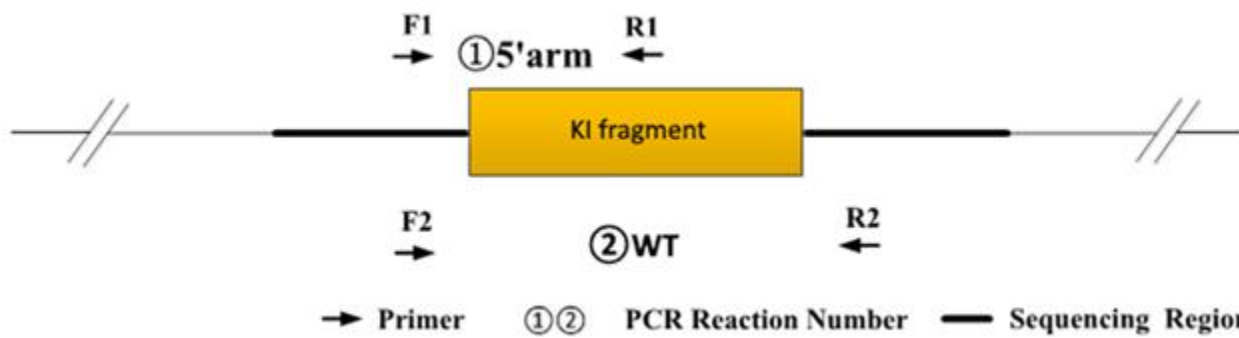


Genotyping Report

Strain ID	T056491	Strain Type	KI(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Tianjiao Wang	Gene Name	<i>CAG-LSL-mPreIp-flag-PolyA.H11-Myh6-CreERT2</i>		

1. Strategy of Genotyping



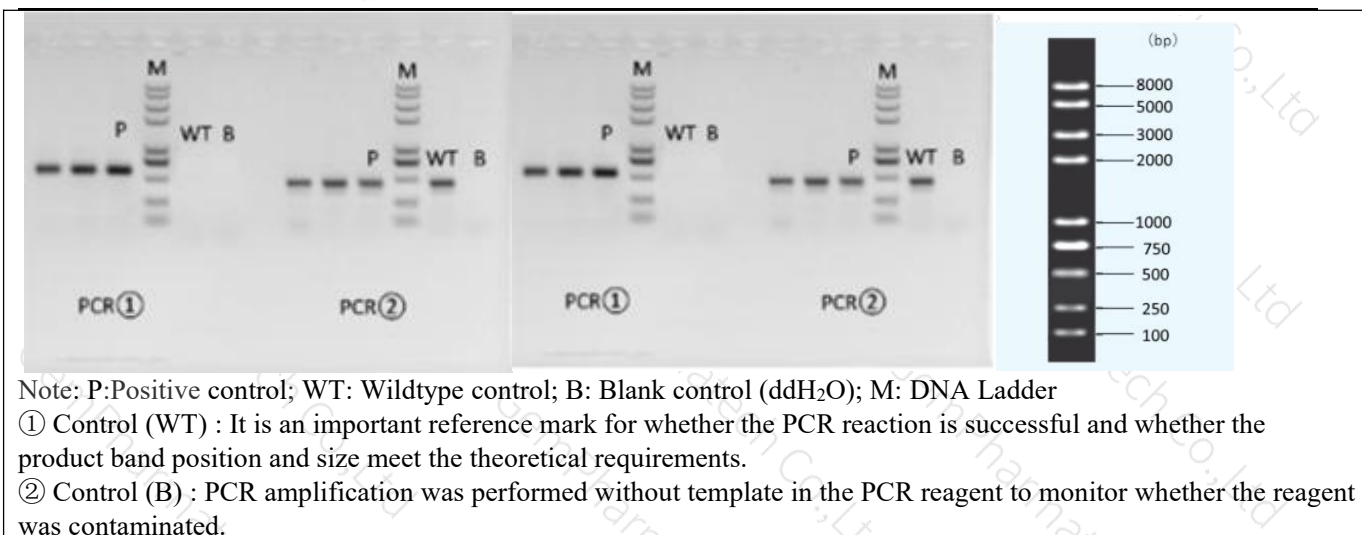
Wild type: ①PCR reaction obtains none band; ②PCR reaction obtains a WT band.
Heterozygote: ①PCR reaction obtains a Targeted band; ②PCR reaction obtains a WT band.
Homozygote: ①PCR reaction obtains a Targeted band; ②PCR reaction obtains none band.
Note: The sizes of WT and Targeted band are shown below. For ②PCR reaction, because the WT band is much smaller than the target band, it is likely to produce dominant amplification, the reaction is only used to judge whether there is a WT allele.

2. Primer Information

PCR No.	Primer No.	Sequence	Band Size
① 5'arm	T054788-F1	TTCTCCTCGGATCTGGAAAACG	KI:627bp
	T054788-R1	TCGGGTGAGCATGTCTTTAATCT	
② WT	T054788-F2	CCCAAAGTCGCTCTGAGTTGTTA	WT:479bp
	T054788-R2	TCGGGTGAGCATGTCTTTAATCT	
① 5'arm	T004858- F1	GGCACATGAGTAACAAAGGCATG	KI:553bp
	T004858-R1	AGCCAACCTTTGTTTCATGGCAG	
② WT	T004858- F2	CAGCAAAACCTGGCTGTGGATC	WT:412bp
	T004858-R2	ATGAGCCACCATGTGGGTGTC	

3. Gel Image & Conclusion

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4. PCR Condition

PCR Reaction Component			
Seg.	reaction component		Volume (μl)
1	2 × Rapid Taq Master Mix (Vazyme P222)		12.5
2	ddH2O		9.5
3	Primer A(10pmol/μl)		1
4	Primer B(10pmol/μl)		1
5	Template(≈100ng/μl)		1
PCR program ① priority selection			
Seg.	Temp.	Time	Cycle
1	95℃	5min	20×
2	98℃	30s	
3	65℃* (-0.5℃/cycle)	30s	
4	72℃	45s*	
5	98℃	30s	20×
6	55℃*	30s	
7	72℃	45s*	
8	72℃	5min	
9	10℃	hold	
PCR program ② the second choice			
Seg.	Temp.	Time	Cycle
1	95℃	5min	35×
2	98℃	30s	

3	58℃*	30s	
4	72℃	45s*	
5	72℃	5min	
6	10℃	hold	

Note*: Annealing temperature and extension time can be determined according to the actual amplification situation and amplification enzyme efficiency.