Table S1 More detailed information about the PCR steps.

|  |  |  |
| --- | --- | --- |
| First PCR |  | Second PCR |
| 5xBuffer | 10μL |  | 5xBuffer | 8μL |
| dNTP (10mM) | 1μL |  | dNTP (10mM) | 1μL |
| DNA polymerase | 1U |  | DNA polymerase | 0.8 U |
| F/R Inner primer(10 uM) | Each 1μL |  | F/R Outer primer(10 uM) | Each 1μL |
| Template | 5ng-50ng |  | Template | 5uL |
| ddH2O | fill up to 50 μL |  | ddH2O | fill up to e0 μL |
| First PCR system |  | Second PCR system |
| PCR instrument(Applied Biosystems 9700 , USA) |  | PCR instrument(Applied Biosystems 9700 , USA) |
|  | 94℃ | 2min |  |  | 94℃ | 2min |
| 33 cycles | 94℃ | 30s |  | 8 cycles | 94℃ | 30s |
| 55℃ | 30s |  | 56℃ | 30s |
| 72℃ | 30s |  | 72℃ | 30s |
|  | 72℃ | 5min |  |  | 72℃ | 5min |
|  | 10℃ | heat preservation |  |  | 10℃ | heat preservation |

The electrophoresis (1.5% agarose gel in 0.5\*TBE) and gel extraction kit (Axygen Biosciences, USA) can separate and purify the PCR products. F/R Inner primer: 5'-TTCCCTACACGACGCTCTTCCGATCT3';5'-GAGTTCCTTGGCACCCGAGAATTCCA3'. F/R Outer primer: 5'-AATGATACGGCGACCACCGAGATCTACACTCTTTCCCTACACGACGCTC -3';5'-CAAGCAGAAGACGGCATACGAGATGTGACTGGAGTTCCTTGGCACCCGAGA-3.