

Supplementary information

Table S1. Sequences of gene E and N of SARS-CoV-2 used as positive controls.

Gene E target sequence	Gene N target sequence
TGAGTACGAACTTATGTACTCATTCGTTT	ACCAGGAACTAATCAGACAAGGAACT
CGGAAGAGACAGGTACGTTAATAGTTAA	GATTACAAACATTGGCCGCAAATTGC
TAGCGTACTTCTTTTTCTTGCTTTCGTGGT	ACAATTTGCCCCCAGCGCTTCAGCGTT
ATTCTTGCTAGTTACACTAGCCATCCTTA	CTTCGGAATGTCGCGCATTGGCATGGA
CTGCGCTTCGATTGTGTGCGTACTGCTGC	AGTCACACCTTCGGGAACGTGGTTGAC
AATATTGTTAACGTGAGTCTTGTAACC	CTACACAGGTGCCATCAAATTGGATG
TTCTTTTACGTTTACTCTCGTGTTAAAAA	ACAAAGATCCAAATTTCAAAGATCAA
TCTGAA	GTC

Table S2. LAMP primer sequences.

Gene E	Sequence
E1-F3	TGAGTACGAACTTATGTACTCAT
E1-B3	TTCAGATTTTAAACACGAGAGT
E1-FIP	ACCACGAAAGCAAGAAAAAGAAGTTCGTTTCGGAAGAGACAG
E1-BIP	TTGCTAGTTACACTAGCCATCCTTAGGTTTACAAGACTCACGT
E1-LF	CGCTATTAACATTAACG
E1-LB	GCGCTTCGATTGTGTGCGT
Gene N	Sequence
N2-F3	ACCAGGAACTAATCAGACAAG
N2-B3	GACTTGATCTTTGAAATTTGGATCT
N2-FIP	TTCCGAAGAACGCTGAAGCGGAACTGATTACAAACATTGGCC

N2-BIP	CGCATTGGCATGGAAGTCACAATTTGATGGCACCTGTGTA
N2-LF	GGGGGCAAATTGTGCAATTTG
N2-LB	CTTCGGGAACGTGGTTGACC
Hs_rActin	Sequence
ACT-F3	AGTACCCCATCGAGCACG
ACT-B3	AGCCTGGATAGCAACGTACA
ACT-FIP	GAGCCACACGCAGCTCATTGTATCACCAACTGGGACGACA
ACT-BIP	CTGAACCCCAAGGCCAACCGGCTGGGGTGTGAAGGTC
ACT-LF	TGTGGTGCCAGATTTTCTCCA
ACT-LB	CGAGAAGATGACCCAGATCATGT

Table S3. Reproducibility data. The reproducibility test was performed with a serial dilution of each positive control (pUC17_N and pUC17_E) at a final concentration of 10^3 (level 1), 10^4 (level 2) and 10^5 (level 3) copies per μL , each.

Run data		Level 1 - 10^3	Level 2 - 10^4	Level 3 - 10^5
Operator and date	Positive results	Replicates	Replicates	Replicates
Operator 1 - Dec 10, 2020	100%	10/10	10/10	10/10
Operator 2 - Dec 11, 2020				
Operator 3 - Dec 14, 2020				
Operator 1 - Dec 10, 2020	100%	10/10	10/10	10/10
Operator 2 - Dec 11, 2020				
Operator 3 - Dec 14, 2020				

Operator 1 - Dec 10, 2020					
Operator 2 - Dec 11, 2020	100%	10/10	10/10	10/10	
Operator 3 - Dec 14, 2020					

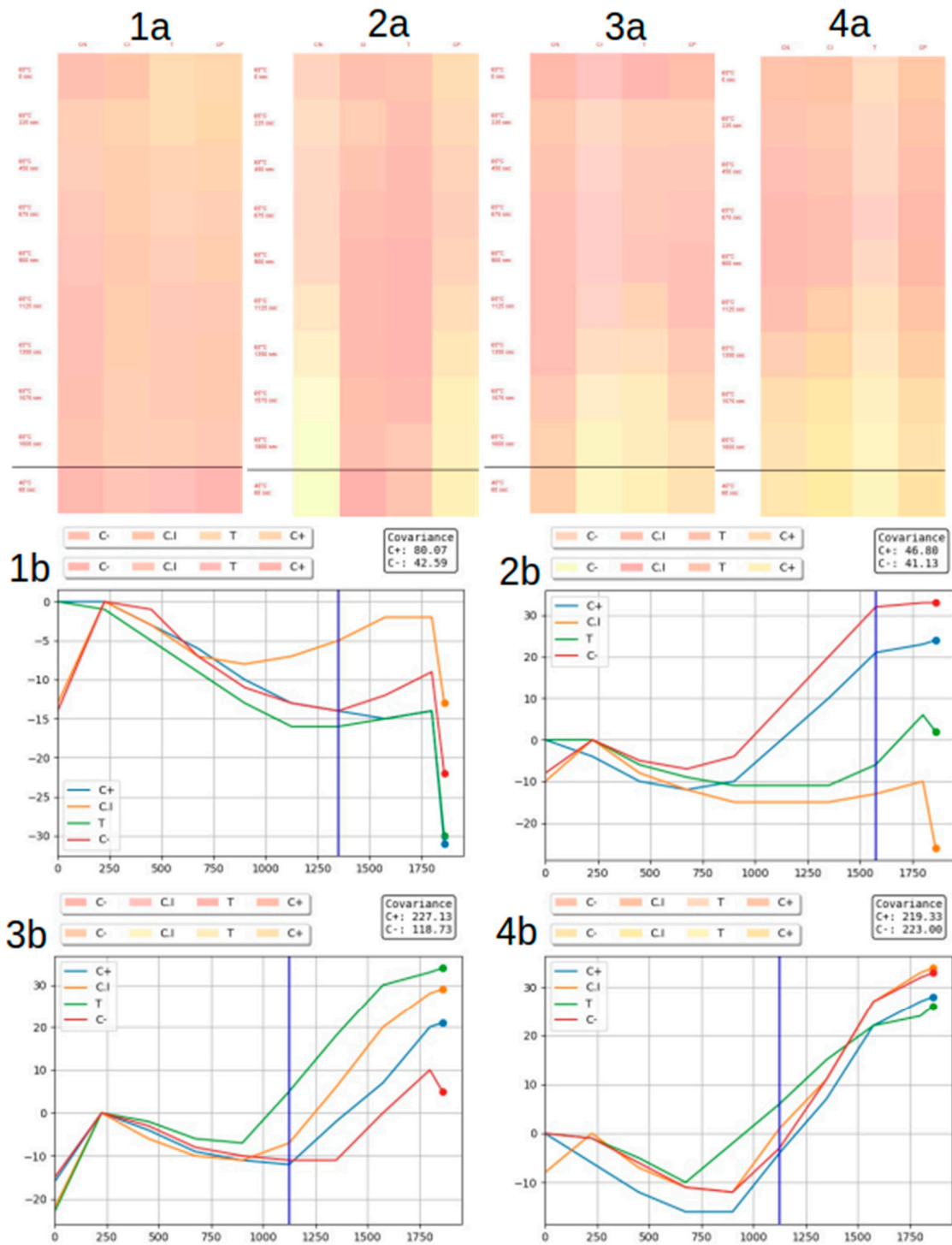


Figure S1. Software images obtained from the Limit of Detection (LoD) test. Figure 1a-b represents the non-template control. Figure 2a-b represents the 2.44×10^2 and 1.22×10^2 (left to right) genomic copies per reaction. Figure 3a-b represents the 9.77×10^2 and 4.88×10^2 (left to right) genomic copies per reaction. Figure 4a-b represents the 1.95×10^3 genomic copies per reaction.

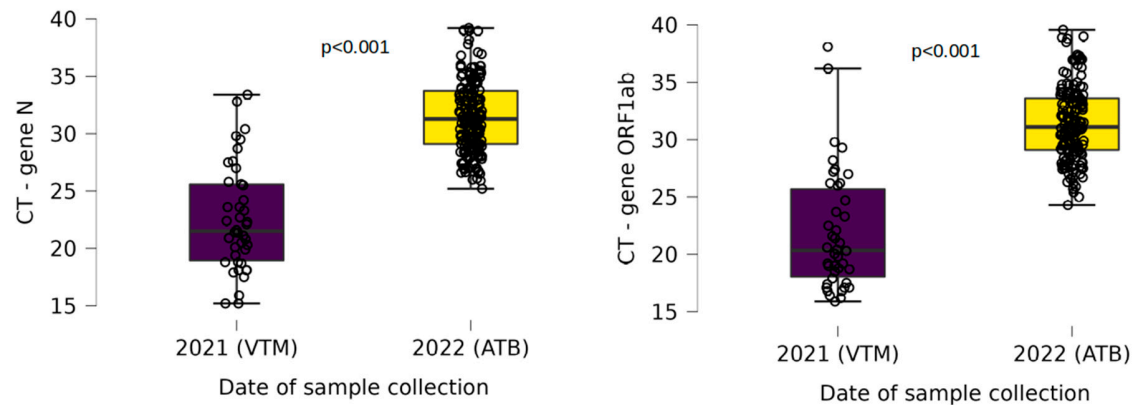


Figure S2. RT-qPCR CTs for samples collected in viral transport media (VTM) or antigen-test buffer (ATB). CTs for the N gene are shown in the left image, and the ORF1ab gene is in the right image. For both genes, we saw a statistically significant difference between sample solutions ($p < 0.001$).



Figure S3. Omicron variant mutation at E gene (T9I). In purple is the sequence of our FIP primer targeting the E gene, demonstrating that the mutation of the Omicron variant affected the primer annealing.