广州市微生物研究所 GUANG ZHOU INSTITUTE OF MICROBIOLOGY

检测报告 TEST REPORT

| Report Number Name of Sample | KY20200051 | | | | |
|---------------------------------|---------------------|--|--|--|--|
| Name of Sample | Air Purifier | | | | |
| Applicant | LightAir Holding AB | | | | |

GUANG ZHOU INSTITUTE OF MICROBIOLOGY TEST REPORT

Date Received: Feb. 13, 2020

| Name of Sample | Air Purifier | Source of Sample | Delivery | | | | | |
|------------------------|--|--------------------|------------------------|--|--|--|--|--|
| Applicant | LightAir Holding AB | Client | Wilson Lam | | | | | |
| Manufacturer | Sino Vantage Industrial Ltd. | Brand | LightAir | | | | | |
| Type and Specification | Mini100WEWE | Quantity of Sample | 1PC | | | | | |
| Date of Production | - OF | State of Sample | Machine | | | | | |
| Batch Number | = 30 | Packing of Sample | In box | | | | | |
| Sample Picture | 1. Refer to GB/T 18801-2015 Air cle | Ughtar | | | | | | |
| Standard and Methods | Refer to GB/1 18801-2013 Air CR Refer to <technical forevaluation="" standard="" td="" test<=""><td></td><td>Air disinfection effec</td></technical> | | Air disinfection effec | | | | | |
| Items of Analysis | Purification Effect of Airborne Virus Aerosol (Influenza A virus A/PR8/34 H1N1) | | | | | | | |
| itellis of Allalysis | | | 110.0 1 1111.11 | | | | | |

To be continued



GUANG ZHOU INSTITUTE OF MICROBIOLOGY TEST REPORT

Date Received: Feb. 13, 2020 Date Analyzed: Feb. 18, 2020

Test Method for Purification Effect of Airborne Virus Aerosols

1. Test Equipment

1) Strain: Influenza A virus A/PR8/34 H1N1

2) Cells: MDCK

2. Test Conditions

1) Environment temperature: (23~25) °C

2) Environment relative humidity: (50~60) %

3) Test time: 60min

4) The volume of the test chamber: 10 m³

5) Machine setting: "The highest gear".

Test Results

| Number of Sample | Virus | Test Number | Control Group | | | Test Group | | |
|------------------|--------------------|----------------|--------------------------------------|---------------------------------------|---------------------------------|--------------------------------------|---------------------------------------|-----------------------------|
| | | | 0 min (TCID ₅₀ /m³) | 60 min (TCID ₅₀ /m³) | Natural Decay Rate (%) | 0 min (TCID ₅₀ /m³) | 60 min (TCID ₅₀ /m³) | Purification Rate (%) |
| | | 1 | 2.85×10 ⁶ | 9.01×10 ⁵ | 68.39 | 1.33×10 ⁶ | 9.07×10 ³ | 97.84 |
| KY20200051-1 | A/PR8/34 (H1N1) | 2 | 6.24×10 ⁶ | 2.26×10 ⁶ | 63.78 | 6.24×10 ⁶ | 1.33×10 ⁵ | 94.12 |
| | | 3 | 1.33×10 ⁶ | 4.22×10 ⁵ | 68.27 | 1.33×10 ⁶ | 1.33×10 ⁴ | 96.85 |

*** End of report***





Statements

- 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 responsibility 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 report and the name of the inspection unit shall not be used for product labels, advertisements, awards and merchandise publicity.
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- 6. The test data and results of items which are not accredited by CMA, only used as scientific research, teaching or internal quality control.
- 7. Any ambiguity by the language which used in the report, the Chinese shall prevail.

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