

GENOTYPING BY PCR PROTOCOL
MUTANT MOUSE REGIONAL RESOURCE CENTER: UC DAVIS
 2795 2nd Street, Suite 400, Davis, CA 95618
mmrrc@ucdavis.edu
 530-754-MMRRC

NAME OF PCR: STOCK Tg(Neurod1-EGFP)1Gsai/Mmcd MMRRC # 000329-UCD

Protocol:

Reagent/ Constituent	Volume (μL)
Water	11.275
10x Buffer	2.5
MgCl ₂ (stock concentration is 25mM)	1.7
Betaine (stock concentration is 5M)	6.5
dNTPs (stock concentration is 10mM)	0.5
DMSO	0.325
Primer 1 (stock concentration is 10μM) Neurod1 (00329) F1	0.5
Primer 2 (stock concentration is 10μM) GFP R2	0.5
Taq Polymerase (5Units/μL)	0.2
DNA extracted with ☐ NaOH ☒ Proteinase K ☐ Other:	1.0
TOTAL VOLUME OF REACTION:	25μL

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. **Betaine and DMSO are absolutely required for this PCR to work. Also, may adjust MgCl₂ to increase reaction or decrease non specific amplifications.

Strategy:

Steps	Temp (°C)	Time (m:ss)	# of Cycles
1. Initiation/Melting HOT START? ☐	94	5:00	1
2. Denaturation	94	0:15	} 40x
3. Annealing	65 to 55 (↓1°C/cycle)	0:30	
4. Elongation		0:40	
5. Amplification	72	5:00	1
6. Finish	4	Hold	n/a

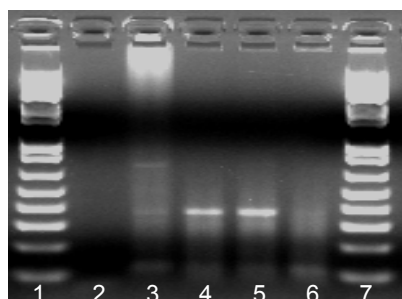
Primers:

Name	Nucleotide Sequence (5' - 3')
1: Neurod1 (00329) F1	TAG GAT TAG GGA GAG GGA GCT GAA
2: GFP R2	TAG CGG CTC AAG CAC TGC A

Electrophoresis Protocol:

Agarose: 1.5% V: 90
 Estimated Running Time: 90 min.

Primer Combination	Band	Genotype
1 and 2	380 bp	transgenic
Tg copy # ~ copies/genome		



Lanes
 1 & 7: 1 kb+ ladder
 (Invitrogen, Cat. #10787-026)
 2: H₂O
 3: WT Control
 4 & 5: Neurod1 tg/+
 6: Other GENSAT line