



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

500

Report Number

QX20210681

Name of Sample

UVC Air Disinfection Unit

Applicant

Signify Luminaires (Shanghai) Co.,

检验检测专用章







GUANG ZHOU INSTITUTE OF MICROBIOLOGY CO., LTD.

TEST REPORT

Date Received: Jul. 26, 2021 Date Analyzed: Aug. 04, 2021

		Date Allai	yzed. Aug. 04, 2021		
Name of Sample	UVC Air Disinfection Unit	Source of Sample	Delivery		
Applicant	Signify Luminaires (Shanghai) Co., Ltd.	Client	Kang Yangying		
Manufacturer	Zhejiang Howell Illuminating Technology Co., Ltd.	Brand			
Type and Specification	UVCA110	Quantity of Sample	1PC		
Date of Production	2021-5-28	State of Sample	Machine		
Batch Number	<u>~</u> *	Packing of Sample	In box		
Standard and Methods	GB 28235-2020 Hygienic requirements for ultraviolet appliance of disinfection				
Items of Analysis	Ultraviolet Intensity	D.A.	30		
八	Applicant Address: 2F,Building 1, No.2555, Hechuan Road, Minhang District, Shanghai;				
Remarks	Manufacturer Address: No.1228 Tanjialing West Road, Lanjiang Street, Yuyao, Ningbo				
30	Zhejiang, China.	W ₂₂			

To be continued







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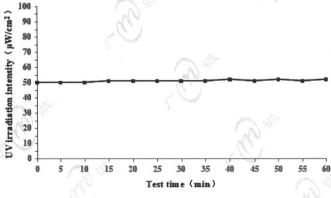
Method for Testing Ultraviolet Intensity:

- 1. Test Equipment
 - Model ST-512 ultraviolet radiometer, sensibility: 1μW/cm²
- 2. Test Procedure
 - 1) Test the background concentration.
 - 2) After the sample is opened and stabilized, the ultraviolet intensity is measured at the center 1m away from the sample lamp tube with the ultraviolet illuminance meter. A point is tested every 5 minutes for 60 minutes continuously. The final result is the average value of all test data.

Test Results

Test Results			M	
Number of Sample	Items	Vertical distance between test point and lamp (m)	Test Time (min)	Results (µW/cm²)
	D. 20.		0	50
· ·		Co	5	50
14 m	- B		10	50
	W.		15	51
- A	5		20	51
W.			25	_{3Y} 51 ₹
0.000102101		ar .	30	51
QX20210518-1 U	ltraviolet Intensity		35	51
yes		200	40	52
		- 0	45	51
	20 m	Park	50	52
		No	55	51
N Err		- San Anna	60	52
	do		Average value	51

UV irradiation intensity curve



End of report

Editor The Checker A Issue

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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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