

**EMMA ID:** 01784

**Gene:** *Csrp2*

**Common name:** *CSR2 ko*

**Allele:** *Csrp2*<sup>tm1Rawe</sup>

## Genotyping Information

Genotyping by end-point PCR based on gel is composed of a genespecific short range PCR using primers on wild type allele and a mutant allele-specific short range PCR. The combined results show the genotype of the mice. For example: mutant positive, wild type positive = Heterozygous.

### PCR primer pairs and expected size bands

| Assay    | Forward Primer | Reverse Primer | Expected Size Band (bp) |
|----------|----------------|----------------|-------------------------|
| Wildtype | 31             | 30             | 218                     |
| Mutant   | 31             | Neo            | 422                     |

### Primer sequences

| Primer Name | Sequence 5' --> 3'                  |
|-------------|-------------------------------------|
| 31          | 5' CTA CCT TCC CAG CTC CAA TGA TC 3 |
| 30          | 5' CAG CAG TAG AGC TCC GAA GCT CC   |
| Neo         | 5'CTGCTCTTACTGAAGGCTCTT3'           |

### PCR setup (Qiagen, Hot Start Plus)

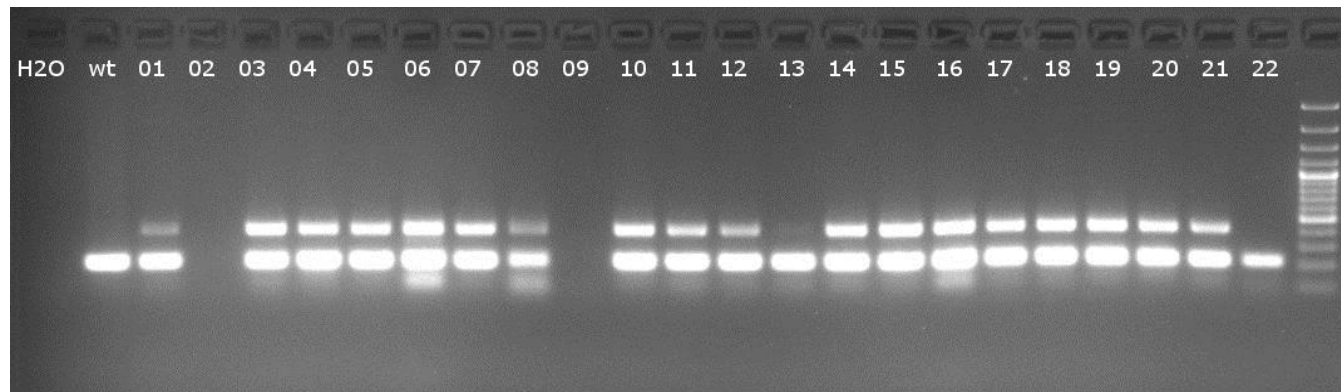
| Component         | Volume (µl) 1x | Final conc. |
|-------------------|----------------|-------------|
| DNA (~ 50-100 ng) | 2              |             |
| Q-Solution (5x)   | 2,5            | 0,5         |
| PCR-Buffer (10x)  | 2,5            | 1           |

### Amplification conditions

| PCR Settings                                       | Temperature (°C) | Time   | # of cycles |
|--|------------------|--------|-------------|
| 1 Denaturation<br>(Melting)                        | 95°C             | 5 min  | 1           |
| 2 Amplification<br>(Melting, Annealing,<br>Polym.) | 94°C             | 30 sec | 39          |
|  | 50°C             | 45 sec |             |
|  | 72°C             | 45 sec |             |
| 3 Polymerisation                                   | 72°C             | 10 min | 1           |
| 4 Cooling  | 12°C             | hold   | 1           |

These PCR conditions have been optimized for our methods and preparation kits. Adaptions may be required.

### Gel Image



Separated by gel electrophoresis on a 2% agarose gel.