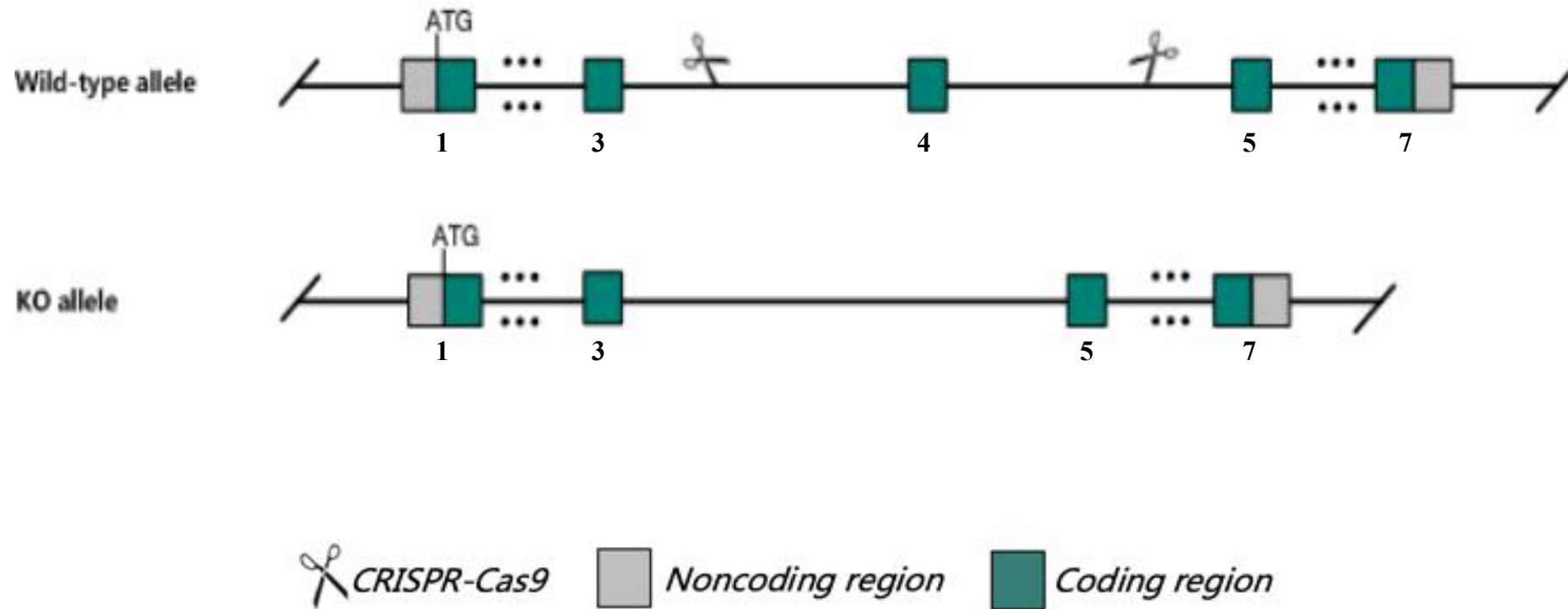
Project Type

- Cas9-KO

Genetic Background

- C57BL/6JGpt

Strain Strategy



Technical Information

- The *Chp1* gene has 5 transcripts. According to the structure of *Chp1* gene, exon4 of *Chp1*-201 (ENSMUST00000014221.13) transcript is recommended as the knockout region. The region contains 128bp coding sequence. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Chp1* 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

Chp1 calcineurin-like EF hand protein 1 [*Mus musculus* (house mouse)]

[Download Datasets](#)

Gene ID: 56398, updated on 5-Mar-2024

Summary

Official Symbol	Chp1 provided by MGI
Official Full Name	calcineurin-like EF hand protein 1 provided by MGI
Primary source	MGI:MGI:1927185
See related	Ensembl:ENSMUSG00000014077 AllianceGenome:MGI:1927185
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	Chp; p24; vac; Cahp; Sid470p; 1500003O03Rik
Summary	Predicted to enable several functions, including calcium ion binding activity; calcium-dependent protein binding activity; and microtubule binding activity. Predicted to be involved in several processes, including microtubule cytoskeleton organization; regulation of protein modification process; and regulation of protein transport. Predicted to act upstream of or within protein transport. Predicted to be located in several cellular components, including Golgi membrane; microtubule cytoskeleton; and transport vesicle. Is expressed in genitourinary system. Human ortholog(s) of this gene implicated in spastic ataxia. Orthologous to human CHP1 (calcineurin like EF-hand protein 1). [provided by Alliance of Genome Resources, Apr 2022]
Expression	Ubiquitous expression in subcutaneous fat pad adult (RPKM 137.3), mammary gland adult (RPKM 107.2) and 28 other tissues See more
Orthologs	human all

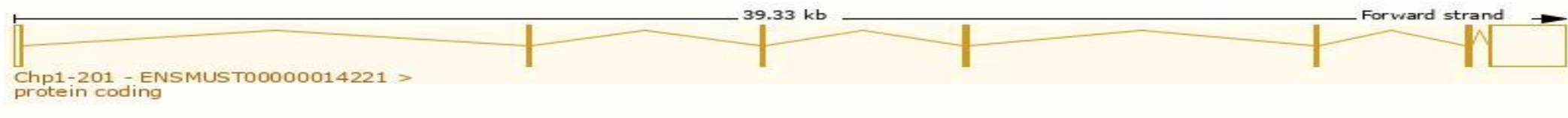
Source: <https://www.ncbi.nlm.nih.gov/>

Transcript Information

The gene has 5 transcripts, all transcripts are shown below:

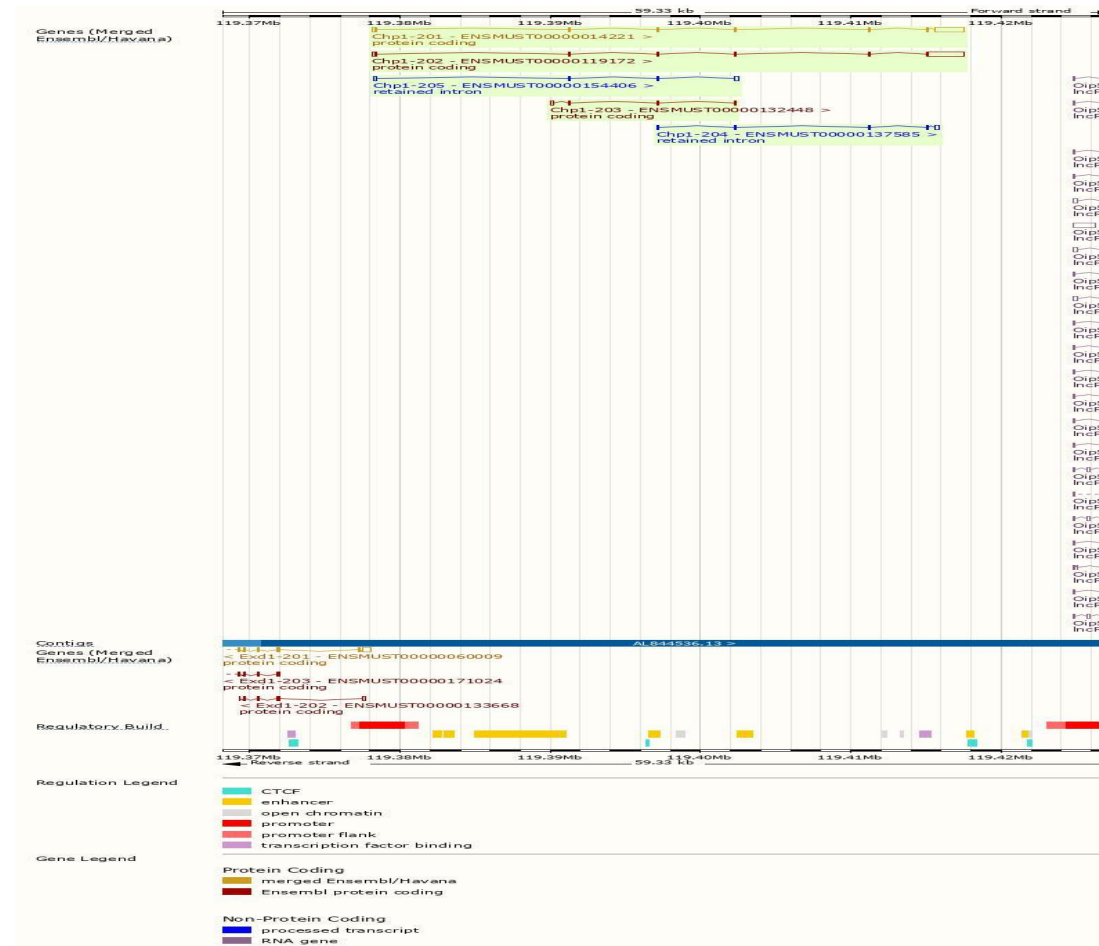
Transcript ID	Name	bp	Protein	Biotype	CCDS	UniProt Match	Flags
ENSMUST00000014221.13	Chp1-201	2609	195aa	Protein coding	CCDS16604	P61022	Ensembl Canonical GENCODE basic APPRIS P1 TSL:1
ENSMUST00000119172.2	Chp1-202	3027	191aa	Protein coding		B0R091	GENCODE basic TSL:2
ENSMUST00000132448.8	Chp1-203	518	33aa	Protein coding		B0R092	TSL:5 CDS 3' incomplete
ENSMUST00000137585.3	Chp1-204	738	No protein	Retained intron		-	TSL:2
ENSMUST00000154406.8	Chp1-205	543	No protein	Retained intron		-	TSL:2

The strategy is based on the design of *Chp1*-201 transcript, the transcription is shown below:

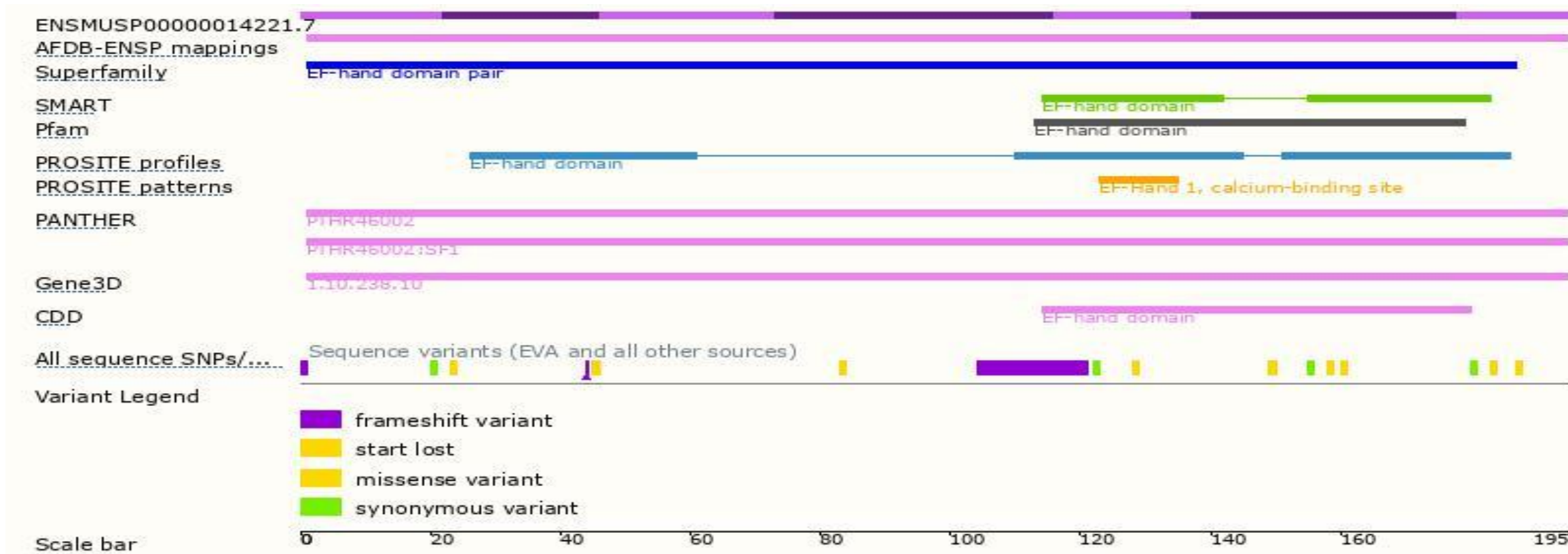


Source: <https://www.ensembl.org>

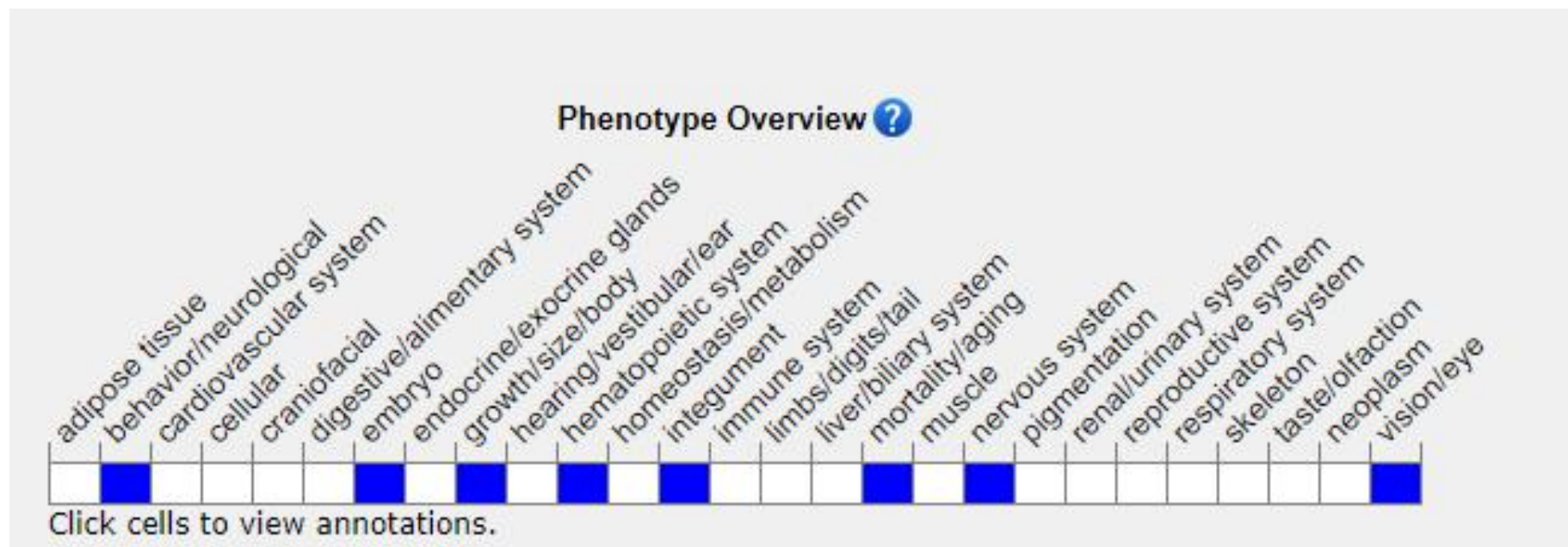
Genomic Information



Protein Information



Mouse Phenotype Information (MGI)



- Mice homozygous for an ENU mutation display ataxia and progressive Purkinje cell axonal dystrophy.

Important Information

- *Chp1* is located on Chr2. 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.

References



<https://www.mousephenotype.org/data/genes/MGI:1927185>

Mutation details: The L1L2_Bact_P cassette was inserted at position **119401802** of Chromosome 2 upstream of the critical exon(s) (Build GRCm39). The cassette is composed of an FRT site followed by lacZ sequence and a loxP site. This first loxP site is followed by a neomycin resistance gene under the control of the human beta-actin promoter, SV40 polyA, a second FRT site and a second loxP site. A third loxP site is inserted downstream of the targeted exon(s) at position **119402773**. The critical exon(s) is/are thus flanked by loxP sites. A "conditional ready" (floxed) allele can be created by flp recombinase expression in mice carrying this allele. Subsequent cre expression results in a knockout mouse. If cre expression occurs without flp expression, a reporter knockout mouse will be created. Further information on targeting strategies used for this and other IKMC alleles can be found at http://www.informatics.jax.org/mgihome/nomen/IKMC_schematics.shtml (J:157065)

<https://www.informatics.jax.org/allele/MGI:4842153>