Six1 Cas9-KO Strategy Rond almakech Co. / Ky

Designer: Conplainax Ch

Qiong Zhou ary Co. (x

Project Overview



Project Name

Six1

Project type

Cas9-KO

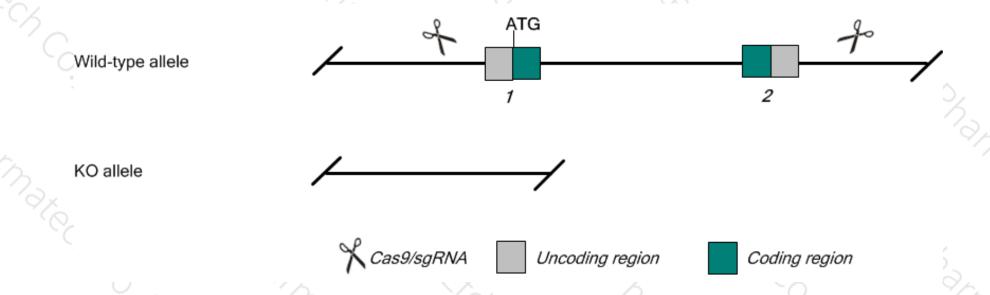
Animal background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Six1 gene. The schematic diagram is as follows:



Technical routes



- The Six1 gene has 4 transcripts. According to the structure of Six1 gene, exon1-2 of Six1-201 (ENSMUST00000050029.7) transcript is recommended as the knockout region. The region contains all coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Six1* 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.

Notice



- According to the existing MGI data, Homozygous inactivation of this gene causes perinatal lethality associated with severe muscle hypoplasia, rib defects, absence of kidneys and thymus, craniofacial anomalies, as well as defects in neurogenesis and ear, nasal, and gland development. Heterozygotes may show variable hearing loss.
- ➤ The *Six1* gene is located on the Chr12. 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)



Six1 sine oculis-related homeobox 1 [Mus musculus (house mouse)]

Gene ID: 20471, updated on 19-Mar-2019

Summary

Official Symbol Six1 provided by MGI

Official Full Name sine oculis-related homeobox 1 provided by MGI

Primary source MGI:MGI:102780

See related Ensembl: ENSMUSG00000051367

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 BB138287

Expression Biased expression in limb E14.5 (RPKM 14.6), CNS E11.5 (RPKM 6.0) and 7 other tissues See more

Orthologs human all

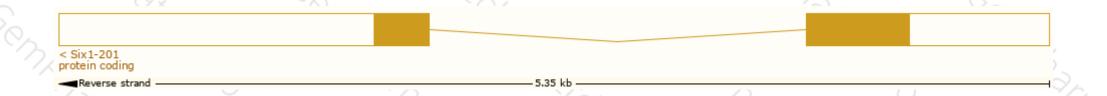
Transcript information (Ensembl)



The gene has 4 transcripts, and all transcripts are shown below:

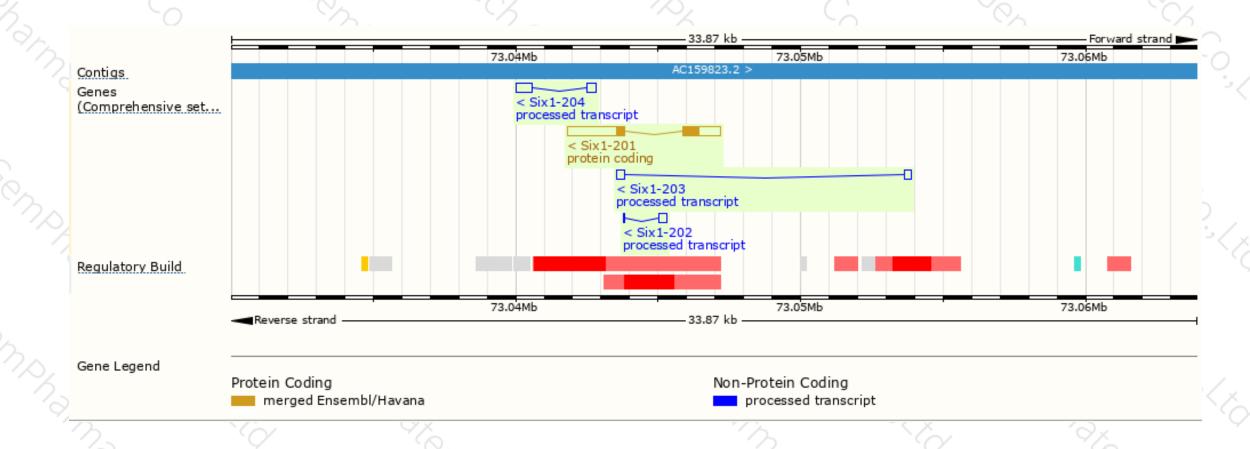
Name 🌲	Transcript ID 👙	bp 🌲	Protein 🌲	Biotype	CCDS	UniProt 🍦	Flags 🌲
Six1-201	ENSMUST00000050029.7	3316	<u>284aa</u>	Protein coding	<u>CCDS25973</u> ₽	<u>Q3√2C3</u> ₽ <u>Q62231</u> ₽	TSL:1 GENCODE basic APPRIS P1
Six1-204	ENSMUST00000176310.1	885	No protein	Processed transcript	-	-	TSL:3
Six1-203	ENSMUST00000176091.1	528	No protein	Processed transcript	-	-	TSL:2
Six1-202	ENSMUST00000175677.1	334	No protein	Processed transcript	-	-	TSL:3

The strategy is based on the design of Six1-201 transcript, The transcription is shown below



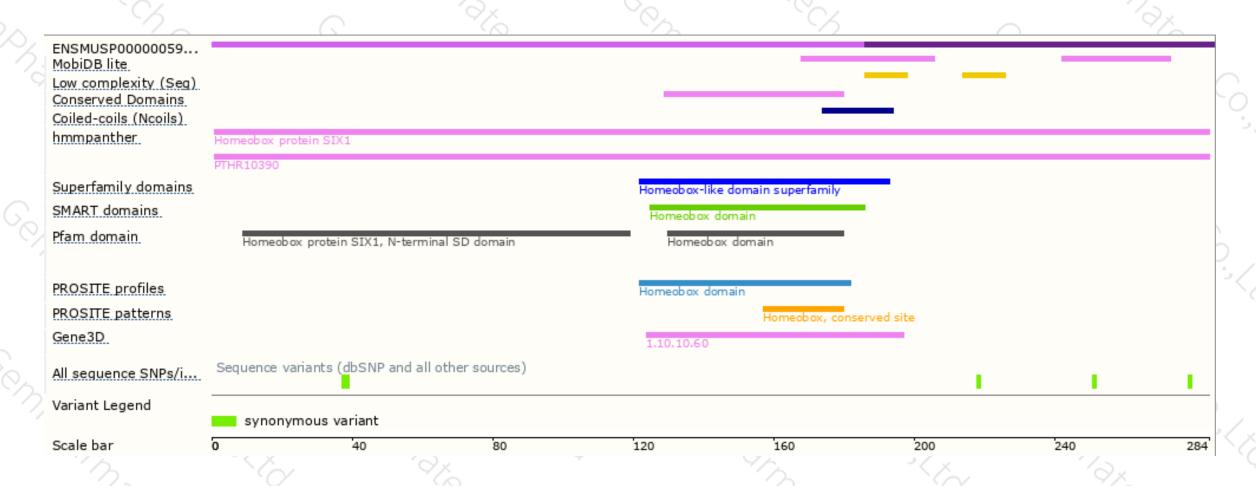
Genomic location distribution





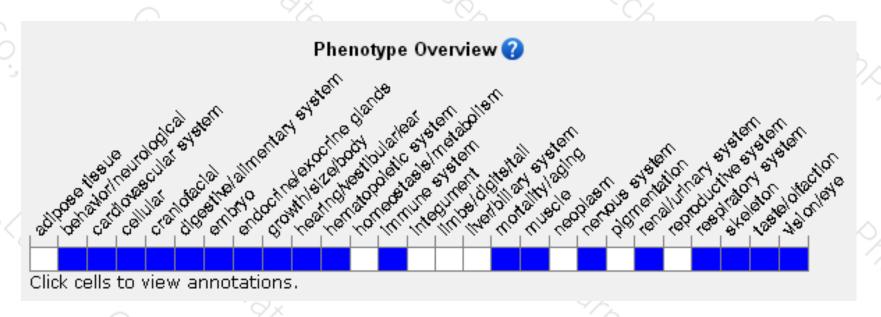
Protein domain





Mouse phenotype description(MGI)





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

Homozygous inactivation of this gene causes perinatal lethality associated with severe muscle hypoplasia, rib defects, absence of kidneys and thymus, craniofacial anomalies, as well as defects in neurogenesis and ear, nasal, and gland development. Heterozygotes may show variable hearing loss.

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





