



**Certificate  
of Analysis**

Verified by:  
Jessie Ho

Quality Release  
Date: June 17, 2020

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**Product Name:            Coronavirus Environmental Test Kit**

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Catalog Num:	T1401S	<b>Chai Inc.</b>
Lot Num:	9787882	Santa Clara, CA 95050
Analysis Date:	June 17, 2020	Phone: (650) 779-5577
Expiration Date:	June 12, 2021	
Storage:	Store at -20 °C	

**Kit Components**

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Sahara One-Step RT-qPCR Master Mix with UNG

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One-Step DNA/RNA Extraction Buffer

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Coronavirus Environmental Oligo Mix

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Cofactor Buffer

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DNase/RNase-Free Distilled Water

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SARS-CoV-2 N Positive Control

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Environmental swabs, sterile

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**Quality Control Data**

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**Sahara One-Step RT-qPCR Master Mix with UNG**

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**RT-qPCR Functional Assay**

**Result**

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A quantitative detection of log-fold serial dilutions of a control RNA as template. Linear regression analysis of quantification cycle ( $C_q$ ) versus log input quantity must give an amplification efficiency of between 90–110% and coefficient of determination ( $R^2$ )  $\geq 0.99$ .

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PASS

**PCR Carry-Over Digestion Functional Assay**

**Result**

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A quantitative detection of log-fold serial dilutions of a control amplicon as template in presence and absence of Uracil-DNA Glycosylase (UNG). Absolute value of  $C_q$  with and without UNG was monitored. UNG must result in a significant shift in  $C_q$  ( $\geq 10$ ), or no amplification.

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## One-Step DNA/RNA Extraction Buffer

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### Physical Appearance

### Result

One-Step DNA/RNA Extraction Buffer must have a clear, colorless appearance.

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### pH

### Result

One-Step DNA/RNA Extraction Buffer must have a pH between 5.6–6.4.

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### Quantification of Extracted RNA

### Result

A quantitative detection of extracted viral RNA from *Escherichia coli* bacteriophage MS2 (ATCC 15597-B1) as control. A quantitative detection of the extracted viral RNA reading the absorbance at OD260 and RT-qPCR must confirm high yield extraction of viral RNA.

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PASS

## Coronavirus Environmental Oligo Mix

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### Functional Assay

### Result

A quantitative detection of log-fold serial dilutions of SARS-CoV-2 gRNA as template. Linear regression analysis of  $C_q$  versus log input quantity must give an amplification efficiency of between 90–110% and coefficient of determination ( $R^2$ )  $\geq 0.99$ . Internal Control (IC) must give similar  $C_q$  for all dilutions of SARS-CoV-2 gRNA (<2  $C_q$  difference).

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## Cofactor Buffer

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### Physical Appearance

### Result

Cofactor Buffer must have a clear, colorless to slightly pink appearance.

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### Performance

### Result

Performance of the Cofactor Buffer is evaluated through RT-qPCR amplification of a control RNA template. The Cofactor Buffer must show similar performance and result in similar  $C_q$  (<1  $C_q$  difference) compared to control Cofactor Buffer.

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## SARS-CoV-2 Positive Control

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### Sequence Verification

### Result

The sequence of the SARS-CoV-2 N Positive Control is verified with a quantitative detection of log-fold serial dilutions. The SARS-CoV-2 N Positive Control must give amplification using the Coronavirus Environmental Oligo Mix.

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PASS