

# Tiam1 Cas9-KO Strategy

Designer: Daohua Xu

Reviewer: Huimin Su

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# **Project Overview**



**Project Name** 

Tiam 1

**Project type** 

Cas9-KO

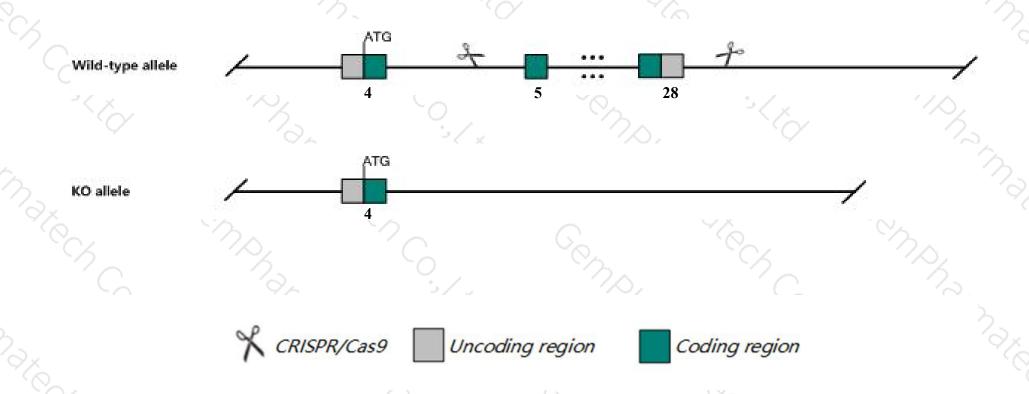
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Tiam1* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Tiam1* gene has 15 transcripts. According to the structure of *Tiam1* gene, exon5-exon28 of *Tiam1-201* (ENSMUST00000002588.10) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Tiam1* gene. The brief process is as follows: CRISPR/Cas9 system

### **Notice**



- ➤ According to the existing MGI data, Mice homozygous for a targeted null allele display resistance to chemically-induced tumors, however, tumors that do develop progress to malignancy. Mice homozygous for a gene trap allele display anencephaly, exencephaly and/or neural tube defects.
- > The *Tiam1* gene is located on the Chr16. 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)



#### Tiam1 T cell lymphoma invasion and metastasis 1 [Mus musculus (house mouse)]

Gene ID: 21844, updated on 31-Jan-2019

#### Summary

☆ ?

Official Symbol Tiam1 provided by MGI

Official Full Name T cell lymphoma invasion and metastasis 1 provided by MGI

Primary source MGI:MGI:103306

See related Ensembl:ENSMUSG00000002489

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 Al847750, D16lum10, D16lum10e

Expression Broad expression in cerebellum adult (RPKM 9.6), frontal lobe adult (RPKM 5.9) and 22 other tissuesSee more

Orthologs human all

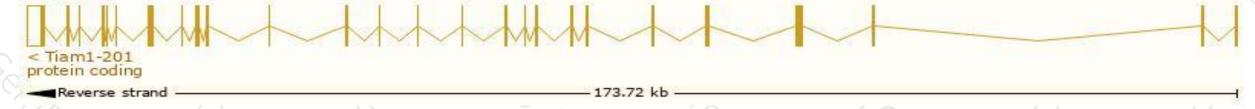
# Transcript information (Ensembl)



#### The gene has 15 transcripts, all transcripts are shown below:

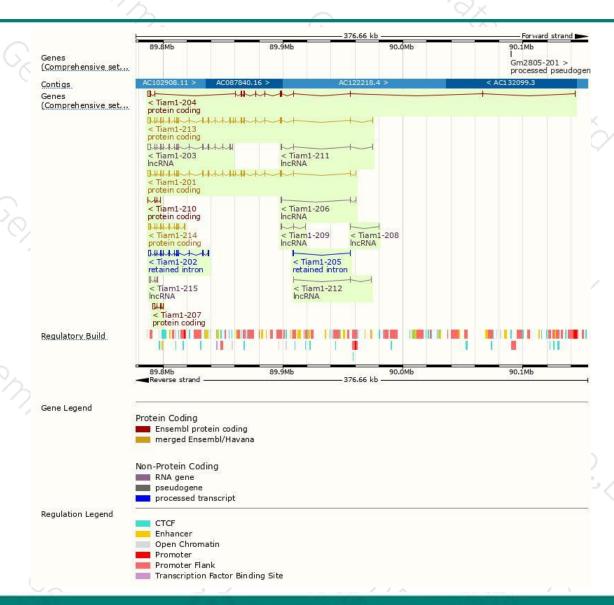
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Tiam1-201	ENSMUST00000002588.10	7289	<u>1591aa</u>	Protein coding	CCDS28314	G3UWG2	TSL:1 GENCODE basic APPRIS P1
Tiam1-213	ENSMUST00000163370.7	7255	<u>1591aa</u>	Protein coding	CCDS28314	G3UWG2	TSL:1 GENCODE basic APPRIS P1
Tiam1-214	ENSMUST00000164263.8	3569	<u>622aa</u>	Protein coding	CCDS49904	E9Q1R7	TSL:1 GENCODE basic
Tiam1-204	ENSMUST00000114124.8	5203	900aa	Protein coding	-	Q6P1D6	TSL:5 GENCODE basic
Tiam1-207	ENSMUST00000134021.1	1770	<u>168aa</u>	Protein coding		J3QQ55	CDS 5' incomplete TSL:1
Tiam1-210	ENSMUST00000144691.8	321	92aa	Protein coding	-	J3QMH1	CDS 5' incomplete TSL:5
Tiam1-202	ENSMUST00000089084.11	3495	No protein	Retained intron			TSL:1
Tiam1-205	ENSMUST00000124433.7	667	No protein	Retained intron	-	2	TSL:1
Γiam1-203	ENSMUST00000114122.7	4596	No protein	IncRNA	5	5	TSL:1
Tiam1-215	ENSMUST00000178095.1	1465	No protein	IncRNA	-	-	TSL:1
Tiam1-211	ENSMUST00000151655.7	726	No protein	IncRNA		2	TSL:2
Tiam1-209	ENSMUST00000139544.1	503	No protein	IncRNA	-	2	TSL:3
Tiam1-206	ENSMUST00000130951.7	461	No protein	IncRNA	5	5	TSL:2
Γiam1-212	ENSMUST00000155915.1	446	No protein	IncRNA		-	TSL:2
Tiam1-208	ENSMUST00000136760.1	241	No protein	IncRNA		-	TSL:5

The strategy is based on the design of *Tiam1-201* transcript, The transcription is shown below



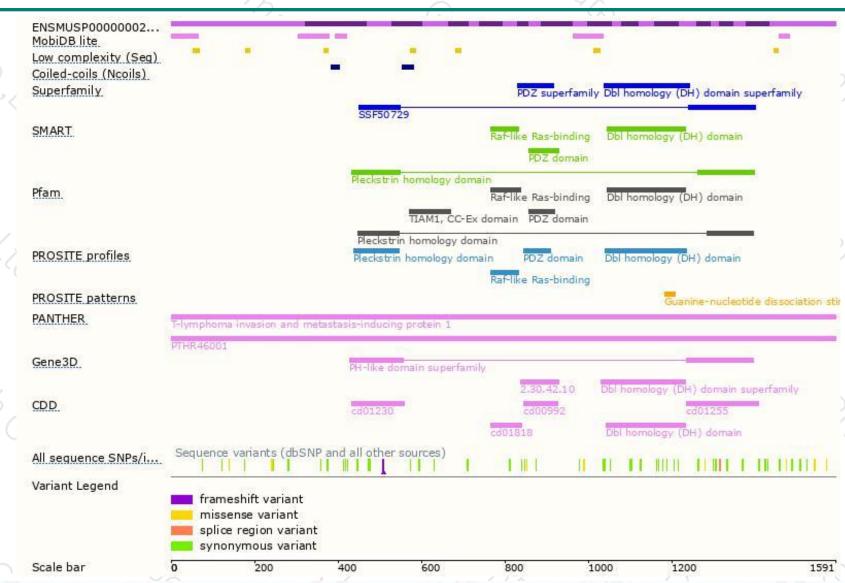
### Genomic location distribution





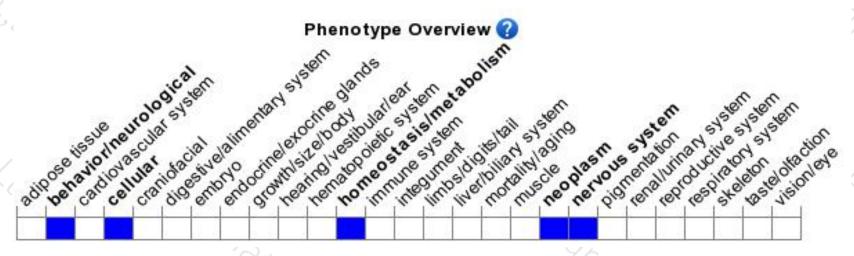
### Protein domain





# 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 targeted null allele display resistance to chemically-induced tumors, however, tumors that do develop progress to malignancy. Mice homozygous for a gene trap allele anencephaly, exencephaly and/or neural tube defects.



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





