



Number: GZHT02354657

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|-----------------------|--------------|---------------------|--------------|
| Report Ref: | GZHT02354657 | | |
| Date Received: | Oct 26, 2020 | Date Issued: | Nov 05, 2020 |

| | | | |
|----------------------|--|--|--|
| Company Name: | BAODING YINHONG YUHE MEDICAL DEVICE MANUFACTURING CO., LTD | | |
| Address: | NANLONGSHAN VILLAGE, DAWANGDIAN INDUSTRIAL PARK, XUSHUI DISTRICT BAODING CITY, HEBEI PROVINCE, CHINA | | |
| Contact Name: | Xiaoming Xu | | |

| | |
|---|--|
| The Following Sample Was Submitted And Identified By/On Behalf Of The Applicant As: | |
| End Uses | : Non-Sterile Medical Face Mask |
| Ratings | : Type IIR |
| Sample Name | : Disposable Medical Mask(Non-Sterile) |
| No. Of Sample | : One (50 pieces) |
| Size | : 17.5cmX9.5cm |
| Colour | : Black |
| Standard | : EN 14683:2019+AC:2019 |
| Brand Name | : YINHONYUHE |
| Manufacturer | : BAODING YINHONG YUHE MEDICAL DEVICE MANUFACTURING CO., LTD |
| Date received/ Test Started | : Oct 26, 2020 |
| Ref | : Type No.: YH/YY-2099R |

Test was conducted on specific items, at our client's request.

Approved by:

Jana

Emica Yu

Sr. Manager

Assistant Supervisor

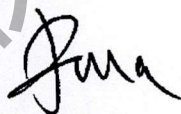


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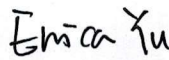
Original Sample Photo



Approved by:



Sr. Manager



Assistant Supervisor





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Summary of testing:

With reference to following standard:

- EN 14683:2019+AC:2019 Medical face masks – Requirements and test methods Type IIR

Materials Used in The Submitted Sample Were Found To Comply With The Type IIR Requirements of EN 14683:2019+AC:2019 with respect to Microbial Cleanliness, Bacterial Filtration Efficiency, Differential Pressure and Splash Resistance Pressure tests.

ORIGINAL

仅限查阅资质时使用

Approved by:

Sr. Manager

Assistant Supervisor



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Tests Conducted (As Requested By The Applicant)

1 Splash Resistance Pressure (ISO 22609:2004):

Synthetic Blood Surface Tension: 0.042N/m, Distance Between Blow Head Front End And Target Area: 305 mm, Artificial Blood Volumes: 2 mL, Blood Pressure: 16.0 kPa, Velocity: 550 cm/s, Without Targeting-plate Used

Condition test specimens for a minimum of 4 hours in an environment of temperature (21±5) °C and relative humidity (85±5)% and conduct the test within 1 minute of removal from conditioning chamber.

Test Environment Condition: Temperature 24.0°C, Relative Humidity 86.0%

| Tested Sample | Observation | Pass/Fail | Performance Requirement for Medical Face Mask |
|---------------|----------------|-----------|---|
| Specimen (1) | No Penetration | Pass | Type IIR No Penetration at 16.0 kPa |
| Specimen (2) | No Penetration | Pass | |
| Specimen (3) | No Penetration | Pass | |
| Specimen (4) | No Penetration | Pass | |
| Specimen (5) | No Penetration | Pass | |
| Specimen (6) | No Penetration | Pass | |
| Specimen (7) | No Penetration | Pass | |
| Specimen (8) | No Penetration | Pass | |
| Specimen (9) | No Penetration | Pass | |
| Specimen (10) | No Penetration | Pass | |
| Specimen (11) | No Penetration | Pass | |
| Specimen (12) | No Penetration | Pass | |
| Specimen (13) | No Penetration | Pass | |
| Specimen (14) | No Penetration | Pass | |
| Specimen (15) | No Penetration | Pass | |
| Specimen (16) | No Penetration | Pass | |
| Specimen (17) | No Penetration | Pass | |
| Specimen (18) | No Penetration | Pass | |
| Specimen (19) | No Penetration | Pass | |
| Specimen (20) | No Penetration | Pass | |
| Specimen (21) | No Penetration | Pass | |
| Specimen (22) | No Penetration | Pass | |
| Specimen (23) | No Penetration | Pass | |
| Specimen (24) | No Penetration | Pass | |
| Specimen (25) | No Penetration | Pass | |

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Tests Conducted (As Requested By The Applicant)

| | | |
|---|-----------------|------|
| Specimen (26) | No Penetration | Pass |
| Specimen (27) | No Penetration | Pass |
| Specimen (28) | No Penetration | Pass |
| Specimen (29) | No Penetration | Pass |
| Specimen (30) | No Penetration | Pass |
| Specimen (31) | No Penetration | Pass |
| Specimen (32) | No Penetration | Pass |
| Conclusion*: | Accepted | |
| * = An acceptable quality limit of 4.0% is met for a single sampling plan when 29 or more of the 32 tested specimens show "pass" results. | | |

- 2 Differential Pressure (EN 14683:2019+AC:2019 Annex C):
Air flow: 8L/min, Test area diameter 25 mm, Test area: 4.9 cm².

| Tested Sample | Result (Pa/cm ²)* | | | | | Performance Requirement for Medical Face Mask (Pa/cm ²) |
|--|-------------------------------|------------|------------|------------|------------|---|
| | Specimen 1 | Specimen 2 | Specimen 3 | Specimen 4 | Specimen 5 | |
| Location 1 | 42.5 | 43.9 | 37.8 | 42.7 | 44.1 | Type IIR < 60 |
| Location 2 | 40.0 | 31.6 | 42.2 | 34.7 | 44.5 | |
| Location 3 | 46.5 | 40.8 | 40.9 | 42.2 | 55.3 | |
| Location 4 | 38.9 | 37.5 | 37.3 | 37.4 | 39.0 | |
| Location 5 | 33.9 | 32.5 | 33.4 | 37.4 | 43.2 | |
| Average | 40.4 | 37.3 | 38.3 | 38.9 | 45.2 | |
| * = All the locations were evenly taken from the main mask body. | | | | | | |

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Tests Conducted (As Requested By The Applicant)

3 Microbial Cleanliness

As Per EN 14683:2019+AC:2019 Medical Face Masks-Requirements and Test Methods Annex D.

| Test Item | Result (cfu/g) | | | | | Limit (cfu/g) |
|-----------------------|------------------|--------------|--------------|--------------|--------------|---------------|
| | Specimen (1) | Specimen (2) | Specimen (3) | Specimen (4) | Specimen (5) | |
| Microbial Cleanliness | <1 | <1 | <1# | <1 | <1 | Type IIR ≤30 |

Remark:

cfu = colony forming unit

≤ = Not more than

= No colony was detected at the extraction liquid of the samples.

Remark: This test item was conducted in Caipin Road, Guangzhou Science City, GETDD, Guangzhou, Guangdong.

4 Bacterial Filtration Efficiency (BFE)

Test Method: EN 14683: 2019+AC: 2019 Annex B

Summary of Test Method:

A specimen of the mask material is clamped between a six-stage cascade impactor and an aerosol chamber. The bacterial aerosol is introduced into the aerosol chamber using a nebulizer and a culture suspension of Staphylococcus aureus. The aerosol is drawn through the medical face mask material using a vacuum attached to the cascade impactor. The six-stage cascade impactor uses six agar plates to collect aerosol droplets which penetrate the medical face mask material. Control samples are collected with no test specimen clamped in the test apparatus to determine the upstream aerosol counts. The agar plates from the cascade impactor are incubated for (48 ± 4) h and counted to determine the number of viable particles collected.

The bacterial filtration efficiency (BFE) of the mask is given by the number of colony forming units passing through the medical face mask material expressed as a percentage of the number of colony forming units present in the challenge aerosol.

Conditioning of the Specimens: 4 h at (21 ± 5) °C and (85 ± 5) % relative humidity

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Tests Conducted (As Requested By The Applicant)

Test Condition:

Biological Aerosol: *Staphylococcus aureus* (ATCC 6538)

Testing side: Inside of the test specimen was facing towards the challenge aerosol

Test area: 78 cm²

Flow rate: 28.3 L/min

The average plate count results of the positive controls: 2.0×10³ CFU

The average plate count results of the negative controls: < 1 CFU

Mean particle size (MPS): 2.7 μm

Incubation condition: (37 ± 2) °C for (20 to 52) h

Number of test specimens: 5

Test Procedure:

1. Preparation of the bacterial challenge: Dilute the culture in peptone water to achieve a concentration of approximately 5×10⁵ CFU/mL.
2. Deliver the challenge to the nebulizer using a peristaltic or syringe pump. Connect tubing to nebulizer and peristaltic pump and into the challenge suspension; purge tubing and nebulizer of air bubbles.
3. Perform a positive control run without a test specimen clamped into the test system to determine the number of viable aerosol particles being generated.
4. Initiate the aerosol challenge by turning on the air pressure and pump connected to the nebulizer.
5. Immediately begin sampling the aerosol using the cascade impactor. Adjust the flow rate through the cascade impactor to 28.3 L/m.
6. Time the challenge suspension to be delivered to the nebulizer for 1 min.
7. Time the air pressure and cascade impactor to run for 2 min.
8. At the conclusion of the positive control run, remove plates from the cascade impactor.
9. Place new agar plates into the cascade impactor and clamp the test specimen into the top of the cascade impactor, with the inside oriented toward the challenge as intended.
10. Repeat the challenge procedure for each test specimen and positive control sample.
11. Perform a negative control sample by collecting a 2 min sample of air from the aerosol chamber. No bacterial challenge should be pumped into the nebulizer during the collection of the negative control sample.
12. Incubate agar plates at (37 ± 2) °C for (48 ± 4) h.
13. Count each of the six-stage plates of the cascade impactor.
14. Total the counts from each of the six plates for the test specimens and positive controls. Calculate the filtration efficiency percentages.

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Tests Conducted (As Requested By The Applicant)

Calculation:

The Bacterial Filtration Efficiency (BFE), was calculated as a percentage using the following equation:

$$\% \text{ BFE} = (C-T)/C \times 100$$

where,

- C = Average plate counts total for test controls;
- T = Plate count total for the test specimen.

Test Result:

| Tested Specimen | Result | | Performance Requirement in EN 14683: 2019+AC: 2019 (% BFE) |
|-----------------|---------------------------------|---|--|
| | The Total Plate Count (T) (CFU) | Bacterial Filtration Efficiency (BFE) (%) | |
| Specimen (1) | 3 | 99.9 | Type IIR: ≥ 98 |
| Specimen (2) | 3 | 99.9 | |
| Specimen (3) | 6 | 99.7 | |
| Specimen (4) | 5 | 99.8 | |
| Specimen (5) | 10 | 99.5 | |

Remarks:

CFU = Colony Forming Unit

This item was conducted in Caipin Road, Guangzhou Science City, GETDD, Guangzhou, Guangdong.

End of Report

This report is made solely on the basis of your instructions and/or information and materials supplied by you. It is not intended to be a recommendation for any particular course of action. Intertek does not accept a duty of care or any other responsibility to any person other than the Client in respect of this report and only accepts liability to the Client insofar as is expressly contained in the terms and conditions governing Intertek's provision of services to you. Intertek makes no warranties or representations either express or implied with respect to this report save as provided for in those terms and conditions. We have aimed to conduct the Review on a diligent and careful basis and we do not accept any liability to you for any loss arising out of or in connection with this report, in contract, tort, by statute or otherwise, except in the event of our gross negligence or wilful misconduct. No copy of the test report(except for full text copy) shall be made without the written approval by Intertek.

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