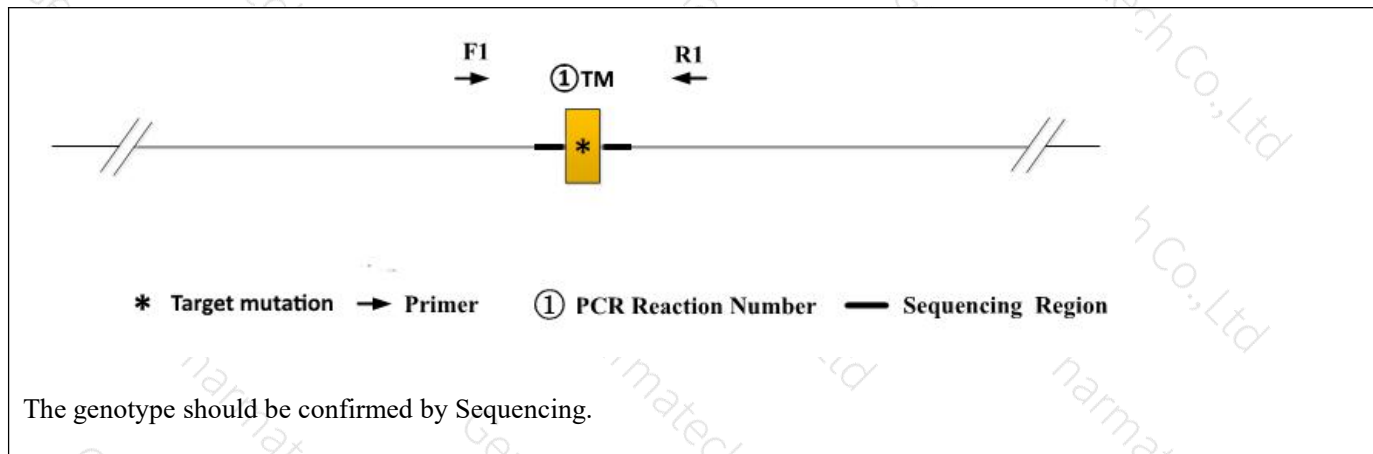


Genotyping Report

Strain ID	T057525	Strain Type	KI(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Tianjiao Wang	Gene Name	<i>Fbxo47-p.F384L</i>		

1. Strategy of Genotyping



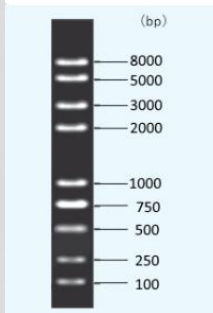
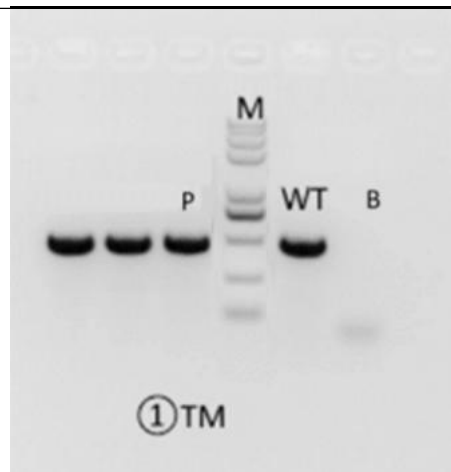
2. Primer Information

PCR No.	Primer No.	Primer Name	Sequence	Band Size
①TM	F1	T057525-F1	GCTATGCTATGTCACAATCCTGCC	WT:468bp Targeted:468bp
	R1	T057525-R1	TGGAAAGCAGAGGCAGTCAGTAGA	

Sequencing:

Primer No.	Sequence	
T057525-F1	GCTATGCTATGTCACAATCCTGCC	①TM

3. Gel Image



Note: P: samples; WT: Wildtype control; B: Blank control (ddH₂O); M:DNA Ladder

① Control (WT) : It is an important reference mark for whether the PCR reaction is successful and whether the product band position and size meet the theoretical requirements.

② Control (B) : PCR amplification was performed without template in the PCR reagent to monitor whether the reagent was contaminated.

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(20~80ng/μl)		1
PCR program I 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	15×
6	55℃*	30s	
7	72℃	45s*	
8	72℃	5min	
9	10℃	hold	
PCR program II the second choice			
Seg.	Temp.	Time	Cycle
1	95℃	5min	

2	98℃	30s	35×
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.