

Isl2 Cas9-KO Strategy

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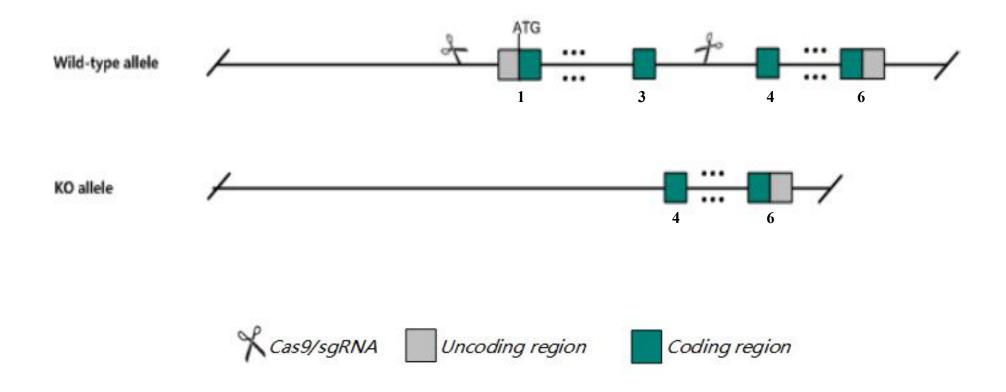


Project Name	Isl2			
Project type	Cas9-KO			
Strain background	C57BL/6JGpt			

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Isl2 gene. The schematic diagram is as follows:





The *Isl2* gene has 5 transcripts. According to the structure of *Isl2* gene, exon1-exon3 of *Isl2-*201(ENSMUST00000034869.11) transcript is recommended as the knockout region. The region contains start codon ATG.Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Isl2* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA 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.



According to the existing MGI data, mutations of this gene result in neonatal lethality, motor neuron migration defects and impaired visceral motor neuron differentiation.

The *Isl2* gene is located on the Chr9. 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

Isl2 insulin related protein 2 (islet 2) [Mus musculus (house mouse)]

Gene ID: 104360, updated on 12-Jan-2021

Summary

Official Symbol	Isl2 provided by MGI					
Official Full Name	insulin related protein 2 (islet 2) provided by MGI					
Primary source	MGI:MGI:109156					
See related	Ensembl:ENSMUSG0000032318					
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	3110001N10Rik, isle					
Expression	Biased expression in CNS E11.5 (RPKM 2.3), colon adult (RPKM 1.4) and 11 other tissuesSee more					
Orthologs	human all					



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Transcript information Ensembl

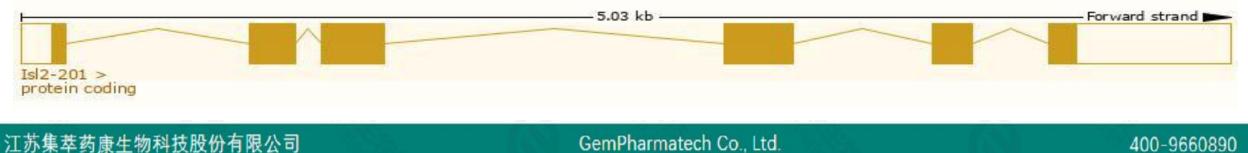


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The gene has 5 transcripts, all transcripts are shown below:

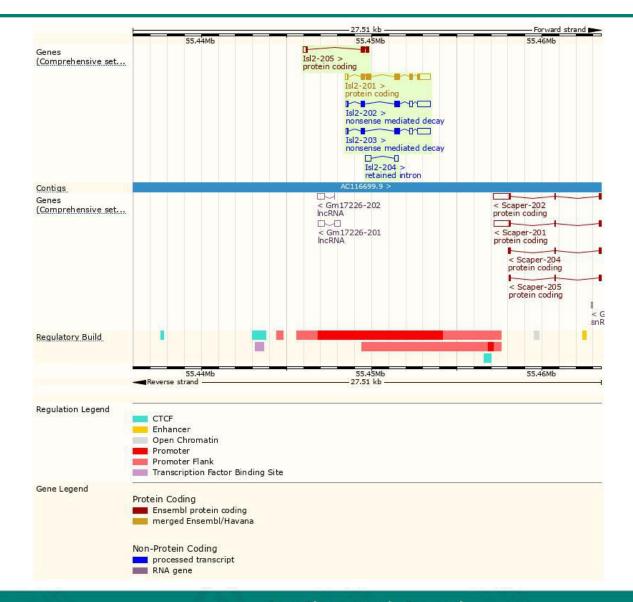
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Isl2-201	ENSMUST0000034869.11	1854	<u>359aa</u>	Protein coding	CCDS23205		TSL:1 , GENCODE basic , APPRIS P1 ,
Isl2-205	ENSMUST00000175950.8	637	<u>151aa</u>	Protein coding	(-)		CDS 3' incomplete , TSL:3 ,
Isl2-203	ENSMUST00000164373.8	1535	<u>174aa</u>	Nonsense mediated decay			TSL:1,
Isl2-202	ENSMUST00000114290.4	1532	<u>174aa</u>	Nonsense mediated decay			TSL:1,
Isl2-204	ENSMUST00000165302.2	529	No protein	Retained intron	(1)		TSL:3,

The strategy is based on the design of *Isl2-201* transcript, the transcription is shown below:



Genomic location distribution





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Protein domain



ENSMUSP00000034 MobiDB lite		
Low complexity (Seq) Superfamily	SSF57716	
SMART	Zinc finger, LIM-type	Homeobox-like domain superfamily
Pfam	Zinc finger, LIM-type	Homeobox domain
PROSITE profiles	Zinc finger, LIM-type	Homeobox domain
PROSITE patterns	Zinc finger, LIM-type	Homeobox, conserved site
PANTHER	PTHR 24204:SF2	
	PTHR24204	
Gene3D	2.10.110.10	1.10.10.50
CDD	cd09366 cd09471	Homeoblox domain
All sequence SNPs/i	Sequence variants (dbSNP and all other sources	
Variant Legend	missense variant	
	synonymous variant	
Scale bar	0 40 80 120	160 200 240 280 359

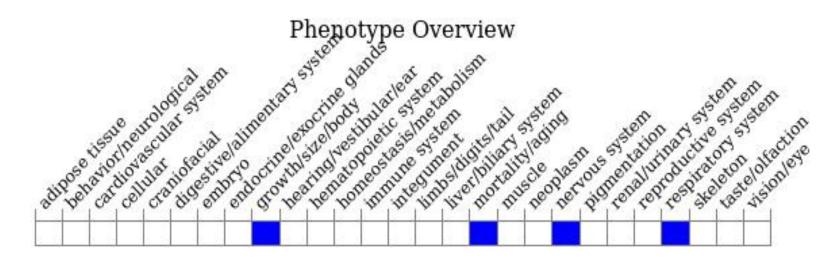
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Mouse phenotype description(MGI)





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, mutations of this gene result in neonatal lethality, motor neuron migration defects and impaired visceral motor neuron differentiation.



If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





