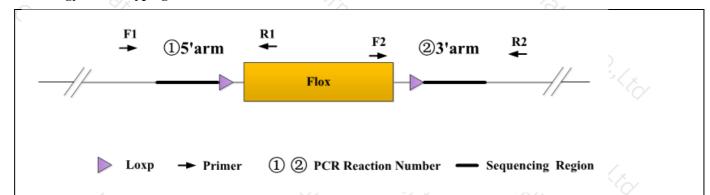
Genotyping Report

Strain ID	T014943	Strain Type	CKO(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Ying Chen	Gene Name	x	Ppp1r15a	C

1. Strategy of Genotyping



Wild type: ①PCR reaction obtains a single WT band; ②PCR reaction obtains a single WT band.

Heterozygote: ①PCR reaction obtains a WT band and a Targeted band; ②PCR reaction obtains a WT band and a Targeted band.

Homozygote: ①PCR reaction obtains a single Targeted band; ②PCR reaction obtains a single Targeted band.

Note: The sizes of WT and Targeted band are shown below.

2. Primer Information

PCR No.	Primer No.	Sequence	Band Size
①(5'arm)	T014943-F1	TGGGAAGGCTAGGAGAGAGCTTA	WT: 238bp Targeted: 343bp
	T014943-R1	GGCTCCACACCTCTCTTATAATGGGT	
	T014943-F2	ATTTATTCAAAGGGCCTAGATTCTGAG	WT: 229bp Targeted: 335bp
②(3'arm)	T014943-R2	CCCTCTGATTCACTCTAAGACACAGATG	

3. Gel Image & Conclusion



Note: P: Positive control; WT: Wildype 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

Seg.	rea	reaction component		
1		2 × Rapid Taq Master Mix (Vazyme P222)		
2	ddH2O	ddH2O		
3 %	Primer A(10pmol/μl)	6	1 %	
1 %	Primer B(10pmol/μl)	3	70,1	
5	Template(≈100ng/μl)	Template(≈100ng/μl)		
PCR program	$\widehat{\mathbb{D}}$ priority selection	777		
Seg.	Temp.	Time	Cycle	
1 72/	95℃	5min	77/36 53/4	
		- / · · · · · · · · · · · · · · · · · ·		
2	98℃	30s	20×	
3	98℃ 65℃* (-0.5℃/cycle)	30s 30s	20×	
	(0_	12	20x	
1 6	65°C* (-0.5°C/cycle)	30s	20× 20×	
1 C	65°C* (-0.5°C/cycle) 72°C	30s 45s*	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	
2 3 4 5 6	65°C* (-0.5°C/cycle) 72°C 98°C	30s 45s* 30s	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	
5 %	65°C* (-0.5°C/cycle) 72°C 98°C 55°C*	30s 45s* 30s 30s	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	



Seg.	Temp.	Time	Cycle
1 3/7	95℃	5min C	973 3/50
2 (98℃	30s	35×
3	58℃*	30s	3
4	72℃	45s*	3/
5 7 _{0×}	72℃	5min	(2) (A)
6	10℃	hold	~ C

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