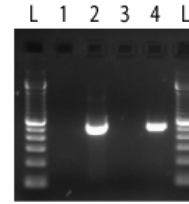


**pEMS1302 Promoterless construct PCR**- contains EGFP<sup>Cre</sup> reporters

**MiniPromoter:** none  
**pEMS#:** 1302  
**Expected product size (bp):** 544

<b>Reaction components</b>	<b>Vol/Rxn (µl)</b>
H <sub>2</sub> O	15.15
10X PCR buffer*	2.5
50 mM MgCl <sub>2</sub> *	0.75
2.5 mM dNTPs**	2
10 µM oEMS2364	1.25
10 µM oEMS2378	1.25
Taq Pol. (5 U/µl)*	0.1
DNA***	2
Total Volume of Rxn:	25



1- WT mouse DNA  
 2- Control plasmid DNA (pEMS1302)  
 3- No Template  
 4- Knock-in ESC line  
 L- 100 bp ladder  
 expected band size = 544 bp

\* Taq Polymerase set from Invitrogen (Cat no.18038-042)

\*\* dNTPs from Invitrogen (Cat no.10297-018)

\*\*\*Approximately 100 ng DNA used for samples, approximately 5 ng used for plasmid control

Samples run on a 2% agarose gel (containing SYBRsafe (Invitrogen Cat no. S33102))

<b>Cycling conditions:</b>	<b>Step</b>	<b>Temp</b>	<b>Time</b>	<b>Note</b>
	1	94°C	3 min	
	2	94°C	1 min	
	3	61°C	1 min	
	4	72°C	45 sec	reapeat steps 2-4 34 times
	5	72°C	5 min	
	6	4°C	hold	

**Primers:**

<b>Name</b>	<b>Sequence</b>	<b>T<sub>m</sub> (°C)</b>	<b>Notes</b>
<b>oEMS2364</b>	5'- GCGTATCACGAGGCCCTTTC -3'	56.0	Sense primer located in Vector backbone 5' to MCS
<b>oEMS2378</b>	5'- CTTCAGCTCGATGCGGTTCA -3'	56.2	Anti-sense primer located in EGFP sequence