

GENOTYPING BY PCR PROTOCOL FORM
MUTANT MOUSE REGIONAL RESOURCE CENTER: UC DAVIS

Investigator/PI: MMRRRC

Address: 2795 2nd street suite 400 Davis, CA 95618

Contact: Reneé Araiza

Telephone: 530-757-3284

FAX: 530-757-3284

email: mmrrc@ucdavis.edu

**PCR Protocol: STOCK Tg(Grp-cre)KH107Gsat MMRRRC:031182-UCD &
 STOCK Tg(Grp-cre)KH288Gsat MMRRRC:031183-UCD**

Reagent/ Constituent	Volume (µL)
Water	11.275
10x Buffer	2.5
25 mM MgCl ₂	1.7
5 M Betaine	6.5
10 mM dNTPs	0.5
DMSO	0.325
Primer 1: (20uM)	0.5
Primer 2: (20uM)	0.5
Taq Polymerase-5 Units/µl	0.2
DNA Sample	1.0
TOTAL VOLUME OF REACTION:	25 µl

Comments on protocol:

- Use Touch-Down cycling protocol-first 10 cycles anneal at 65° C decreasing in temperature by 1.0° C; next 30 cycles anneal at 55° C.
- Betaine/DMSO is standardized due to high GC content in promoter regions and protocol may be tested without. Also, may adjust MgCl₂ to increase reaction or decrease non specific amplifications.

Strategy:

Steps	Temp (°C)	Time (m:ss)	# of Cycles
1. Initiation/Melting	94	5:00	1
2. Denaturation	94	0:15	\
3. Annealing steps 2-4 will cycle in sequence	65 to 55 (↓1°C/cycle)	0:30	> x 40
4. Extension	72	0:40	/
5. Final Extension	72	5:00	1
6. Finish	4	Hold	--

Primers:

Primer Name	Nucleotide sequence (5'– 3')
1: Grp F	GGGACAACGCACTCTCAGCCTA
2: CreGS R1	CggCAAACggACAgAAGCATT

Electrophoresis Protocol:

% Agarose: 1.5 Volts : 90

Estimated Running Time (min): 90

Primer Combinations	Band size (bp)	genotype
1 & 2	230	transgenic

