

Supplementary data

Table S1. PCR mixture for 1 reaction

| Component | Volume (μ L) |
|---|-------------------|
| 10x <i>Taq</i> DNA polymerase buffer with KCl | 1.25 |
| 25 mM MgCl ₂ | 0.75 |
| 10 mM dNTPs | 0.50 |
| 10 μ M forward primer | 0.25 |
| 10 μ M reverse primer | 0.25 |
| Nuclease free water | 9.40 |
| Total | 12.5 |

Table S2. PCR thermal cycles

| Step | Temperature (°C) # | Time |
|----------------------|--------------------|----------------|
| Initial denaturation | 95 | 10 min |
| 10 cycles of | | |
| 1. Denaturation | 95 | 30 s |
| 2. Annealing | 55 | 30 s |
| 3. Extension | 72 | 1 min per 1 kb |
| Final extension | 72 | 5 min |

Table S3. Oligonucleotide primer sequences used in this study

| Primer | Sequence (5'-3') | Annealing temperature (°C) |
|----------------------|----------------------------|----------------------------|
| T7 promotor | TAATACGACTCACTATAGGG | 55 |
| T7 terminator | GCTAGTTATTGCTCAGCGG | 55 |
| R1 | CCATGATTACGCCAAGCTTGGAGCC | 55 |
| R2 | GCTAGATTCAAAACAGCAGAAAGG | 55 |
| SARS-CoV-2 E forward | ACAGGTACGTTAATAGTTAATAGCGT | 60 |
| SARS-CoV-2 E reverse | ATATTGCAGCAGTACGCACACA | 60 |