

# **Brca1** Cas9-KO Strategy

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## **Project Overview**



**Project Name** 

Brca1

**Project type** 

Cas9-KO

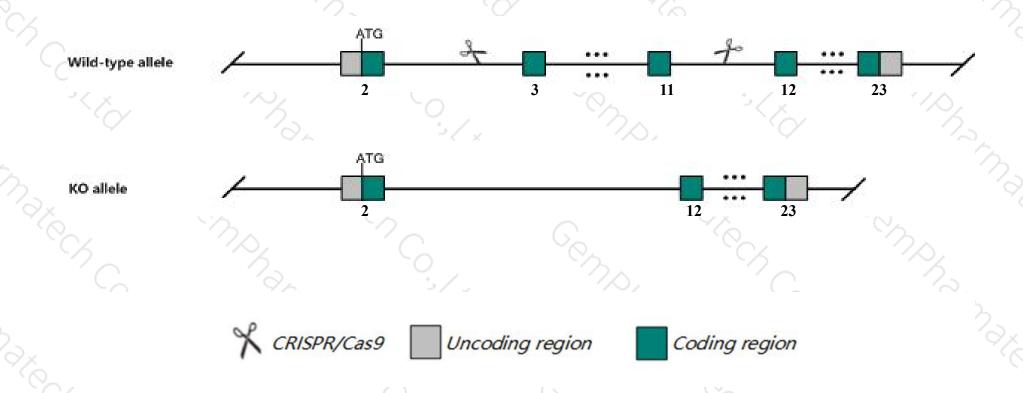
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Brca1* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Brca1* gene has 7 transcripts. According to the structure of *Brca1* gene, exon3-exon11 of *Brca1-201* (ENSMUST00000017290.10) transcript is recommended as the knockout region. The region contains 3973bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Brca1* gene. The brief process is as follows: CRISPR/Cas9 system

### **Notice**



- > According to the existing MGI data, Homozygous null mutants are embryonic lethal with abnormalities including growth retardation, neural tube defects, and mesoderm abnormalities; conditional mutations cause genetic instability and enhanced tumor formation; mutants with truncated BRCA1 protein survive, have a kinky tail, pigmentation anomalies, male infertility and increased tumor incidence.
- The *Brca1* 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.

### Gene information (NCBI)



#### Brca1 breast cancer 1, early onset [Mus musculus (house mouse)]

Gene ID: 12189, updated on 9-Apr-2019

#### Summary

☆ ?

Official Symbol Brca1 provided by MGI

Official Full Name breast cancer 1, early onset provided by MGI

Primary source MGI:MGI:104537

See related Ensembl:ENSMUSG00000017146

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

Expression Biased expression in liver E14 (RPKM 5.9), CNS E11.5 (RPKM 5.7) and 12 other tissuesSee more

Orthologs <u>human</u> all

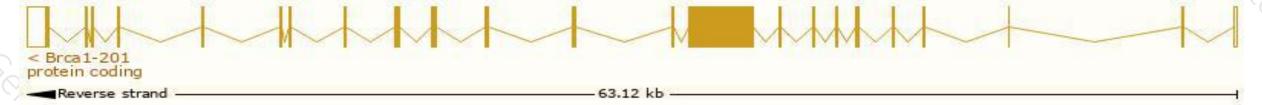
## Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

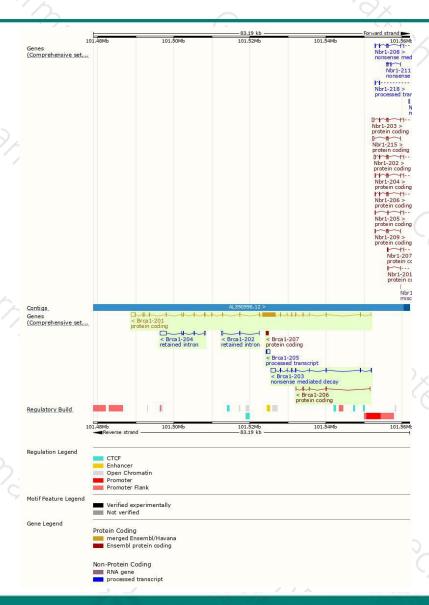
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Brca1-201	ENSMUST00000017290.10	6572	1812aa	Protein coding	CCDS25474	P48754	TSL:1 GENCODE basic APPRIS P1
Brca1-207	ENSMUST00000191198.1	531	<u>177aa</u>	Protein coding		A0A087WPE1	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete TSL:3
Brca1-206	ENSMUST00000190862.1	355	<u>79aa</u>	Protein coding	-	A0A087WP26	CDS 3' incomplete TSL:5
Brca1-203	ENSMUST00000142086.2	1917	<u>75aa</u>	Nonsense mediated decay	20	A0A087WPK5	TSL:1
Brca1-205	ENSMUST00000188168.1	852	No protein	Processed transcript	-	5	TSL:3
Brca1-204	ENSMUST00000156843.1	2136	No protein	Retained intron			TSL:1
Brca1-202	ENSMUST00000131460.1	539	No protein	Retained intron		20	TSL:3

The strategy is based on the design of *Brca1-201* transcript, The transcription is shown below



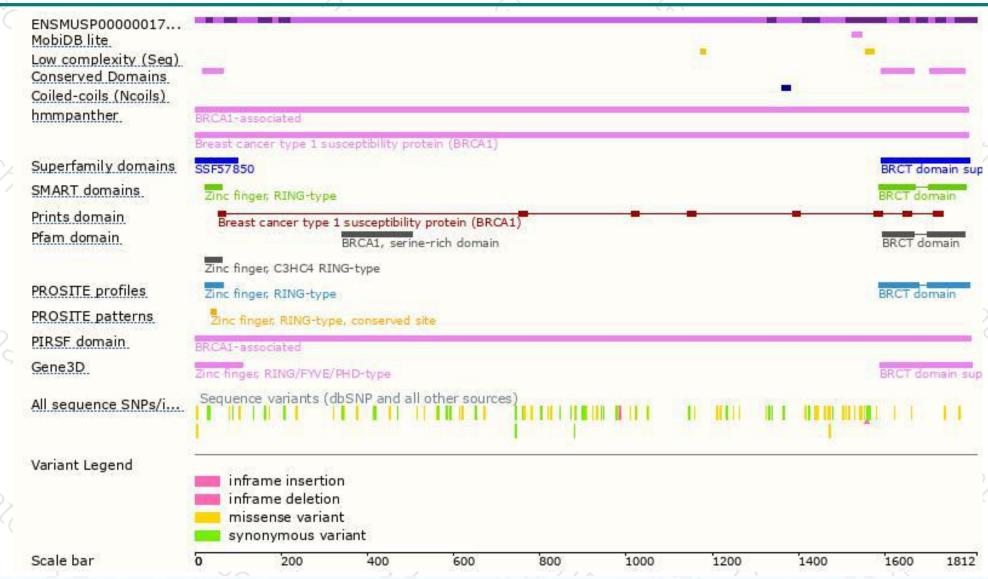
### Genomic location distribution





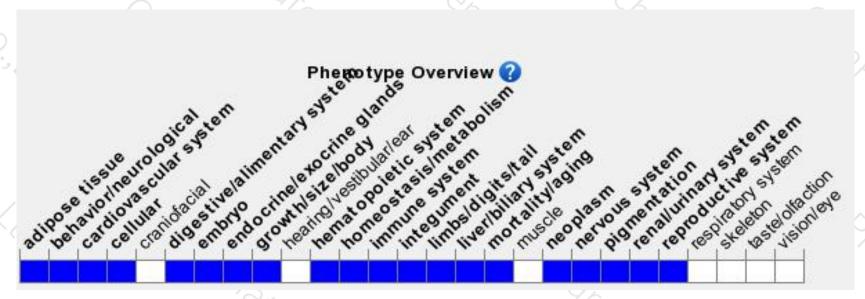
### Protein domain





### 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, Homozygous null mutants are embryonic lethal with abnormalities including growth retardation, neural tube defects, and mesoderm abnormalities; conditional mutations cause genetic instability and enhanced tumor formation; mutants with truncated BRCA1 protein survive, have a kinky tail, pigmentation anomalies, male infertility and increased tumor incidence.



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





