

GENOTYPING BY PCR PROTOCOL

KOMP Repository: UC DAVIS

Protocol: CR985 Ces1d

| 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 |

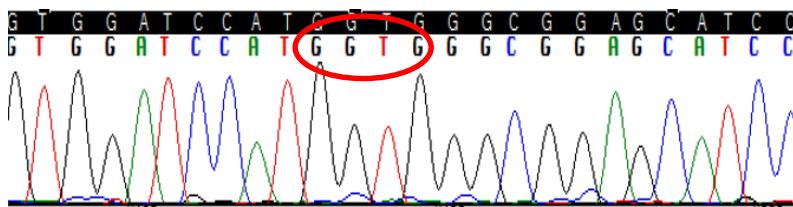
Primers:

| Name | Nucleotide Sequence (5' - 3') | Agarose: 1.5% V: 90 |
|------------------|-------------------------------|---------------------------------------|
| 1. Ces1d CR985-F | GGTTATGTTCTTGTCCAGGTGATGG | Estimated Running Time: 90 min |
| 2. Ces1d CR985-R | GTCACCACCAACCATTTCATGG | Primer Combination Band (bp) Genotype |
| | | 1 & 2 116 wildtype |
| | | 1 & 2 102 mutant |

Frameshift Indel

Ces1d Exon 4

14bp Deletion



Contig gtggatccatgg**aggtggactggtggtgg**tgggcggagcatcc

Allele Description: Exon 4 (ENSMUSE00001059198) received a 14bp deletion (**aggtggactggtggtgg**) from the Ces1d gene (ENSMUST00000034172.7) 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.

