

## 广州市微生物研究所有限公司

GUANG ZHOU INSTITUTE OF MICROBIOLOGY CO., LTD.

# 检测报告

TEST REPORT

Report Number

QX20210652

Name of Sample

UVC Air Disinfection Unit

Applicant

Signify Luminaires (Shanghai) Co., Ltd.









# GUANG ZHOU INSTITUTE OF MICROBIOLOGY CO., LTD. TEST REPORT

Date Received: Aug. 20, 2021 Date Analyzed: Aug. 23, 2021

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Name of Sample	UVC Air Disinfection Unit	Source of Sample	Delivery
Applicant	Signify Luminaires(Shanghai) Co., Ltd.	Client	- 8
Manufacturer	Zhejiang Howell Illuminating Technology Co., Ltd.	Brand	
Type and Specification	UVCA110	Quantity of Sample	1PC
Date of Production	2021.05.28	State of Sample	Machine
Batch Number	{ C	Packing of Sample	In box
Standard and Methods	<technical disinfection="" for="" standard="">2002-2.</technical>	1.3 Air disinfection effect	evaluation test
Items of Analysis	Simulated Field Test (Staphylococcus albus 803	32)	0
Remarks	Applicant Address: 2F,Building 1, No.2555, He Manufacturer Address: No.1228 Tanjialing We Zhejiang, China.		- 1

\*\*\*To be continued\*\*\*







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#### Method for Testing Air Disinfection:

1. Test Equipment

1) Strain: Staphylococcus albus

2) Microbial aerosol generator: TK-3

3) Culture media: NA

4) Sampling equipment: six-stage sieve sampler

2. Test Conditions

1) The volume of the test chamber: 10 m<sup>3</sup>

2) Environment temperature: (20~25) °C

3) Environment humidity: (50~70) % RH

3. Operation Conditions of the Machine

Set the switch to position "The highest wind speed".

4. Test Procedures

- 1) Get a bacteria slant culture (4~5 generation) which is incubated at 37 °C for 24 h, wash the culture from this slant with 10 mL NB, filter the liquid culture by aseptic cotton buds, and dilute this inoculum with NB to suitable concentration. Then make atomized bacterial suspension.
- 2) The equipment is placed in the two test chambers respectively, close the door, and open the HEPA filter. Simultaneously operate the environmental control devices until the experimental cabin temperature to be (20~25) °C, relative humidity to be (50~70) %RH, Turn off the chamber environmental control system.
- 3) Release microbial aerosol: turn on the microbial aerosol generator, then turn on the ceiling fan, turn off the fan after 5 min, and let stand for 5 min.
- 4) At the same time, the test group and the control group were sampled with six-stage sieve sampler.
- 5) The test group started the sample and sampled after 120 min of action, and the control group also sampled in the corresponding time period.
- 6) Choose 2 NA plates (the same batch) as the negative control, and culture them on the same condition with the samples.
- 7) Run the test three times.
- 5. Computational Formula

Natural decay rate 
$$N_t(\%) = \frac{V_0 - V_t}{V_0} \times 100$$

Where:  $V_0$  = Original Bacteria Count of Control group;  $V_t$  = Bacteria Count after Treatment of Control group.

Killing Rate 
$$K_t(\%) = \frac{V_1 \times (1 - N_t) - V_2}{V_1 \times (1 - N_t)} \times 100$$

Where:  $V_1$  = Original Bacteria Count of test group;  $V_2$  = Bacteria Count after Treatment of test group. \*\*\*To be continued\*\*\*







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Test results

A)	30			Control Group			Test Group			
	Number of Sample Test Stra	Test Strain	Test Time (min)	Test Number	Count	Bacteria Count after Treatment $V_t$ (cfu/m <sup>3</sup> )	Natural Decay Rate $N_t$ (%)	Original Bacteria Count V <sub>1</sub> (cfu/m³)	Bacteria Count after Treatment $V_2$ (cfu/m³)	Killing Rate $K_t$ (%)
	8	6		1	1.15×10 <sup>5</sup>	8.63×10 <sup>4</sup>	24.96	1.19×10 <sup>5</sup>	<7	>99.99
. (	QX20210638-1	Staphylococcu albus	s 120	2	1.19×10 <sup>5</sup>	8.85×10 <sup>4</sup>	25.63	1.16×10 <sup>5</sup>	C 1	>99.99
1 1/4	· · · · · · · · · · · · · · · · · · ·		3	1.15×10 <sup>5</sup>	8.86×10 <sup>4</sup>	22.96	1.22×10 <sup>5</sup>	<7	>99.99	

Note: No microorganisms grew in the negative control group.

\*\*\*End of report\*\*\*

Editor Checker Checker Elssuer

Date Reported











### Statements

- 1. The report would be invalid under the following conditions: altered, added, deleted, copied, without the special seal for inspection or signatures by approver.
- 2. For the received sample, the sample information in the report is claimed by the applicant, the inspection unit is not responsible for its authenticity. The report is responsible for the received sample only.
- 3. If there is any objection to the inspection report, it should be presented to the inspection unit within 15 working days from the issuance date, otherwise the report shall be deemed as having been accepted. Microbiological item is not subjected to retest.
- 4. The items marked with "\*" in the report are not accredited by CNAS or CMA. The items marked with "#" are accredited by CNAS. The items marked with "+" are accredited by CMA.
- 5. The test data and results of items which are not accredited by CMA, only can be used as scientific research, teaching or internal quality control.
- 6. Any ambiguity by the language which used in the report, the Chinese shall prevail.

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