

# ***Cd84 Cas9-CKO Strategy***

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**Reviewer:**

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# Project Overview

**Project Name**

***Cd84***

**Project type**

**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

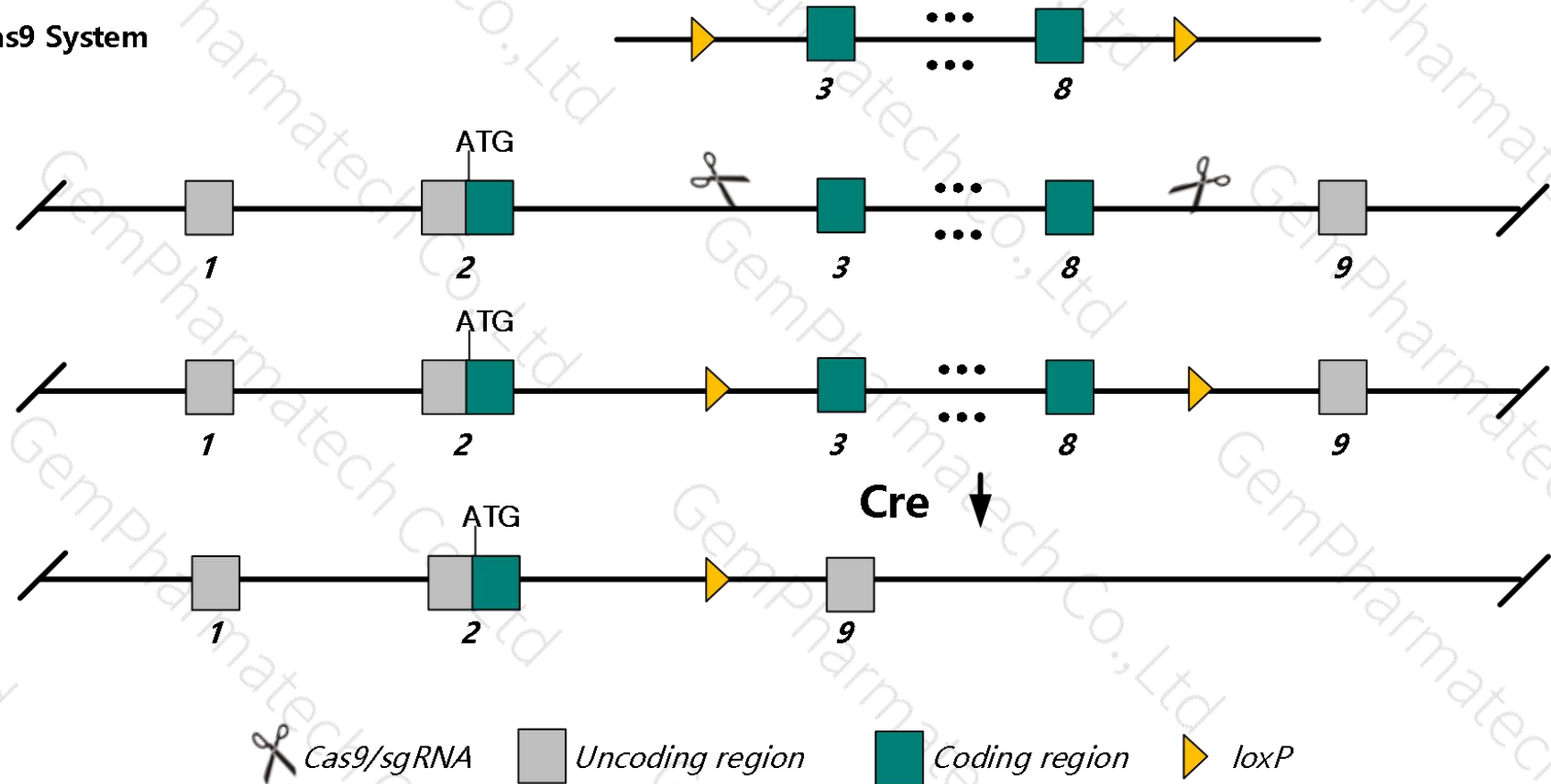
This model will use CRISPR/Cas9 technology to edit the *Cd84* gene. The schematic diagram is as follows:

## Donor and CRISPR/Cas9 System

### Wild-type allele

### Floxed allele

### KO allele



# Technical routes

- The *Cd84* gene has 5 transcripts. According to the structure of *Cd84* gene, exon3-exon8 of *Cd84-205* (ENSMUST00000155802.7) transcript is recommended as the knockout region. The region contains most coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Cd84* 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 knock-out allele of this gene show defects in T follicular helper function and germinal center formation. Mice homozygous for a different knock-out allele display normal platelet physiology and thrombus formation.
- The *Cd84* gene is located on the Chr1. 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)

## Cd84 CD84 antigen [ *Mus musculus* (house mouse) ]

Gene ID: 12523, updated on 10-Oct-2019

### Summary

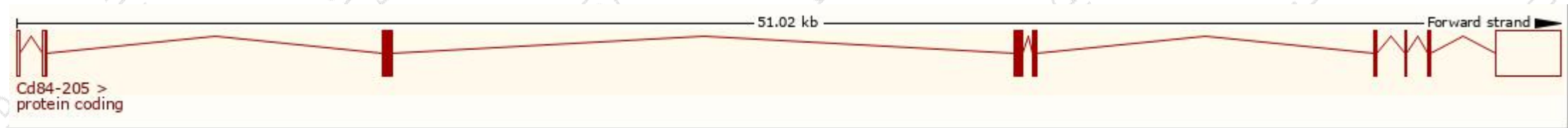
Official Symbol	Cd84 provided by <a href="#">MGI</a>
Official Full Name	CD84 antigen provided by <a href="#">MGI</a>
Primary source	<a href="#">MGI:MGI:1336885</a>
See related	<a href="#">Ensembl:ENSMUSG00000038147</a>
Gene type	protein coding
RefSeq status	VALIDATED
Organism	<a href="#">Mus musculus</a>
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	CDw84; SLAMF5; A130013D22Rik
Expression	Biased expression in thymus adult (RPKM 2.1), spleen adult (RPKM 1.0) and 14 other tissues <a href="#">See more</a>
Orthologs	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

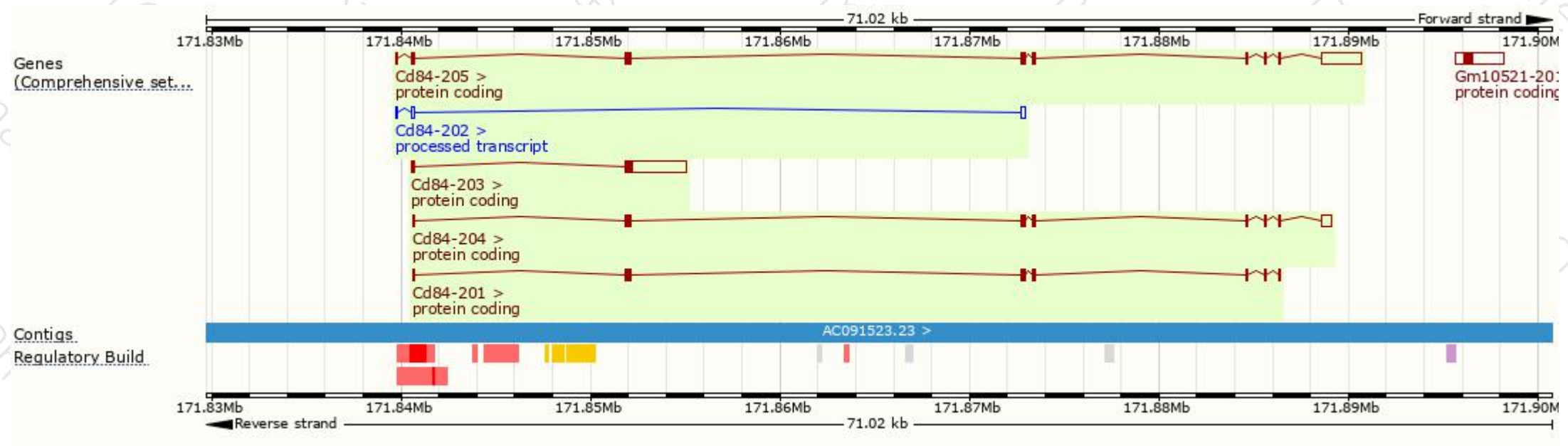
The gene has 5 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cd84-203	<a href="#">ENSMUST00000135386.1</a>	3434	<a href="#">140aa</a>	Protein coding	<a href="#">CCDS56657</a>	<a href="#">A0A0R4J1S4</a>	TSL:1 GENCODE basic APPRIS ALT2
Cd84-205	<a href="#">ENSMUST00000155802.7</a>	3291	<a href="#">329aa</a>	Protein coding	<a href="#">CCDS15503</a>	<a href="#">A0A0R4J0K5</a>	TSL:1 GENCODE basic APPRIS P3
Cd84-204	<a href="#">ENSMUST00000136479.7</a>	1503	<a href="#">328aa</a>	Protein coding	<a href="#">CCDS69979</a>	<a href="#">E9Q9E8</a>	TSL:1 GENCODE basic APPRIS ALT2
Cd84-201	<a href="#">ENSMUST00000042302.6</a>	1019	<a href="#">329aa</a>	Protein coding	<a href="#">CCDS15503</a>	<a href="#">A0A0R4J0K5</a>	TSL:1 GENCODE basic APPRIS P3
Cd84-202	<a href="#">ENSMUST00000128189.1</a>	454	No protein	Processed transcript	-	-	TSL:3

The strategy is based on the design of *Cd84-205* transcript,The transcription is shown below

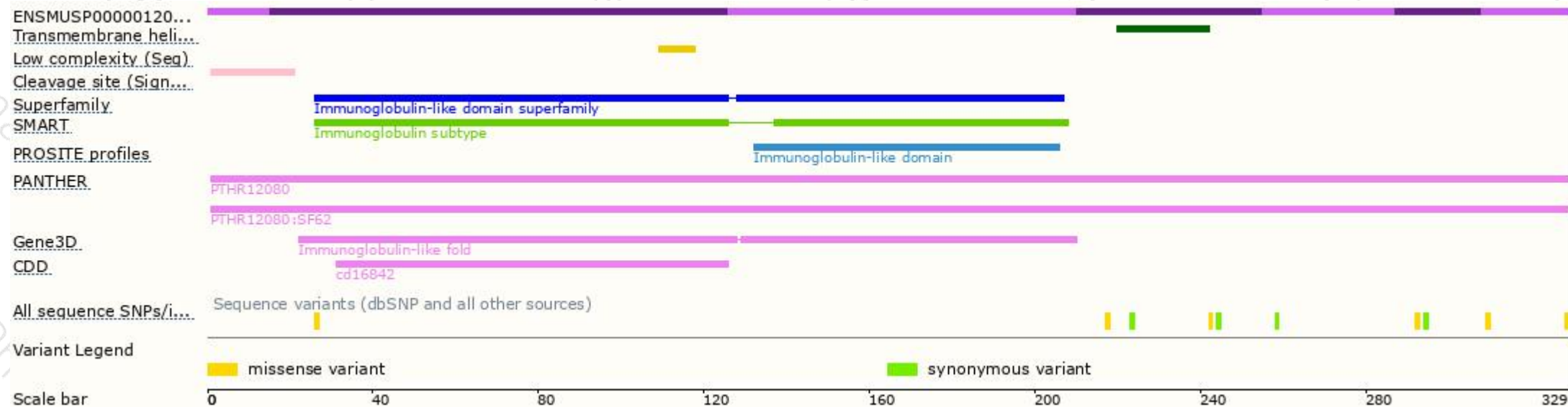


# Genomic location distribution

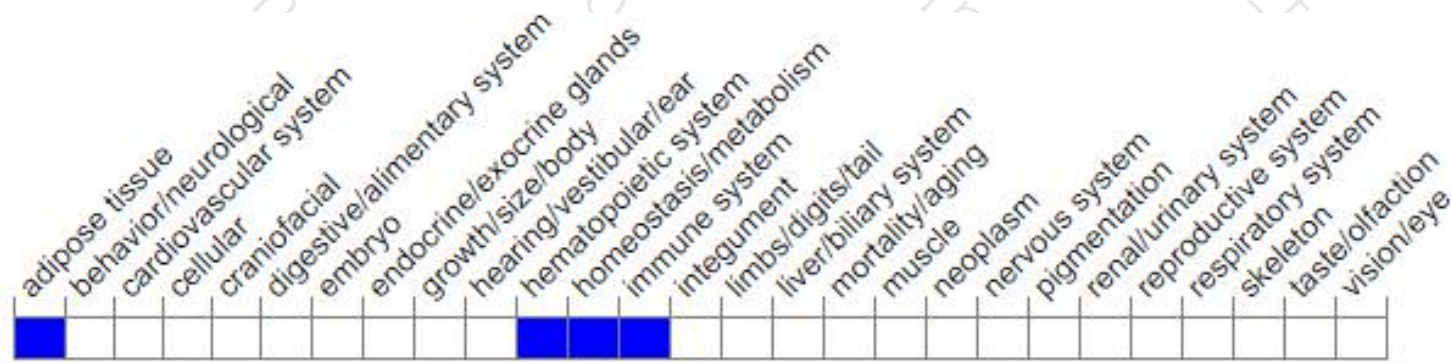




# Protein domain



# 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 knock-out allele of this gene show defects in T follicular helper function and germinal center formation. Mice homozygous for a different knock-out allele display normal platelet physiology and thrombus formation.

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

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