

Test Report

No.T32020280526SN

Date: Oct 23, 2020

Page 2 of 2

Test Results:

Viral filtration efficiency (VFE) With reference to ASTM F2101-19

Test Side : White Side (Inside)

Pre-Conditioning : Minimum of 4 hours at 21±5°C and 85±5% R.H.

Test Condition : 21±2 °C and 30-50% R.H.

VFE Test Area : ~40 cm²

VFE Flow Rate : 28.3 Litre per minute (L/min)

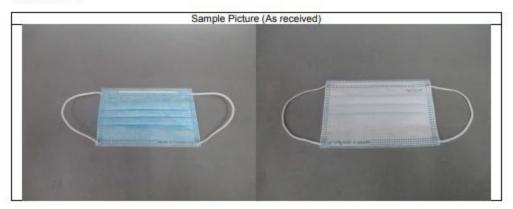
Mean Particle Size : 2.7 µm

Positive Control Average : 2.4 x 10³ PFU Negative Monitor Count : <1 PFU

Test Specimen	Percent VFE (%)	
1	>99.9	
2	99.9	
3	>99.9	
4	99.9	
5	99.9	

Note: The above test results was conducted by a SGS assessed competent subcontractor laboratory

Sample Photo:



SGS authenticate the photo on original report only

*** End of Report ***

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Unless otherwise stated the results shown in this test report refer only to the sample(s) tested and such sample(s) are retained for 30 days only.

905 thing long Limited | Office: Units: 303 & 305, 30F Suiteting 22E, Phase 3, HK Science Park, New Territories, Hong Kong 1 (852) 2334 4481 | f (852) 2754 3125 | e-mistg/his/Sogs.com Laboratory, 10F, Units: 16-29 30F, 44F & 54F, On Wall Centre, 25 Lot Yo Read, Favling, New Territories, Hong Kong, www.spigroup.com.htm.



Testimonial of Test Results

Our Reference: CFD083/2020

Company Name & Address

Ever Sharp (China) Limited

No.1 11/F Topsail Plaza 11 On Sum Street Shatin HK

Product Description

Medical Face Mask

Conclusion(s)

This is to testify that the test results of the above submitted sample conform jointly to the test requirements of **ASTM F2100-19 Level 3**:

- Bacteria Filtration Efficiency (BFE)
- Particulate Filtration Efficiency (PFE) at 0.1micron
- Differential Pressure
- Resistance to Penetration by Synthetic Blood at 160mmHg
- Flammability to Class 1 according to the test reports (HC20080563 issued on 2020-08-28)

Date of Tests

2020-08-13 to 2020-08-28

Test Report Number(s)

HC20080563

Authorized Signatory: _

U Dan

Name: LAU Yuk Kuen, Joey

Date of Issued: 2020-09-01



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- The Company shall not be called or be liable to be called to give evidence or testimony on the Report in a court of law without its prior written consent, unless required by the relevant governmental authorities, laws or court orders.
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- 12 . Issuance records of the Report are available on the internet at www.stc.group. Further enquiry of validity or verification of the Reports should be addressed to the Company.



Sponsor: Yang Dexing Xiantao Dexing Plastic Products Co., Ltd No.3 Shengli Up Street, Penchang Town Xiantao, Hubei Province, 433018 CHINA



Bacterial Filtration Efficiency (BFE) Final Report

Test Article: Product Name: Meltalown Filter Fabric

Type: DX-BFE-20 Lot #DX180626

Study Number: 1070289-S01 Study Received Date: 09 Jul 2018

Testing Facility: Nelson Laboratories, LLC

6280 S. Redwood Rd.

Salt Lake City, UT 84123 U.S.A.

Test Procedure(s): Standard Test Protocol (STP) Number: STP0004 Rev 15

Deviation(s): None

Summary: The BFE test is performed to determine the filtration efficiency of test articles by comparing the bacterial control counts upstream of the test article to the bacterial counts downstream. A suspension of *Staphylococcus aureos* was aerosolized using a nebulizer and dolivered to the test article at a constant flow rate and fixed air pressure. The challenge delivery was maintained at $1.7 - 2.7 \times 10^3$ colony forming units (CFU) with a mean particle size (MPS) of $3.0 \pm 0.3 \, \mu m$. The aerosols were drawn through a six-stage, viable particle, Andersen sampler for collection. This test method complies with ASTM F2101-14, EN 14683:2014, Annex B, and AS4381:2015.

All text method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test Side: Either

BFE Test Area: ~40 cm²

BFE Flow 536: 28.3 Liters per minute (L/min)

Conditioning Parameters: 85 ± 5% relative humidity (RH) and 21 ± 5°C for a minimum of 4 hours

Positive Control Average: 1.8 x 10³ CFU
Negative Monitor Count: <1 CFU

MPS: 2.9 µm







Study Director

801-290-7500

Janelle R. Bentz M.S.

Study Completion Date

1070289-S01

nelsonlabs.com

sales@nels or labs com

g

FRT0004-0001 Rev 19







Test Article Number	Percent BFE (%)
1	99.9
2	A6.3
3	99.7
4	99.8
5	99.6

The filtration efficiency percentages were calculated using the following equation:

$$\% BFE = \frac{C - T}{C} \times 100$$

C = Positive control average

T = Plate count total recovered downstream of the test article Note: The plate count total is available upon request







Sponsor:
Yang Dexing
Xiantao Dexing Plastic Products Co., Ltd.
No.3 Shengli Up Street, Pengonang Town
Xiantao, Hubei Province, 433018
CHINA



Latex Particle Challenge Final Report

Test Article: Product Name: Meltblown Filter Fabric

Type: DX-PFE-25 Lot #DX180626 1070288-S01

Study Number: 1070288-S01 Study Received Date: 09 Jul 2018

Testing Facility: Nelson Laboratories, LLC 6280 S. Redwood Rd.

Salt Lake City, UT 84123 U.S.A.

Test Procedure(s): Standard Test Protocol (STP) Number: STP0005 Rev 05

Deviation(s): None

Summary: This procedure was performed to evaluate the non-viable particle filtration efficiency (PFE) of the test anicle. Monodispersed polystyrene latex spheres (PSL) were nebulized, dried, and passed through the test article. The particles that passed through the test article were enumerated using a laser particle counter.

Three one-minute counts were performed, with the test article in the system, and the results averaged. Three one-minute control counts were performed, without a test article in the system, before and after each test article and the counts were averaged. Control counts were performed to determine the average number of particles delivered to the test article. The filtration efficiency was calculated using the average number of particles penetrating the test article compared to the average of the control values.

The procedure employed the basic particle filtration method described in ASTM F2299, with some exceptions; notably the procedure incorporated a non-neutralized challenge. In real use, particles carry a charge, thus this challenge represents a more natural state. The non-neutralized aerosol is also specified in the FDA guidance document on surgical face masks. All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test Side: Either Side
Area Tested: 91.5 cm²
Particle Size: 0.1 µm

Laboratory Conditions: 20°C, 33% relative humidity (RH) at 1119; 21°C, 33% RH at 1259

Average Filtration Efficiency. 99.86% Standard Deviation: 0.021

Results:

Test Article Manual Test	Average Test Article Counts	Average Control Counts	Filtration Efficiency (%)
11/1/49	20	12,834	99.84
2	19	13,772	99.86
3	13	10.913	99.88





Study Director

Brandon L. Williams

Study Completion Date



nelsor lab. .com

sales@nelsonlabs.com

osm

FRT0005-0001 Rev 4



IFICATE OF COMPLIANCE



This is to certify that

The Cleanroom at

Unit 1-7, 15-21, 11/F, Topsail Plaza 11 On Sum Street, Shatin, N.T., Hong Kong SAR

of Ownership

Ever Sharp (China) Limited

Complies with the Acceptance Criteria of

BS EN ISO 14644-1: 2015: Class 7

(Ambient Particle Concentration, HEPA filter leakage, Air Change Per Hour & Pressure Differential)

As detailed in

Test Report No.: RP2000218

Date of Certification: 24/07/2020

Lab (Asia) Lio

Date of issue:

Date of expiry: 24/07/2021

08/08/2020

Certified by:

Ir Donney Man Wai Leung Biosafety Consultant, MSc, MBA, BSc **NEBB Certified Cleanroom Professional**

The Lab (Asia) Ltd.

Tel: +(852) 2470 2588 Fax: +(852) 2470 2589

22 San Hi Tsuen Street, Ping Shan, N.T., Hong Kong. Email: info@thelab.asia Website: www.thelab.asia

Exp. 3/3... DONNEY M. LEUNG



The Lab (Asia) Ltd. is a member of the SGS Group.



Test Report

No.T32020280526SN

Date: Oct 23, 2020

Page 1 of 2

EVER SHARP (CHINA) LIMITED

FLAT 1-7, 15-21, 11/F TOPSAIL PLAZA, 11 ON SUM STREET, SHATIN, N.T., HONG KONG

The following samples were submitted and identified by/on behalf of the client as:

WELLMAN 惠民 DISPOSABLE 3-PLY MEDICAL FACE MASK

Case No. : CA320202822749

Lot No. / Batch Code : NOT PROVIDED

: BLUE MASK Sample Description

: 10 PCS **Quantity Submitted**

Manufacturer : EVER SHARP (CHINA) LIMITED

Country of Origin : HONG KONG Sample Receiving Date : AUG 20, 2020

Testing Period : AUG 20 TO OCT 23,Oct 23, 2020

Test Requested	Conclusion
Viral filtration efficiency (VFE) (With reference to ASTM F2101) See Reference to ASTM F2101)	

******** FOR FURTHER DETAILS, PLEASE REFER TO THE FOLLOWING PAGE(S) ********

Signed for and on behalf of SGS Hong Kong Ltd.

Au Kam Chi, Gigi Technical Manager

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Test Report No.T32020280526SN Date: Oct 23, 2020 Page 2 of 2

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Test Side White Side (Inside)

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: 21±2 °C and 30-50% R.H. **Test Condition**

VFE Test Area : ~ 40 cm²

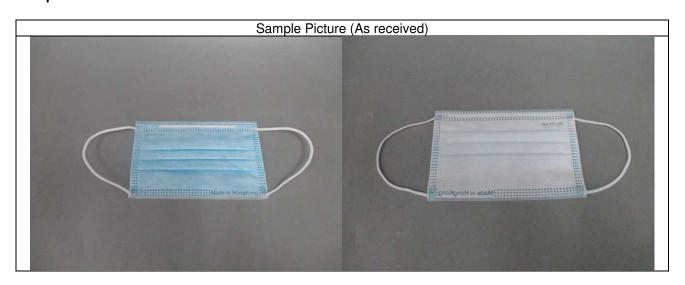
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Mean Particle Size : 2.7 μm
Positive Control Average : 2.4 x 10³ PFU
Negative Monitor Count : <1 PFU

T . C .	D 17/25 (0/)
Test Specimen	Percent VFE (%)
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2	99.9
3	>99.9
4	99.9
5	99.9

Note: The above test results was conducted by a SGS assessed competent subcontractor laboratory

Sample Photo:



SGS authenticate the photo on original report only

*** End of Report ***

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