

GENOTYPING BY PCR PROTOCOL

KOMP Repository: UC DAVIS

Protocol:CR804 Ano3

Reagent/Constituent	Volume (μ L)
Water	10.275
10x Buffer	2.5
MgCl ₂ (stock concentration is 25mM)	1.7
Betaine (stock concentration is 5M) <i>Optional</i>	6.5
dNTPs (stock concentration is 10mM)	0.5
DMSO <i>Optional</i>	0.325
Primer 1. (stock concentration is 20 μ M)	0.5
Primer 2. (stock concentration is 20 μ M)	0.5
Primer 3. (stock concentration is 20 μ M)	0.5
Primer 4. (stock concentration is 20 μ M)	0.5
Taq Polymerase 5Units/ μ L	0.2
DNA (example) extracted w/ "Qiagen DNeasy columns or other similar silica based kits"	1.0
TOTAL VOLUME OF REACTION:	25.000 μL

Comments on protocol:

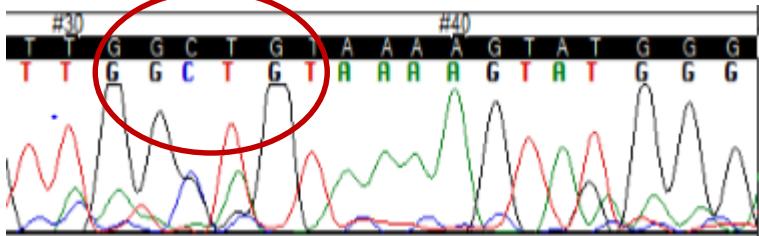
- Protocol may work with other DNA extraction methods.
- Use Touch-Down cycling protocol-first 10 cycles anneal at 65°C decreasing in temperature by 1.0°C; next 30 cycles anneal at 55°C.
- Betaine and DMSO have been standardized due to high GC content. Protocol may be tested without. Also, may adjust MgCl₂ to increase reaction or decrease non-specific amplifications.

Strategy:

Steps	Temp (°C)	Time (m:ss)	# of Cycles
1. Initiation/Melting HOT START? <input type="checkbox"/>	94	5:00	1
2. Denaturation	94	0:15	
3. Annealing steps 2-3-4 cycle in sequence	65 to 55 (↓1°C/cycle)	0:30	40x
4. Elongation	72	0:40	
5. Amplification	72	5:00	1
6. Finish	15	∞	n/a

Primers:

Electrophoresis Protocol			Frameshift Indel		
Name	Nucleotide Sequence (5' - 3')	Agarose: 1.5% V: 90	Primer Combination	Band (bp)	Genotype
1. Ano3_CR804-F	GTTCCAAGTCTAATGATGCCCTCTCC	Estimated Running Time: 90 min			
2. Ano3_CR804-R	TGTTCAGCCTCTCTGCATACTTGC		1 & 2	123	wildtype
			1 & 2	113	mutant



NTC	B:4
WT	B:5
CR804-624	B:6
CR804-625	B:7
CR804-626	B:8
CR804-627	B:9
CR804-628	B:10
CR804-629	B:11

Frameshift Indel 10bp Deletion

Contig atactttaca**gccagcaatt**gccaa

Allele Description: Exon 6 (ENSMUSE00001272495) received a 10bp deletion (**gccagcaatt**) from the Ano3 gene (ENSMUST00000099623.9) using CRISPR Cas9 gene editing technology in mouse zygotes. This causes a frameshifted transcript followed by early termination signal. Subsequent founders were backcrossed to C57BL/6N to produce sequence confirmed heterozygous animals.

