

GENOTYPING BY PCR PROTOCOL FORM

MUTANT MOUSE REGIONAL RESOURCE CENTER: UC DAVIS

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DNA Extraction Method: NaOH _____ Proteinase K X Other _____

Protocol: NAME OF PCR: C57BL/6J-Slfn2^{m1Btlr}/Mmcd (*elektra*), MMRRC #031705-UCD

Reagent/ Constituent	Volume (uL)
DNA Sample	1
10x Buffer (contains 15mM MgCl ₂)	5
dNTPs (stock concentration is 25mM)	0.4
Primer 1 (stock concentration is 20 uM)	1
Primer 2 (stock concentration is 20 uM)	1
Taq Polymerase	2.5
water	39.1
TOTAL VOLUME OF REACTION:	
	50 ul

Comments on protocol (e.g., different concentration of MgCl₂, etc): *Elektra* genotyping is performed by amplifying the region containing the mutation using PCR, followed by sequencing of the amplified region to detect the single nucleotide transversion. Use primers 1 & 2 for amplification and 3 & 4 for sequencing.

Strategy:

Steps	Temp (°C)	Time (min)	# of Cycles
1. Initiation/Melting HOT START?..CHECK HERE []	94	2	1
2. Denaturation	94	30 seconds	
3. Annealing } steps 2-3-4 will cycle in sequence	56	30 seconds	30
4. Elongation	72	1 min	
5. Amplification (i.e., 72°C, 10 min)	72	7	1
6. Finish (i.e., 4°C, indefinite)	4	n/a	n/a

Primers:

Primer Name	Nucleotide Sequence (5' - 3')
1: Slfn2_elektra_F	GCTGGGTCTAAATCTCGGAGCAATC
2: Slfn2_elektra_R	TTCCCGCCAAATCATCCTGGAGTC
3: Slfn2_elektra_seqF	AATTCGGGAGGTGGAGCAATC
4: Slfn2_elektra_seqR	TTCTCCTCACAGAAGTGAGTGAC

Electrophoresis Protocol:

% Agarose: _____ V : _____

Estimated Running Time (min): _____

Primer combination	Band (bp)	genotype
(i.e. 1&2)	1310	
(i.e. 3&4)		
(i.e. 1&2&3)		

Mutation site (red) and flanking sequence:

WT acatttata**t**ctacgtgaa
elektra acatttata**a**ctacgtgaa