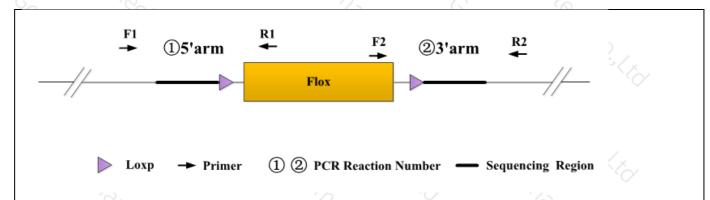


# **Genotyping Report**

Strain ID	T008476	Strain Type	CKO(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Tiantian Sun	Gene Name	-3-<->	bclaf1	

## 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.

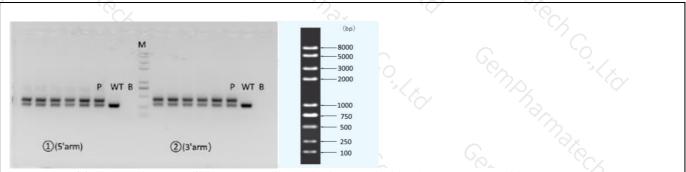
Homozygote: ①PCR reaction obtains a single Targeted band; ②PCR reaction obtains a single Targeted band.

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

#### 2. Primer Information

PCR No.	Primer No.	Sequence	Band Size	
①(5'arm)	T008476-F1	ATCCGCAGCACAGAGAATAGGAGC	WT: 278bp	
	T008476-R1	GATAAGCAATGAATGAGAGGCCCTG	Targeted: 383bp	
②(3'arm)	T008476-F2	AAAGTGAGTTCCAGGACAGCCAGG	WT: 314bp Targeted: 420bp	
	T008476-R2	TCCCTTGCCACCTCTCCTGCATA		

### 3. Gel Image & Conclusion



Note: P: Positive control; 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 C	omponent		30/x 3/x	
Seg.	reaction o	reaction component		
10	2 × Rapid Taq Master Mix(Vazyn	2 × Rapid Taq Master Mix (Vazyme P222)		
2 %	ddH2O	(%) (%)	9.5	
3 3/2	Primer A(10pmol/μl)	6	3/1 3/x	
4	Primer B(10pmol/μl)	Primer B(10pmol/μl)		
5	Template(≈100ng/μl)	Template(≈100ng/μl)		
PCR program (1	priority selection	3	· 6	
Seg.	Temp.	Time	Cycle	
1	95℃	5min		
2	98℃	30s	20×	
3	65℃*(-0.5℃/cycle)	30s	, (°C)	
4 %	<b>72℃</b>	45s*	79, 'G	
5	98℃	30s	20×	
6	55℃*	30s	79x	
7 %	72°C	45s*	.00	
8	<b>72</b> ℃	5min	8	
9	10℃	hold	7/2 3/6	
PCR program @	the second choice	7). C	9%	
Seg.	Temp.	Time	Cycle	
1	95℃	5min	79. 0.3/x	
2	98℃	30s	35×	
3	58℃*	30s	1°C	
4	72°C	45s*	· 'C	
5 S	<b>72℃</b>	5min	8/2 3/2 3/2 M	
6	10℃	hold	7%	

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