

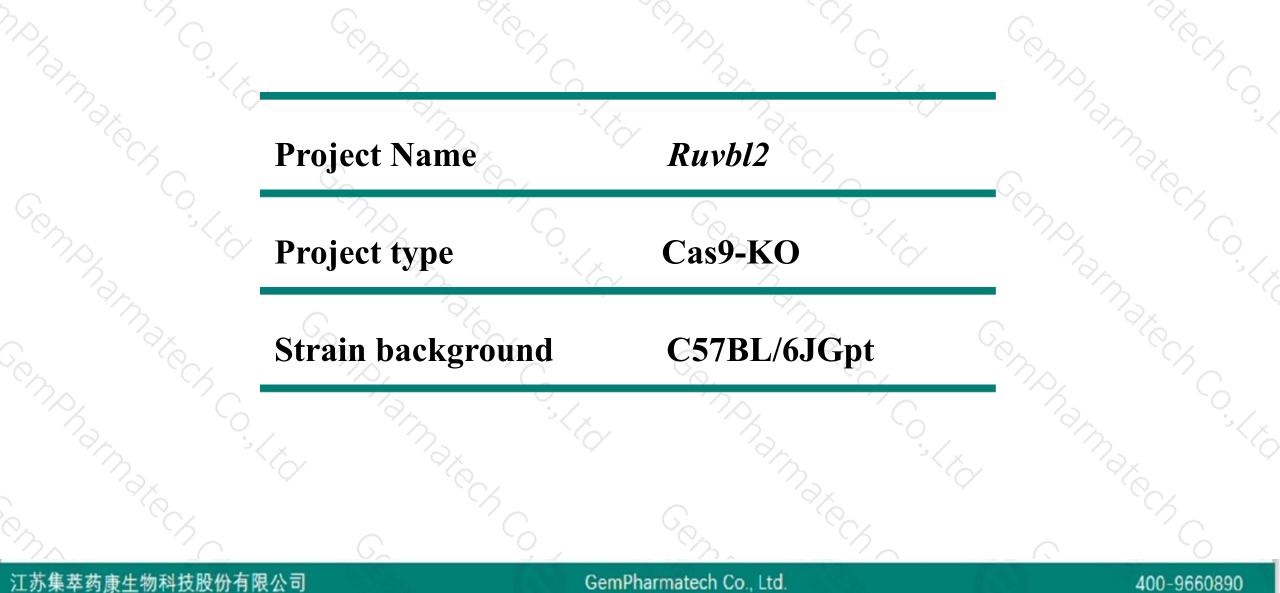
# Ruvbl2 Cas9-KO Strategy Romphamater Control

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## **Project Overview**

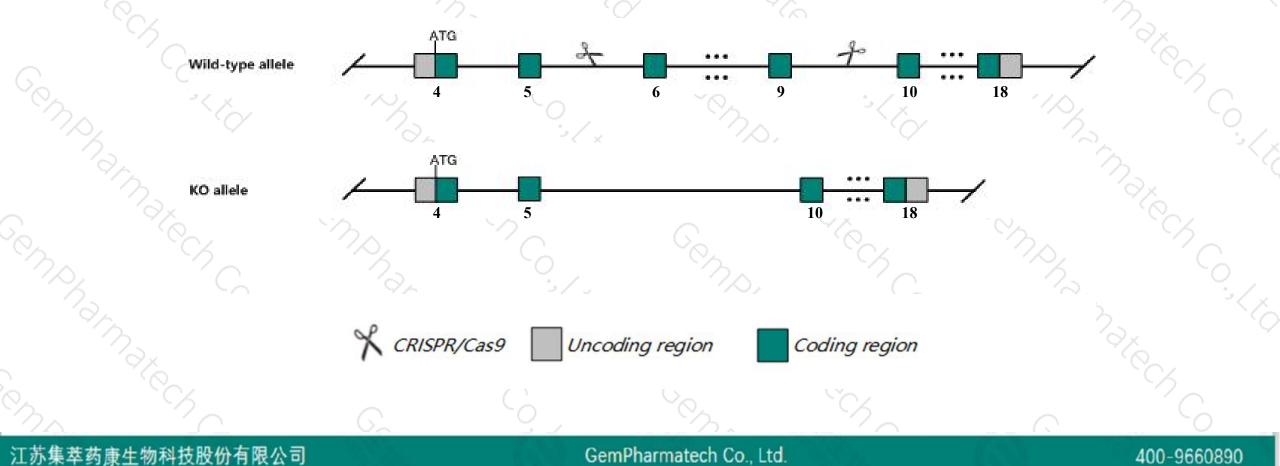




# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Ruvbl2 gene. The schematic diagram is as follows:





- The Ruvbl2 gene has 8 transcripts. According to the structure of Ruvbl2 gene, exon6-exon9 of Ruvbl2-205 (ENSMUST00000211214.1) transcript is recommended as the knockout region. The region contains 395bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Ruvbl2 gene. The brief process is as follows: CRISPR/Cas9 system



- According to the existing MGI data, Mice homozygous for an ENU-induced allele exhibit lethality. Mice heterozygous for a knock-out allele exhibit impaired T cell development and maximal T dependent antibody responses.
- The *Ruvbl2* gene is located on the Chr7. 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)**



\$ ?

## Ruvbl2 RuvB-like protein 2 [Mus musculus (house mouse)]

Gene ID: 20174, updated on 7-Apr-2019

#### Summary

Official Symbol	Ruvbl2 provided by MGI
Official Full Name	RuvB-like protein 2 provided by MGI
Primary source	MGI:MGI:1342299
See related	Ensembl:ENSMUSG0000003868
Gene type	protein coding
<b>RefSeq status</b>	PROVISIONAL
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;
	Muroidea; Muridae; Murinae; Mus; Mus
Also known as	mp47, p47
Expression	Ubiquitous expression in testis adult (RPKM 61.6), CNS E11.5 (RPKM 48.3) and 28 other tissues See more
Orthologs	human all

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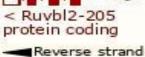
## **Transcript information (Ensembl)**



## The gene has 8 transcripts, all transcripts are shown below:

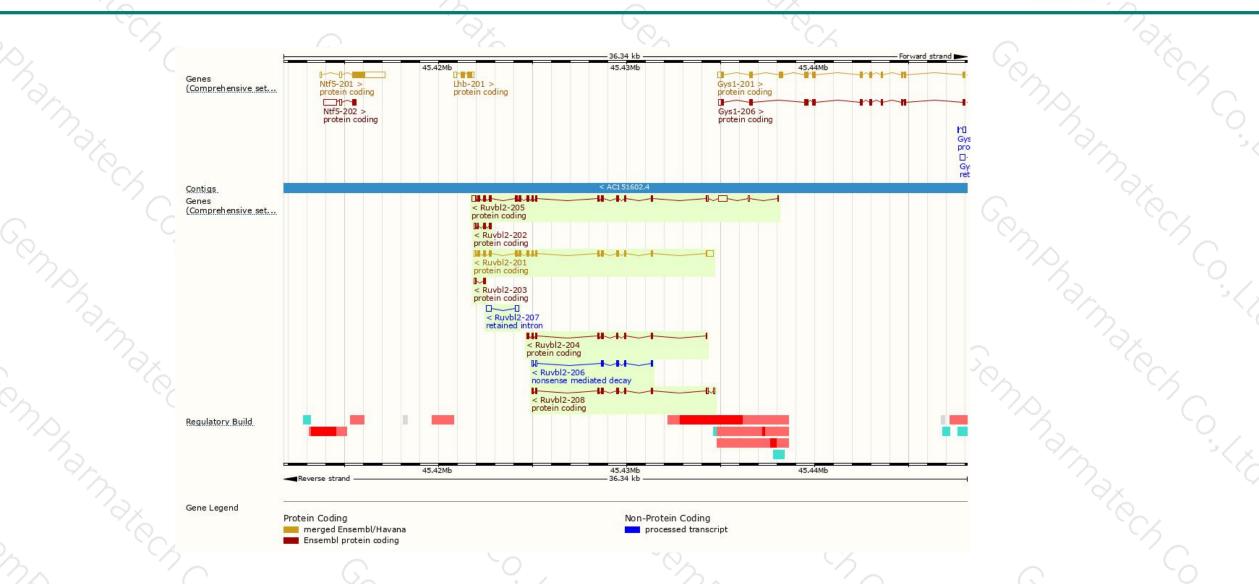
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ruvbl2-205	ENSMUST00000211214.1	2312	<u>463aa</u>	Protein coding	CCDS21243	Q3TXT7 Q9WTM5	TSL:1 GENCODE basic APPRIS P1
Ruvbl2-201	ENSMUST00000107771.11	1834	<u>463aa</u>	Protein coding	CCDS21243	Q3TXT7 Q9WTM5	TSL:1 GENCODE basic APPRIS P1
Ruvbl2-208	ENSMUST00000211666.1	846	<u>221aa</u>	Protein coding	-	A0A1B0GSR4	CDS 3' incomplete TSL:3
Ruvbl2-204	ENSMUST00000210439.1	725	<u>237aa</u>	Protein coding	-	A0A1B0GRW3	CDS 3' incomplete TSL:5
Ruvbl2-202	ENSMUST00000209426.1	396	<u>104aa</u>	Protein coding	5	A0A1B0GR89	CDS 5' incomplete TSL:3
Ruvbl2-203	ENSMUST00000210271.1	219	<u>48aa</u>	Protein coding		A0A1B0GRT1	CDS 5' incomplete TSL:3
Ruvbl2-206	ENSMUST00000211440.1	585	<u>134aa</u>	Nonsense mediated decay	-	A0A1B0GT54	CDS 5' incomplete TSL:5
Ruvbl2-207	ENSMUST00000211478.1	474	No protein	Retained intron	2 2	10 <u>1</u> 20	TSL:3

The strategy is based on the design of Ruvbl2-205 transcript, The transcription is shown below



— 16.34 kb -

# **Genomic location distribution**



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## **Protein domain**



ENSMUSP00000103... Low complexity (Seg) Conserved Domains hmmpanther

Superfamily domains SMART domains Prints domain Pfam domain Gene3D

RuvB-like RuvB-like helicase 2 P-loop containing nucleoside triphosphate hydrolase AAA+ ATPase domain PR01874 TIP49, C-terminal 2.40.50.360 1,10,8,60 3,40,50,300 Sequence variants (dbSNP and all other sources) LП synonymous variant 80 120 160 200 240 280 320 360 400 463 40

All sequence SNPs/i...

0

Variant Legend

Scale bar

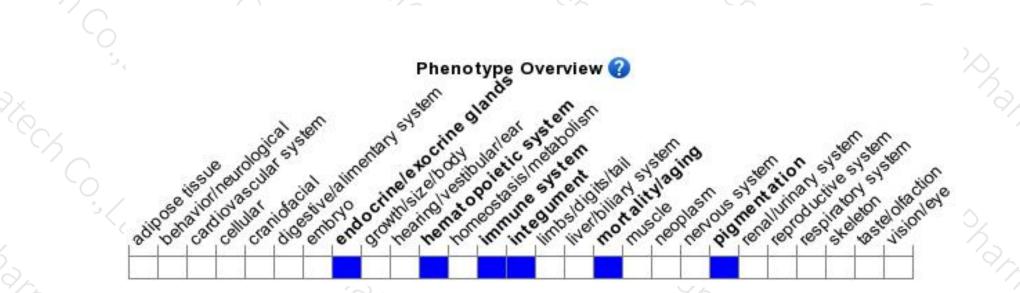
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## Mouse phenotype description(MGI)





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 an ENU-induced allele exhibit lethality. Mice heterozygous for a knock-out allele exhibit impaired T cell development and maximal T dependent antibody responses.



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



