

# Eif4a3 Cas9-KO Strategy

**Designer:** 

**Reviewer:** 

**Design Date:** 

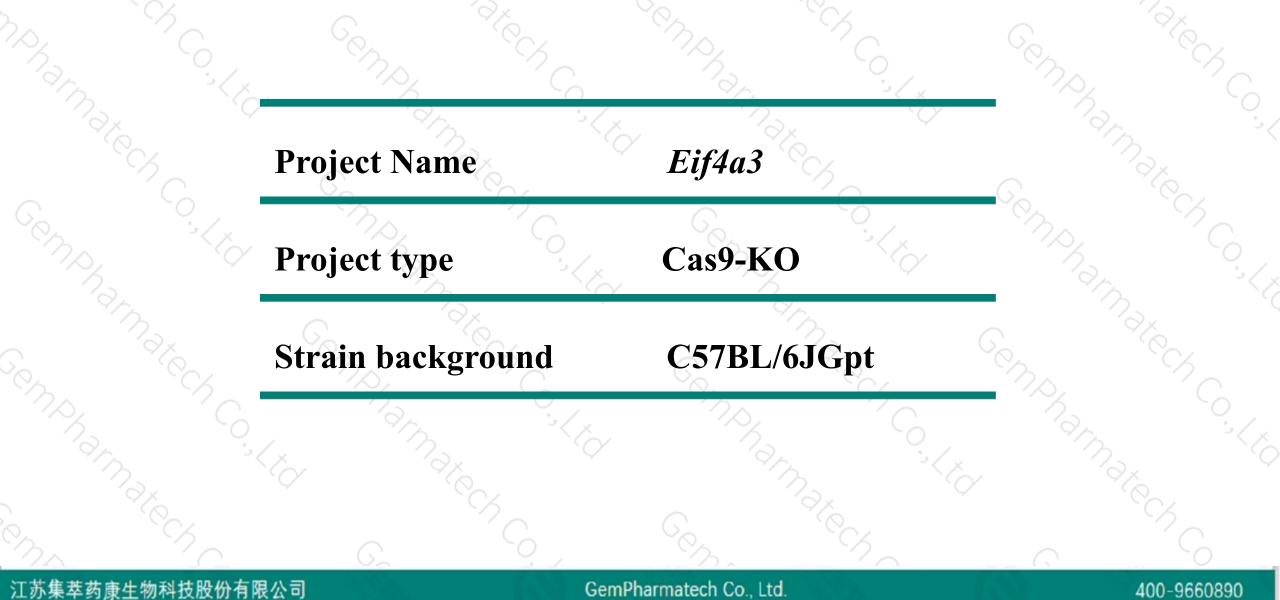
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# **Project Overview**

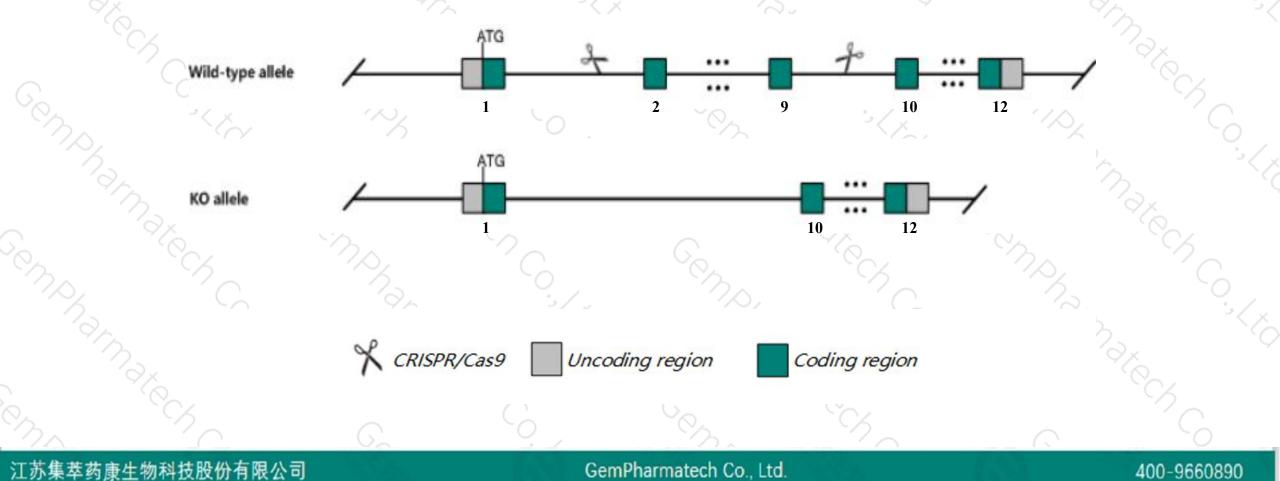




# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Eif4a3* gene. The schematic diagram is as follows:





- The *Eif4a3* gene has 6 transcripts. According to the structure of *Eif4a3* gene, exon2-exon9 of *Eif4a3-201* (ENSMUST0000026667.14) transcript is recommended as the knockout region. The region contains 814bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify *Eif4a3* gene. The brief process is as follows: CRISPR/Cas9 system

- > The *Eif4a3* gene is located on the Chr11. 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.

Notice

# **Gene information (NCBI)**



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#### Eif4a3 eukaryotic translation initiation factor 4A3 [Mus musculus (house mouse)]

Gene ID: 192170, updated on 13-Mar-2020

#### Summary

Official Symbol	Eif4a3 provided by MGI								
Official Full Name	eukaryotic translation initiation factor 4A3 provided by MGI								
Primary source	MGI:MGI:1923731								
See related	Ensembl:ENSMUSG0000025580								
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	2400003O03Rik, Ddx48, elF4A-III, mKIAA0111								
Expression	Ubiquitous expression in CNS E11.5 (RPKM 64.0), limb E14.5 (RPKM 53.1) and 28 other tissues See more								
Orthologs	human all								

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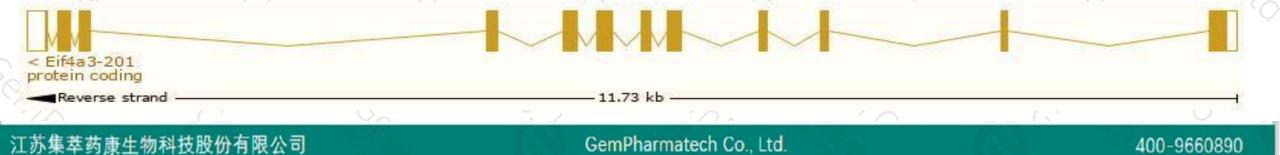
# **Transcript information (Ensembl)**



#### The gene has 6 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Eif4a3-201	ENSMUST00000026667.14	1509	<u>411aa</u>	Protein coding	CCDS25714	Q91VC3	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P
Eif4a3-202	ENSMUST00000106253.1	916	<u>299aa</u>	Protein coding	88	A2AFK7	CDS 3' incomplete TSL:5
Eif4a3-206	ENSMUST00000138849.1	613	No protein	Processed transcript	82	2	TSL:2
Eif4a3-205	ENSMUST00000135294.1	473	No protein	Processed transcript	12		TSL:3
Eif4a3-203	ENSMUST00000129044.7	466	No protein	Processed transcript	17	-	TSL2
Eif4a3-204	ENSMUST00000134743.1	377	No protein	Processed transcript		-	TSL:2

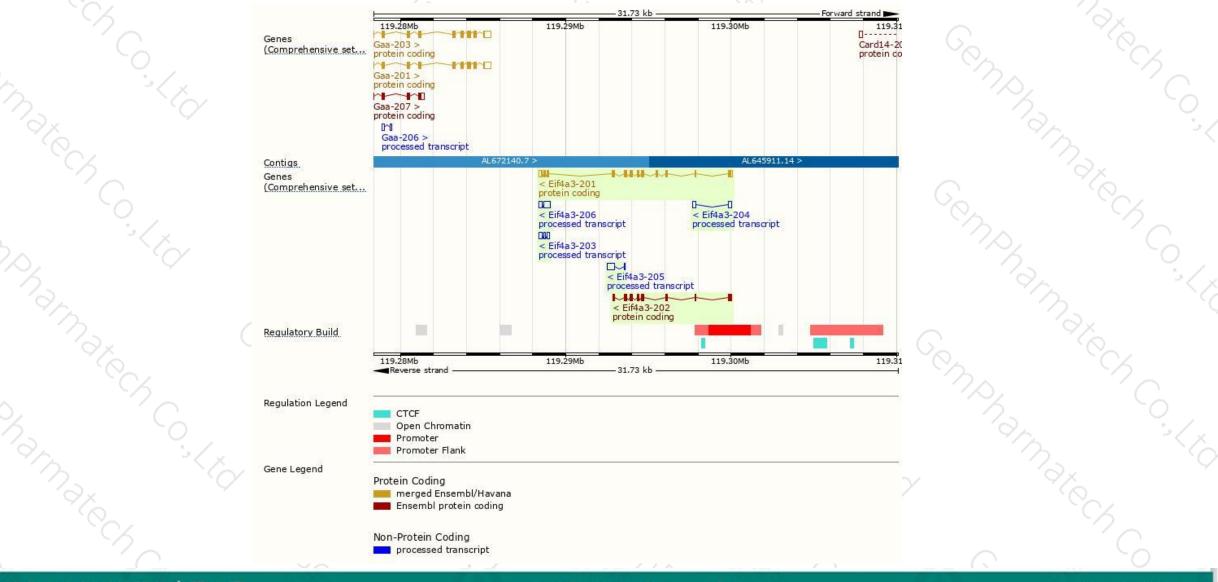
The strategy is based on the design of *Eif4a3-201* transcript, the transcription is shown below:



### **Genomic location distribution**



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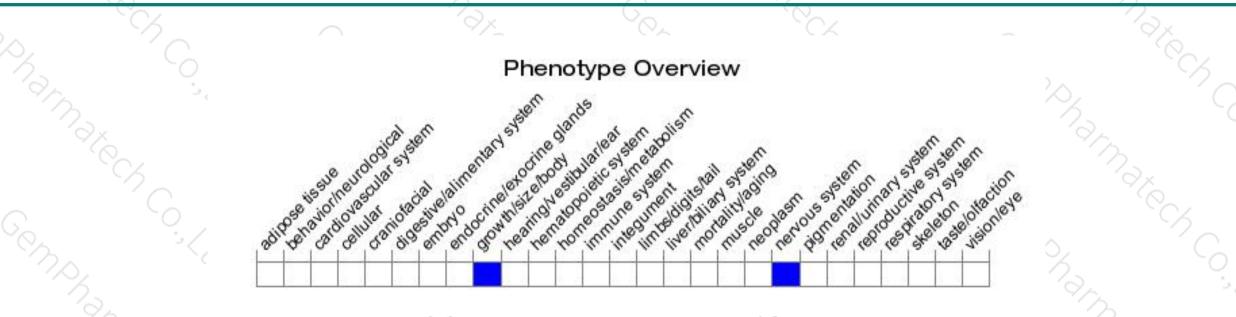
# **Protein domain**



Superfamily	P-loop containing nucleoside triphosphate hydrolase								
SMART	Helicase superfamily 1/2, ATP-binding domain	Helicase, C-terminal							
Pfam.	DEAD/DEAH box helicase domain	Helicase, C-terminal							
PROSITE profiles	RNA helicase, DEAD-box type, Q motif	Helicase, C-terminal	-						
Helicase superfamily 1/2, ATP-binding domain PROSITE patterns ATP-dependent RNA helicase DEAD-box, conserved site									
PANTHER	PTHR24031 PTHR24031:SF57								
Gene3D 3	.40.50.300 cd18045	cd18787							
All sequence SNPs/i Seque	nce variants (dbSNP and all other sources)	(J. (J.) J. J.	) <b>1</b>						
	issense variant /nonymous variant								
Scale bar 0	40 80 120 160	200 240 280 320 360	) 411						

# Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(http://www.informatics.jax.org/).



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



