TEST REPORT



LABORATORY ADDRESS:

Level 4, Block N & O, Faculty of Medicine, University of Malaya 50603 Kuala Lumpur.

Tel: +603-79676670 Email: tidrec@um.edu.my

Website: www.tidrec.com

HEAD OF LABORATORY: Sazaly Abu Bakar, Ph.D., FASc

TEST REPORT NO:	TS4-02	DATE OF ISSUE: 12/10/2020			
CUSTOMER DETAILS					
NAME	Life3 Technology Pte Ltd				
ADDRESS	17 Kaki Buki Place, Singapore 416195				
CONTACT	Dr. Steve Siaw				
SAMPLE & TEST IN	SAMPLE & TEST INFORMATION				
JOB NO.	TS4-0285				
DATE RECEIVED	02.09.2020				
TEST	22.09.2020				
PERFORMED					
ENVIRONMENTAL	Ambient Temperature: Store below 30°C				
CONDITIONS	Relative Humidity: NA				
TYPE OF SAMPLE	Quantum-Ion® E-Series Disinfectant				
	Sample delivered in liquid form in clear plastic bottle (500 ml)				
C	Active ingredients: Titanium Dioxide, Silver and Copper Ion				
	Sample Colour: Translucent White				
SAMPLE ID.	Lot No: FRT/2020/02				
	MGF Date: 1/9/2020				
	EXP Date: 31/8/2022				
TEST METHOD					
(TM)	V	EN14476			
Please tick ($$) at least					
one TM					

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DREC@UM



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RESULTS

Product Suppression Assay

The product suppression assay was performed to accurately determine the activity of the test product at the given contact time. The activity was suppressed by adding cold DMEM+2% FBS, followed by serially diluting it 10-fold in cell culture medium. The suppression of product activity was assayed at 1 min exposure. Results from the suppression assay showed no difference in viral titers compared to controls (Table 1). This suggested that addition of cold media and serial dilution effectively suppresses the product activity, resulting in no reduction of the viral titers.

Table 1: Suppression of product activity

*Contact time (sec)	Interfering substance	Viral Titer [Control] (TCID50/ml)	Viral Titer [After product suppression] (TCID50/ml)	Difference in Viral Titre (TCID50/ml)
60	clean conditions	5.4×10^5	No inhibition	0.00
60	dirty conditions	5.4×10^5	No inhibition	0.00

^{*}Undiluted mixture

Virucidal activity of Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray

The Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray was tested against SARS-CoV-2 in accordance to the European Standard EN14476:2013/FprA1:2015. Manifestation of virus cytopathic effects in cell culture were determined by comparing the study product-treated groups against that of the water - treated controls. The SARS-CoV-2 titer in the control-treated samples under clean and dirty conditions, respectively, were at 5.42 x 10⁵ TCID₅₀/ml. The Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray when tested neat achieved >5 log₁₀ reduction in viral titers when exposed for 1min, 5min and 10min under both clean and dirty condition (Table 2).

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Table 2: Virucidal activity of Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray against SARS-CoV- 2

Virus	The Fusion Resonance Quantum- Ion® E-Series Disinfectant Spray	Log ₁₀ Reduction in viral titers compared to control					
SARS-		Clean Condition			Dirty Condition		
CoV-2	Undiluted	1min	3min	10min	1min	3min	10min
		>5.00	>5.00	>5.00	>5.00	>5.00	>5.00

SUMMARY

The virucidal efficacy of The Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray was tested against SARS- CoV-2 in a suspension assay following the European Standard EN144762013/FprA1:2015 protocol. The Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray when tested undiluted demonstrated potent and rapid virucidal activity of ≥5 log₁₀ reduction of SARS-CoV-2 viral titer in 1 min in both clean and dirty conditions. The Fusion Resonance Quantum-Ion® E-Series Disinfectant Spray, hence, can kill 99.99% SARS-CoV-2.

PREPARED BY:	APPROVED BY:
Name: Pouya Hassandarvish Position: Postdoctoral Research Fellow	Approved Signatory(ies): Sazaly Abu Bakar, Ph.D., FASc Teoh Boon Teong, Ph.D. Juraina Abd Jamil, MMedSc
Date: October 09, 2020	Date: October 09, 2020