

Supplementary information

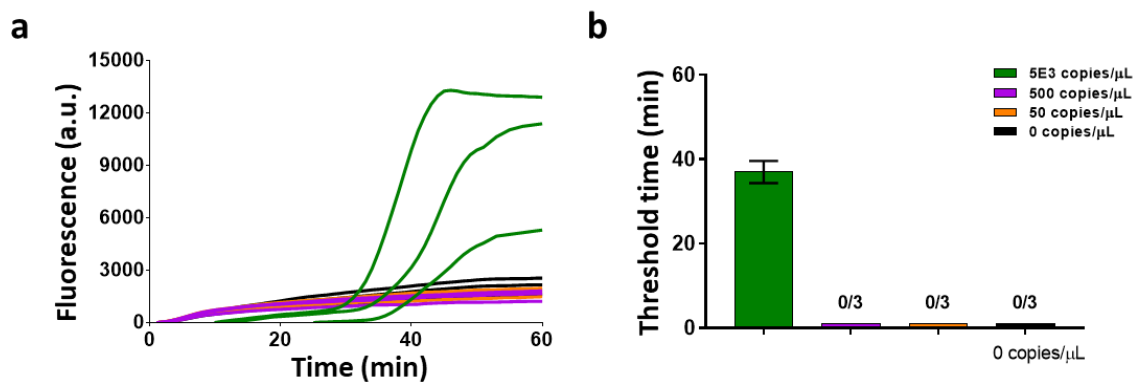


Fig. S1. Characterization of SARS-CoV-2 virus in nasal fluid in a 96 μL reaction. (a-b) Raw fluorescence data and amplification threshold times (for 3 replicates of data) for viral detection in a 96 μL reaction with 50% nasal fluid per reaction. Thermal lysis at 95°C was conducted for 1 min. of the virus in nasal fluid sample before addition of RT-LAMP reagents for the final reaction. The bar graphs show mean and standard deviation. Fraction indicates number of replicates amplified.

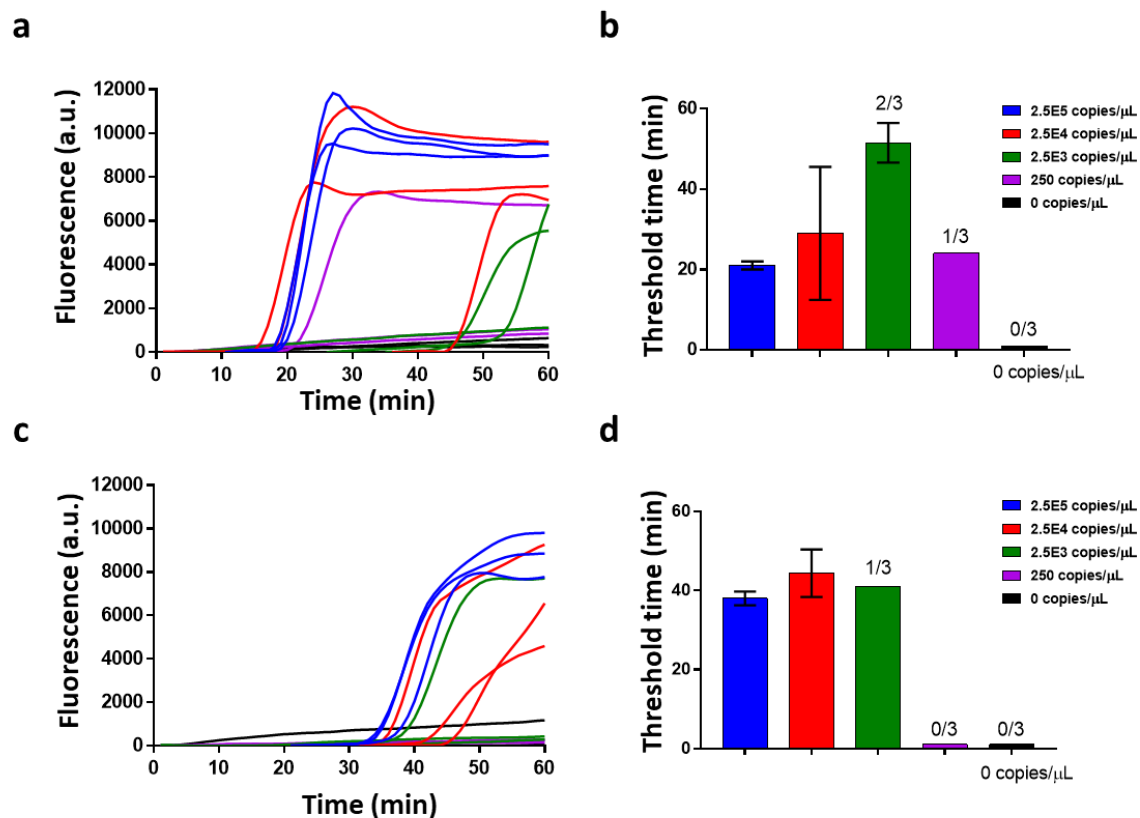
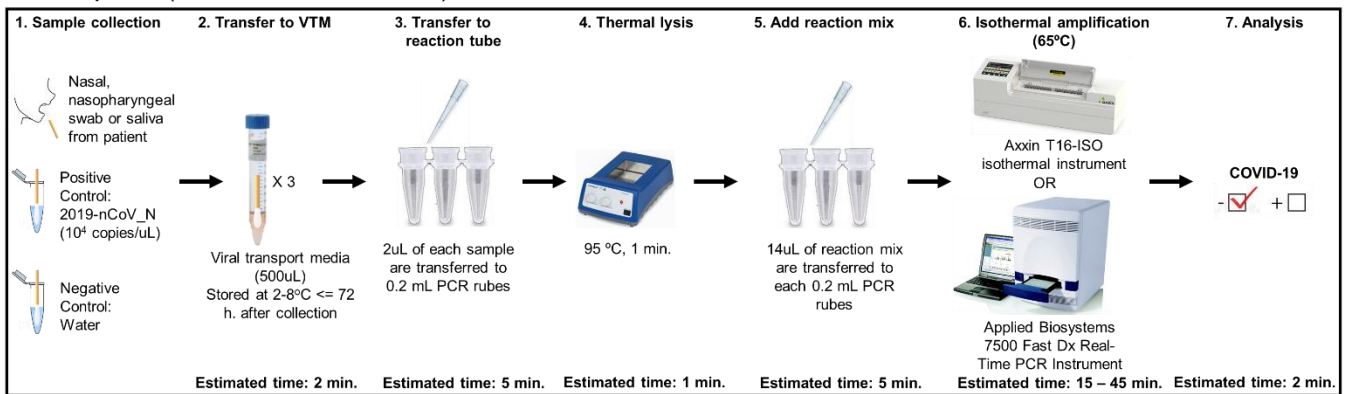


Fig. S2. Characterization of SARS-CoV-2 virus in mock swab samples transported in 100 μ L Viral Transport Media. (a-b) Raw fluorescence data and amplification threshold times (for 3 replicates of data) for viral detection in a 16 μ L reaction with 12.5% VTM per reaction from a 100 μ L VTM sample. **(c-d)** Raw fluorescence data and amplification threshold times (for 3 replicates of data) for viral detection in a 16 μ L reaction with 12.5% VTM per reaction from a 100 μ L VTM sample. The bar graphs show mean and standard deviation.

RT-LAMP protocol (estimated total time: 30 - 60 min.)



RT-PCR protocol (estimated total time: 119 min.)

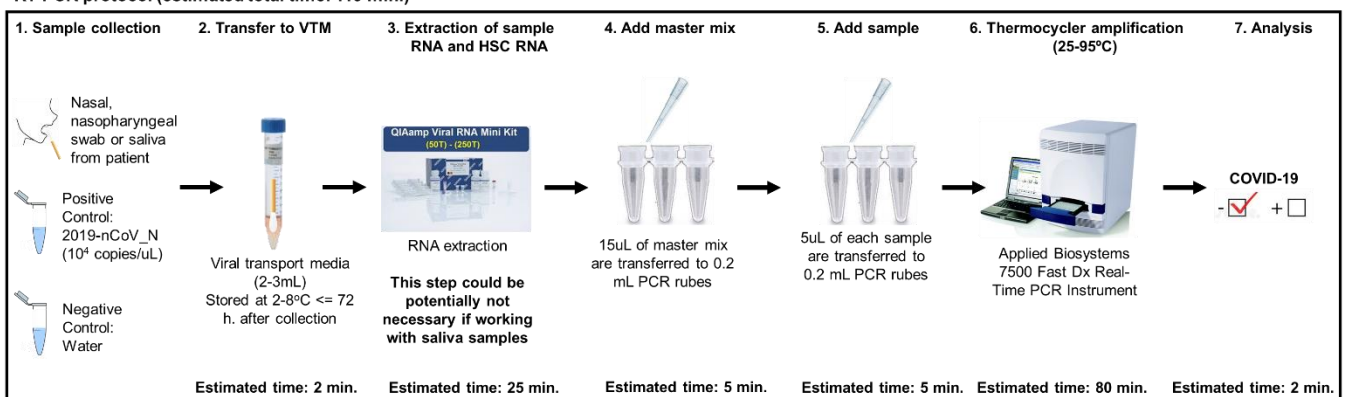


Fig. S3. RT-LAMP assay process flow and comparison side by side with the RT-PCR assay process flow.

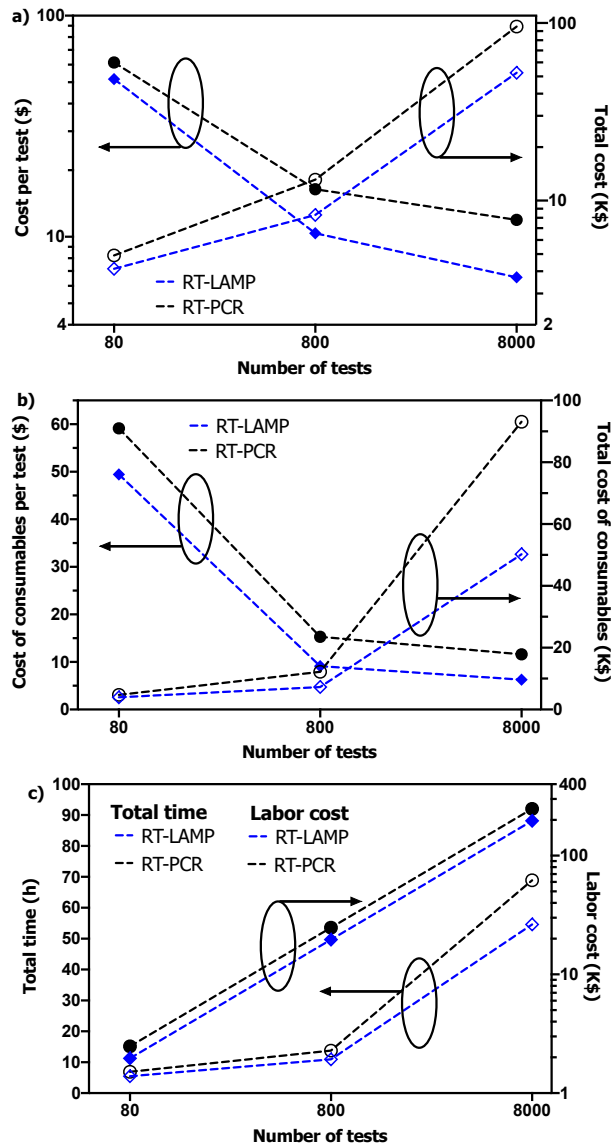


Fig. S4: Resources modeling results. a) Cost per test and total cost (excluding labor) for the three scenarios considered; b) Total cost of consumables and Cost of consumables per test for the 3 models designed; c) Labor cost and Total time for the 3 models designed. The labor cost assumes a gross salary rate = \$1/min.

Table S1. RT-LAMP primer sequences for 12 primer sets (3 primer sets each for 4 target genes)

Gene Orf1a	P1	F3	ATCTAGTTGTAATGGCCTACA	
		B3	ACAAGCACAGGTTGAGAT	
		FIP	TGAGTTTTTCATAAACAGTGCCAAA-CAGGTGGTGTGTTTCAGT	
		BIP	AACCCGTCCTTGATTGGCTT-AATTTAACAATTTCCCAACCGTC	
		Loop F	TGTTAGTTAGCCACTGCGAAGT	
	P2	F3	AAACCCGTCCTTGATTGG	
		B3	CTTTAAGTTTAGCTCCACCAAT	
		FIP	TCACAAGCACAGGTTGAGATAAATT-GAAGGTGTAGAGTTTCTTAGAGAC	
		BIP	GTCACCTGTGCAAAGGAAATTAAG-AGAGTCAGCACACAAAGC	
	P3	F3	CGGTGGACAAATTGTCAC	
		B3	CTTCTCTGGATTTAACACACTT	
		FIP	TCAGCACACAAAGCCAAAAATTTAT-CTGTGCAAAGGAAATTAAGGAG	
		BIP	TATTGGTGGAGCTAAACTTAAAGCC-CTGTACAATCCCTTTGAGTG	
	Gene S	P1	F3	TCTTTCACACGTGGTGTT
			B3	CAGTGGAAGCAAATAAACAC
FIP			GAAAGGTAAGAACAAGTCCTGAGT-TTACCCTGACAAAGTTTTTCAG	
BIP			TTCCAATGTTACTTGGTTCCATGC-GACAGGGTTATCAAACCTCT	
Loop B			TATACATGTCTCTGGGACCAATGG	
P2		F3	GGTGTATTATTACCCTGACAAAG	
		B3	GTACCAAAAATCCAGCCTC	
		FIP	TGGAACCAAGTAACATTGGAAAAGA-TTTTCAGATCCTCAGTTTTACATTC	
		BIP	CTCTGGGACCAATGGTACTAAGAG-GACTTCTCAGTGGAAGCA	
		Loop B	AACCCTGTCCTACCATTTAATGATG	
P3		F3	CCTGACAAAGTTTTTCAGATCC	
		B3	GTACCAAAAATCCAGCCTC	
		FIP	GCATGGAACCAAGTAACATTGGAAA-TCAGTTTTACATTCAACTCAGGA	
		BIP	CTCTGGGACCAATGGTACTAAGAG-GACTTCTCAGTGGAAGCA	
		Loop B	AACCCTGTCCTACCATTTAATGATG	
Gene Orf 8	P1	F3	ACGCCTAAACGAACATGAA	
		B3	AGAACCAGCCTCATCCAG	

		FIP	GGTTGATGTTGAGTACATGACTGTA- CTTGTTTTCTTAGGAATCATCACA
		BIP	ATATGTAGTTGATGACCCGTGTCC- TAAAGGTGCTGATTTTCTAGCT
		Loop F	CTACATTCTTGGTGAAATGCAGCTA
	P2	F3	CTCAACATCAACCATATGTAGT
		B3	CAATTTAGGTTCTCGGCAATT
		FIP	GGTGCTGATTTTCTAGCTCCTACTC- GACCCGTGTCCTATTCAC
		BIP	TGCTGGATGAGGCTGGTTCTA- TGTAAGGTAACAGGAAACTG
	Loop B	ATCACCCATTCAGTACATCGATATC	
	P3	F3	AGCTGCATTTCAACCAAGAA
		B3	CGATATCGATGTAAGTGAATGG
		FIP	TGAATAGGACACGGGTCATCA- GTAGTTTACAGTCATGTAAGTCAA
		BIP	GAGTAGGAGCTAGAAAATCAGCAC- TGATTTAGAACCAGCCTCATC
Gene N	P1	F3	GTTCCCTCATCACGTAGTCG
		B3	GTTTGGCCTTGTTGTTGTT
		FIP	GCCAGCCATTCTAGCAGGAG- CAACAGTTAAGAAATTCAACTCC
		BIP	GATGCTGCTCTTGCTTTGCT- ACCAGACATTTTCTCTCAA
		Loop B	GCTGCTTGACAGATTGAACCAG
	P2	F3	AGACGAATTCGTGGTGGT
		B3	TTGTTAGCAGGATTGCGG
		FIP	TGGCCAGTTCCTAGGTAGT- GACGGTAAAATGAAAGATCTCAG
		BIP	CTTCCCTATGGTGCTAACAAGAC- TGGTGTATTCAAGGCTCC
		Loop B	GGCATCATATGGGTTGCAACTGAG
	P3	F3	GTCATTTGCTGAATAAGCATAT
		B3	GAGTCAGCACTGCTCATG
		FIP	TAAGGCTTGAGTTTCATCAGCCTT- ACGCATACAAAACATTCCCA
		BIP	CAGAGACAGAAGAAACAGCAAAC- GATTGTTGCAATTGTTGGAG
		Loop B	GTGACTCTTCTTCTGCTGCAGATT

Table S2. Supply chain: RT-LAMP and RT-PCR assay

See excel file.

Video S1. Uniform filling of the amplification chambers.

See video

Video S2. Time stamped videos of amplification on the cartridge for 5000 copies/ μ L of virus in VTM.

See video

Video S3. Time stamped videos of amplification on the cartridge for negative control (VTM only).

See video