

# SARS-CoV-2 Variant B.1.1.7 Multiplex RT-qPCR Screening Kit (SCVUK)

Catalog #RU7118 100 samples

### **Product Description**

Coronaviruses are a family of large RNA viruses with size ranging from 26 to 32 kb. These viruses are zoonotic and in humans can cause respiratory infections. As the coronavirus is an RNA virus, it has a relatively high mutation rate resulting in rapid evolution. In December 2019, a new deadly coronavirus known as SARS-CoV-2, which has a high sequence similarity to SARS-CoV, was identified as the cause of the Covid-19 outbreak. Since then, numerous variants of SARS-CoV-2 were reported around the world. Among them, one variant strain first identified in the United Kingdom, known as the B.1.1.7 lineage, has gained enormous attention because it has a much higher infection rate and has become a major strain spreading globally. Among the total of 23 mutations accumulated in the B.1.1.7 lineage, two mutations are thought responsible for the increased transmissibility. The spike-HV 69-70 deletion mutation may promote viral immune escape, and the spike-N501Y mutation may contribute to the higher viral binding affinity to the ACE2 receptor on the human cell membrane. Between these two mutations, the spike-HV 69-70 deletion is considered the hallmark of the B.1.1.7 lineage, as it is not present in other major circulating SARS-CoV-2 strains including 501Y.V2, P.1, and P.2.

ScienCell's SARS-CoV-2 Variant B.1.1.7 Multiplex RT-qPCR Screening Kit (SCVUK) is designed to screen for the possible presence of the B.1.1.7 lineage by detecting the spike-HV 69-70 deletion. Two multiplex primer/probe set components (Cat #7118-REF and #7118-H69V70) are included in the kit. The reference primer/probe set component (Cat #7118-REF) contains 3 primer/probe sets, N1-FAM, N2-FAM, and RP-HEX (Table 1). Among them, N1-FAM and N2-FAM target two regions on the coronavirus SARS-CoV-2 nucleocapsid (N) gene. RP-HEX targets the exon 1 of human RPP30 gene and serves as a control to assess specimen quality. The HV 69-70 primer/probe set component (Cat #7118-H69V70) contains 2 primer/probe sets, H69V70-Del-FAM and H69V70-Present-HEX (Table 2), which target the coronavirus SARS-CoV-2 spike (S) gene with HV 69-70 deleted and present, respectively. For more efficient screening, if the expected mutation rate is low, a pool of up to 10 RNA samples can be used as the template for one qPCR reaction. If results for H69V70-Del-FAM are negative, then all pooled samples do not contain the spike-HV 69-70 deletion mutation. For H69V70-Del-FAM positive pooled samples, the samples should be tested individually to identify the H69V70-Del-FAM positive one(s). Please refer to Tables 6 and 7 for results interpretation.

**Table 1.** Primer/probe set list of the reference primer/probe set component (Cat #7118-REF)

Primer/Probe Set	Primer/Probe Target	Probe Reporter Dye
N1-FAM	SARS-CoV-2 nucleocapsid (N) gene, region 1	FAM
N2-FAM	SARS-CoV-2 nucleocapsid (N) gene, region 2	FAM
RP-HEX	Human RPP30 gene	HEX

**Table 2.** Primer/probe set list of the HV 69-70 primer/probe set component (Cat #7118-H69V70)

Primer/Probe set	Primer/Probe Target	Probe Reporter Dye
H69V70-Del-FAM	SARS-CoV-2 spike (S) gene, HV 69-70 deleted	FAM
H69V70-Present-HEX	SARS-CoV-2 spike (S) gene, HV 69-70 present	HEX

In addition, ScienCell One-Step TaqProbe RT-qPCR master mix (Cat #MB802a), a non-infectious positive control (Cat #7118-Pos), and nuclease-free water (Cat #7118-H2O) are included in the kit. The positive control (Cat #7118-Pos) consists of non-infectious viral RNA fragments of both the original strain and the B.1.1.7 lineage of SARS-CoV-2 spiked into human small airway epithelial cells. It serves to ensure reagents and instruments are working properly.

**Kit Components** 

Cat #	Component	Quantity	Storage
MB802a	One-Step TaqProbe RT-qPCR master mix, 4x	1.5 mL	-20°C
7118-REF	Reference multiplex primer/probe sets, in solution	600 μL	-20°C
7118-H69V70	HV 69-70 multiplex primer/probe sets, in solution	600 μL	-20°C
7118-H2O	Nuclease-free H <sub>2</sub> O	4 mL	4°C
7118-Pos	Positive control (non-infectious; RNA: 500 – 1000 copies/μL, cells: 200 – 300 counts/μL)	50 μL	-80°C

Additional Materials Required (Materials Not Included in Kit)

Component	Recommended
RNA samples	Customers' samples
Viral RNA isolation kit	ScienCell Viral RNA Isolation Kit (ScienCell, Cat #MB891)
qPCR plate or tube	

#### **Quality Control**

The primer/probe sets and the positive control are validated by RT-qPCR. The PCR products are analyzed by gel electrophoresis.

#### **Product Use**

For Research Use Only. Not for use in diagnostic procedures.

#### **Shipping and Storage**

The product is shipped on dry ice. Upon receipt, store the One-Step TaqProbe RT-qPCR master mix (Cat #MB802a) and the primer/probe sets (Cat #7118-REF and #7118-H69V70) at -20°C in a manual defrost freezer, the positive control (Cat #7118-Pos) at -80°C, and nuclease-free  $H_2O$  (Cat #7118-H2O) at 4°C.

#### **Procedures**

*Important: Only* use nuclease-free reagents in PCR applications.

- 1. Prior to use, allow the multiplex primer/probe sets (Cat #7118-REF and #7118-H69V70) to thaw to room temperature in the dark. Shake gently to mix well.
- 2. Centrifuge the vials at 1,500x g for 1 minute.
- 3. Aliquot multiplex primer/probe sets as needed. Store at -20°C in a manual defrost freezer. Avoid repeated freeze-and-thaw cycles. Maintain cold and in the dark when thawed.
- 4. For each test run, two control samples should be included, the non-infectious positive control (Cat #7118-Pos), and H<sub>2</sub>O (Cat #7118-H2O) as the No Template Control (NTC). Prepare two 20 μl RT-qPCR reactions as shown in Table 3 for each control sample, one with the reference primer/probe set component (Cat #7118-REF), and one with the HV 69-70 primer/probe set component (Cat #7118-H69V70).

Table 3.

Control sample (Cat #7118-Pos or 7118-H2O)	5 μ1
Multiplex primer/probe sets (Cat #7118-REF or #7118-H69V70)	6 μl
1-step RT-qPCR Master mix, 4x (Cat #MB802a)	5 μ1
Nuclease-free H <sub>2</sub> O (Cat #7118-H2O)	4 μ1
Total volume	20 μl

5. For each extracted RNA test sample (individual or pooled of up to 10 samples), prepare two 20 μl RT-qPCR reactions as shown in Table 4, one with the reference primer/probe set component (Cat #7118-REF), and one with the HV 69-70 primer/probe set component (Cat #7118-H69V70).

Table 4.

1-step RT-qPCR Master mix, 4x (Cat #MB802a)	5 μ1
Nuclease-free H <sub>2</sub> O (Cat #7118-H2O)  Total volume	4 μl <b>20 μl</b>

- 6. Seal the RT-qPCR reaction wells. Centrifuge the plates or tubes at 1,500x g for 15 seconds.
- 7. Setup RT-qPCR reactions as shown in Table 5.

**Table 5.** Instrument settings for RT-qPCR reactions. Fluorescence data for both FAM and HEX channels should be collected during the data acquisition step.

Step	Temperature	Time	Number of cycles
UNG incubation	25°C	2 min	1
Reverse transcription	50°C	15 min	1
Enzyme activation	95°C	2 min	1

Denaturation	95°C	3 sec	
Annealing and extension	66°C	30 sec	45
Data acquisition	Plate read, detect		

## **Results Interpretation**

**Table 6.** SCVUK kit control sample test results interpretation. A Cq value lower than 40 is considered positive.

Sample	Primer/probe Set	FAM	HEX	Results Interpretation
	7118-REF	+	+	Expected
7110 D	/116-REF	-	-	Reverse transcription and/or PCR failed
7118-Pos	7110 1100170	+	+	Expected
	7118-H69V70	-	-	Reverse transcription and/or PCR failed
	7110 PEF		-	Expected
7110 1120	7118-REF	If anyone is positive		Reagent(s) contaminated
7118-H2O	7110 11(0)170	-	-	Expected
	7118-H69V70	If anyone is positive		Reagent(s) contaminated

**Table 7.** SCVUK kit target sample test results interpretation when control results are as expected. A Cq value lower than 40 is considered positive.

Primer/probe Set	7118-REF		7118-H69V70		Dogulta Intonnuctation	
<b>Detection channel</b>	FAM	HEX	FAM	HEX	- Results Interpretation	
	+	±	+	-	SARS-CoV-2 detected, B.1.1.7 lineage implied	
	+	±	-	+	SARS-CoV-2 detected, NOT B.1.1.7 lineage	
Results	+	±	+	+	SARS-CoV-2 detected, mix of lineages with B.1.1.7 present implied	
Results	+	±	-	-	SARS-CoV-2 detected, possibly a novel lineage	
	-	+	-	-	SARS-CoV-2 NOT detected	
All other combinations		Invalid result				