

(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory) Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-040920-04, Page 01 of 03

### TEST REPORT

FABRIC TEST

Report Code: MS-040920-04

Issued To

Issue Date: 11/09/2020

PART A: Particulars of Sample submitted

A.	Sample Description	1.5	Fabric (with Spunbond)
			Sample ID NMAF01-0409-099-MP
B.	Date of Sample Received		04/09/2020
C.	Date of Commencement of Testing	:	04/09/2020
D.	Date of completion of Testing	;	11/09/2020
E.	Test Method	1:	AATCC-100
F.	Sample submitted By		Customer
G.	Instrument Used		Breathing machine, Aerosol Generator, Pressure Probe etc.

### Initial Concentration of Microbes

S. No.	Name of Test	Unit	Initial Concentration
1.	Total Bacterial Count	cfu/m³	2.31 x10 <sup>4</sup>
2.	Total Fungal Count	cfu/m³	3.14 x 10 <sup>4</sup>
3.	MRSA	efu/m³	1.84 x 10 <sup>4</sup>
4.	Yeast & Mold	cfu/m³	1.76 x 10 <sup>4</sup>

### Test Methodology: (Petri Plate Exposure Technique)

- 1. Air purifier with sample Tested in a chamber of 10ft x 10ft x 10 ft.
- 2. Fan was powered during operation.
- Petriplate having bacterial and fungal growth is exposed to air, where air purifier is running for different time period.
- 4. Exposed petriplate is incubated for colony forming in biological incubator.
- After incubation period, colony of bacteria and fungus is measured by colony counter and recorded.





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### RESULTS

S. No.	Time (Hr)	Observation (cfu/m³)	Reduction Percentage
1.	Initial	2.31 x 10 <sup>4</sup>	=
2.	1	$1.3 \times 10^4$	43.72
3.	4	$8.4 \times 10^3$	63.64
4.	8	$3.6 \times 10^{3}$	84.41
5.	24	76	99.67

2. Total Fungal Count Reduction Rate					
S. No.	Time (Hr)	Observation (cfu/m³)	Reduction Percentage		
1.	Initial	$3.14 \times 10^4$	-		
2.	1	1.86 x 10 <sup>4</sup>	40.76		
3.	4	9.78 x 10 <sup>3</sup>	68.85		
4.	8	4.92 x 10 <sup>3</sup>	84.33		
5.	24	83	99.73		

3. MRSA Count Reduction Rate				
S. No.	Time (Hr)	Observation (pfu)	Reduction Percentage	
1.	Initial	1.82 x 10 <sup>4</sup>	-	
2.	1	1.22 x 10 <sup>4</sup>	32.97	
3.	4	$7.64 \times 10^3$	58.02	
4.	8	$4.5 \times 10^3$	75.27	
5.	24	54	99.69	

4. Yeast Mold Count reduction Rate				
S. No.	Time (Hr)	Observation (pfu)	Reduction Percentage	
1.	Initial	1.76 x 10 <sup>4</sup>	-	
2.	1	$1.34 \times 10^4$	23.86	
3.	4	6.45 x 10 <sup>3</sup>	63.35	
4.	8	4.75 x 10 <sup>3</sup>	73.01	
5.	24	51	99.71	





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Report Code: MS-040920-04, Page 3 of 3

### Summary of Test:

S. No.	Name of Test	Positive Control Sample	Final Count after 24 hrs of Air Purifier	Reduction (%)
1.	Total Bacterial Count (cfu/m³)	2.31 x 10 <sup>4</sup>	76	99.67
2.	Total Fungal Count (cfu/m³)	3.14 x 10 <sup>4</sup>	83	99.73
3.	MRSA	1.82 x 10 <sup>4</sup>	54	99.69
4.	Yeast & Mold	1.76 x 10 <sup>4</sup>	51	99.71

Conclusions: On the basis of above tested parameters, sample having high antimicrobial activity with killing rate 99 % is recorded.

Notes.

cfu: colony forming unit

pfu: plaque forming unit

### Remarks:

Total Bacterial Count contains micrococcus, staphylococcus, Bacillus and Pseudomonas bacteria. Total Fungal Count contains cladosporium, Penicillium and Aspergillus Fungus.

#### Notes:

1. The results given above are related to the tested sample, as received & mentioned Parameters.

2. Responsibility of the Laboratory is limited to the invoiced amount only.

3. This test report will not be generated again, either wholly or in part, without prior written Permission of the laboratory.

Checked by



(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-040920-04, Page 01 of 02

TEST REPORT

FABRIC TEST

:

Report Code: MS-040920-04

Issued To

Issue Date: 23/09/2020

PART A: Particulars of Sample submitted

Α.	Sample Description	1:	Fabric (with Spunbond) Sample ID NMAF01-0409-099-MP
B.	Date of Sample Received		04/09/2020
C.	Date of Commencement of Testing	1 2	04/09/2020
D.	Date of completion of Testing	1.	23/09/2020
E.	Test Method	:	ASTM F2101
F.	Sample submitted By	1:	Customer
G.	Instrument Used	2	Breathing machine, Aerosol Generator, Pressure Probe etc.

### Initial Concentration of Polluting Dust

S. No.	Name Of Test Dust	Unit	Initial Concentration
1.	PM2.5	PPM	984

#### Test Method:

- 1. Air Purifier Tested in a chamber of 20ft x 10ft x 10 ft.
- 2. Fan was powered during operation.
- Concentration is monitored during operation of Air Purifier and also for Control sample without Air Purifier Operation.
- 4. Test Method is in below Table:

PM 2.5	CPCB Volume - 1		
1 171 2.3	CICB volume - 1		





(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

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1. F	M2.5 Removal Perfo	rmance	
S. No.	Time (Hr)	Observation (PPM)	Reduction Latching Percentage
1.	Initial	984	
2.	1	547	44.41
3.	4	284	71.14
4.	8	201	79.57
5.	24	170	98.27

### 2. Differential Pressure

S. No.	Test Parameter	Test Method	Unit	Result	Requirement
1.	Differential Pressure (Breathing Resistance)	EN 14683	Pa/cm <sup>2</sup>	52	•

#### Notes:

- 1. The results given above are related to the tested sample, as received & mentioned Parameters.
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Checked by

Authorized Signatory



(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-040920-04, Page 01 of 02

#### TEST REPORT

FABRIC TEST

Report Code: MS-040920-04

Issued To

Issue Date: 22/09/2020

PART A: Particulars of Sample submitted

A.	Sample Description	1:	Fabric (with Spunboud) Sample ID NMAF01-0409-099-MP
B.	Date of Sample Received		04/09/2020
C.	Date of Commencement of Testing	1 :	04/09/2020
D.	Date of completion of Testing	:	22/09/2020
E.	Test Method	:	ASTM F2101
F.	Sample submitted By	:	Customer
G.	Instrument Used	1	Breathing machine, Aerosol Generator, Pressure Probe etc.

### **Initial Concentration of Microbes**

S. No.	Name of Test	Unit	Initial Concentration
1.	Virus (Bacteriophage Virus)	pfu	1.74 x10 <sup>4</sup>

### Test Methodology: (Petri Plate Exposure Technique)

- 1. Air purifier with filter Tested in a chamber of 10ft x 10ft x 10 ft.
- 2. Fan was powered during operation.
- Petriplate having Bacteriophage Virus exposed to air, where air purifier is running for different time period.
- 4. Exposed petriplate is incubated for colony forming in biological incubator.
- 5. After incubation period, colony of virus is measured by colony counter and recorded.





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Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

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### RESULTS

1. Bacteriophage Virus Killing Rate			
S. No.	Time (Hr)	Observation (pfu)	Reduction Percentage
1.	Initial	1.74 x 10 <sup>4</sup>	-
2.	1	1.34 x 10 <sup>4</sup>	22.99
3.	4	$2.14 \times 10^3$	87.70
4.	8	3.47 x 10 <sup>2</sup>	98.00
5.	24	31	99.82

### Summary of Test:

S. No.	Name of Test	Positive Control Sample	Final Count after 24 hrs of Air Purifier	Reduction (%)
1.	Virus (Bacteriophage Virus) (pfu)	1.74 x 10 <sup>4</sup>	31	99.82

Conclusions: On the basis of above tested parameters, sample having high antimicrobial activity with killing rate 99 % is recorded.

Notes:

pfu: plaque forming unit

### Remarks:

Virus contains Bacteriophase Virus.

#### Notes:

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Checked by

Authorized Signatory



(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-040920-05, Page 01 of 03

TEST REPORT

FABRIC TEST

:

Report Code: MS-040920-05

Issued To

Issue Date: 11/09/2020

PART A: Particulars of Sample submitted

A.	Sample Description	1:	Fabric (with Spunbond)
			Sample ID NMAF01-0409-098-MP
B.	Date of Sample Received		04/09/2020
C.	Date of Commencement of Testing	1:	04/09/2020
D.	Date of completion of Testing	1.2	11/09/2020
E.	Test Method	1:	ASTM F2101
F.	Sample submitted By	:	Customer
G.	Instrument Used	:	Breathing machine, Aerosol Generator, Pressure Probe etc.

### Initial Concentration of Polluting Gases

S. No.	Name Of Test Gas	Unit	Initial Concentration
1.	Benzene (ppm)	PPM	5.4
2.	Hexane (ppm)	PPM	4.0
3.	Mercaptan (ppm)	PPM	5.6
4.	Xylene (ppm)	PPM	5.0
5.	Toluene (ppm)	PPM	4.9
6.	Formaldehyde (ppm)	PPM	5.0

### Test Method:

- 1. Air Purifier with filter Tested in a chamber of 10ft x 10ft x 10 ft.
- 2. Fan was powered during operation.
- Concentration is monitored during operation of fabric and also for Control sample without fabric Operation.
- 4. Test Method is in below Table:





(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-290720-05, Page 2 of 3

Benzene (ppm)	IS:5182 Part-XI
Hexane (ppm)	NIOSH 1501
Mercaptan (ppm)	NIOSH 1501
Xylene (ppm)	NIOSH 1501
Toluene (ppm)	NIOSH 1501
Formaldehyde (ppm)	NIOSH 1501

### RESULTS

1. Benzene Removal Performance				
S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage	
1.	Initial	5.4		
2.	1	3.1	42.59	
3.	4	2.0	62.96	
4.	8	0.64	88.14	
5.	24	0.005	99.90	

S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage
1.	Initial	4.0	-
2.	1	2.6	35.0
3.	4	1.60	60.0
4,	8	0.79	80.25
5.	24	0.0076	99.81

J. IVE	ercaptan Removal Pe	стогнансе	
S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage
1.	Initial	5.6	-
2.	1	3.2	42.86
3.	4	2.9	51.79
4.	8	0.92	83.57
5.	24	0.046	99.18





(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

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S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage
1.	Initial	5.0	·
2.	1	4.2	16.0
3.	4	1.6	68
4.	8	0.58	88.4
5.	24	0.062	98.75

S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage
1.	Initial	4.9	-
2.	1	3.3	32.65
3.	4	1.45	70.41
4.	8	0.37	92.45
5.	24	0.0081	99.83

S. No.	Time (Hr)	Observation (PPM)	Reduction Percentage
1.	Initial	5.0	-
2.	1	3.8	24.0
3.	4	2.9	42.0
4.	8	1.6	68.0
5.	24	0.09	98.20

#### Notes:

1. The results given above are related to the tested sample, as received & mentioned Parameters,

2. Responsibility of the Laboratory is limited to the invoiced amount only.

3. This test report will not be generated again, either wholly or in part, without prior written Permission at the aboratory.

Checked by



(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory) Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

Report Code: MS-040920-06, Page 01 of 02

TEST REPORT

FABRIC TEST

Report Code: MS-040920-06

Issued To

Issue Date: 11/09/2020

PART A: Particulars of Sample submitted

A.	Sample Description	1 7	Fabric (with Spunbond) Sample ID NMAF01-0409-100-MP
B.	Date of Sample Received		04/09/2020
C.	Date of Commencement of Testing	:	04/09/2020
D.	Date of completion of Testing	:	11/09/2020
E.	Test Method	1:	ASTM F2101
F.	Sample submitted By	1:	Customer
G.	Instrument Used	=   1	Breathing machine, Aerosol Generator, Pressure Probe etc.

### Initial Concentration of Polluting Dust

S. No.	Name Of Test Dust	Unit	<b>Initial Concentration</b>		
2.	PM2.5	(microgram/m³)	24		
3.	PM1.0	(microgram/m³)	6		

### Test Method:

- 1. Air Purifier with filter Tested in a chamber of 10ft x 10ft x 10 ft.
- 2. Fan was powered during operation.
- 3. Concentration is monitored during operation of Air Purifier and also for Control sample without Air Purifier Operation.
- 4. Test Method is in below Table:

PM 2.5	CPCB Volume - 1
PM 1.0	Dust Monitor





(An ISO 9001: 2015, ISO 14001:2015, ISO 45001:2018 CertifiedLaboratory)

Laboratory: A-91, Sector 80, Phase-2, Noida-201301, (U.P)

M.: 09911659800, 09958849764, 07210888634

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S. No.	Time (Hr)	Observation (microgram/m <sup>3</sup> )	Reduction Latching Percentage
1.	Initial	24	*
2.	1	22	0.0
3.	4	26	0.0
4.	8	25	0.0
5.	24	24	0.0

2. P	'M1.0 Removal Perfo	rmance	
S. No.	Time (Hr)	Observation (microgram/m³)	Reduction Latching Percentage
1.	Initial	6	-
2.	1	6	0.0
3.	4	4	0.0
4.	8	5	0.0
5.	24	6	0.0

Remarks: On the basis of above test sample has no Pm 2.5 and PM 1.0 Leaching in the Air from Filter installed in Air Purifier.

#### Notes:

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Checked by





# wau Test House Pyt

### ISO CERTIFIED/ISO ACCREDITED NABL LABORATORY

Nature of the Sample:

Customer Ref. No:

NM Graphene Silver Technology

(NMG-TX)

NMAF06-0409-53

Report No:

20092001

Date Of Sampling:

19.09.2020

Sample issue in Lab:

20.09.2020

Test Started On:

20.09.2020

Test Completed on:

23.09.2020

### SAMPLING DETAILS

Sample Packing & Marking:

Plastic Box.

Sample Quantity:

100 ml

### TEST RESULTS:

Modified ISO 18184:2019 Tested method of Three sample against SARS-CoV-2 (COVID-19) at One contact Time of our sample.

Test Virus	Contact Time	Sample ID	Virus Titer (TCID ∞ per Carrier)	Mean Virust Titer (TCIDso per Carrier)	Mean Log10 Virus Titer (TCID <sub>50</sub> per Carrier)	Log <sub>10</sub> Reduction	Percent Reduction
		100	2.31E+06	17-15-2-15			
1	Time Zero	Control	7.39E+06	3.71E+06	7.13	N.A.	N.A.
		sample	1.45E+06				
			7.32E+05	5 4.79E+05			99.99% Kill Rate: 11 second (99%)
	30 Min.	Control sample	4.52E+05		6.18	5.15	
			2.55E+05				
			2.24E+03				
		Rigid Sample-1 (Test)	2.25E+03			3.25	99.57% Kill Rate: 12 second (99%)
SARS-CoV-2 (COVID-19)			2.38E+03				
(COAID-16)			1.93E+04	3.71E+05	3.98		
			7.29E+05	3.712.00			
			7.23E+03				
			2.65E+03				

\* TCID50 Tissue Culture Infectivity Dose at the 50% Endpoint

\* TCIDs Tissue Culture Infectivity Dose at the 50% Enapoint

\* Logio and Percent Reductions for Control sample at 30Min calculated relative to Control sample immediately upon inoculation (Time Zero)

\* Logio and Percent Reduction for the three Test sample at 30 Min calculated relative to Control sample mean viral titer at 30 Min.

\* Logio and Percent Reduction for the three Test sample at 30 Min calculated relative to Control sample mean viral titer at 30 Min.

\* SAMPLE SIZE: DIAMETER = 4.80 +/= 0.1CM.\* PRE INCUBATION C: SARBOURAUD GLUCOSE AGAR.\* PRE INCUBATION D: SARBOURAUD

DILUTE AGENT FOR INOCULATION: PHYSIOLOGIC SALT SOLUTION. INCUBATION: 30 DEG C. SPECIMEN PREPAR

MIN @ 134 DEG C

Reviewed by

\*\*End of Report\*\*

Authorized signatory by Ravi kant Pathak

The results listed refer only to tested sample & applicable parameters endorsement of product is neither inferred.

Total liability of our lab is limited to the invoiced amount & sample will be destroyed after 90 days from the late of test report

unless specified otherwise. The test report is not to be reproduced wholly or in part & cannot be used as evidence in the court of law & should not be used in any advertising media without our special permission in writing.

The test report refers to the sample submitted to us & not drawn by Himway Test House Pvt. Ltd. unless mentioned otherwise.

Regd. Office: Plot No. 327, Street NO. 02, Santi Nagar (Nandgram) Ghaziabad-201003 Mob.: 8448128057, 8448128058 | Email: himwaytesthouse@gmail.com, Visit us: www.himwaytesthouse.com





# uau Test House Pyt

### ISO CERTIFIED/ISO ACCREDITED NABL LABORATORY

Nature of the Sample :

NM Graphene Membrane

Report No:

20092002

19.09.2020 **Date Of Sampling** 

20.09.2020

Customer Ref. No:

NMAF01-0409-106

Sample issue in Lab Test Started On

20.09.2020

Test Completed on

23.09.2020

### SAMPLING DETAILS

Sample Packing & Marking:

Sample Quantity:

Plastic Bag

1 Meter

#### TEST RESULTS:

Modified ISO 18184:2019 Tested method of Three sample against SARS CoV-2 (COVID-19) at One contact Time of our sample.

Test Virus	Contact Time	Sample ID	Virus Titer (TCID ∞ per Carrier)	Mean Virust Titer (TCIDse per Carrier)	Mean Log10 Virus Titer (TCID <sub>50</sub> per Carrier)	Log <sub>10</sub> Reduction	Percent Reduction
		The same	2.33E+06				
	Time Zero	Control	7.38E+06	3.70E+06	7.10	N.A.	N.A.
		sumple	1.40E+06				
			7.29E+05			5.12	99.99% Kill Rate: 13 second (99%)
	30 Min	Control sample	4.49E+05	4.76E+05	6.15		
			2.51E+05				
			2.20E+03				99.57% Kill Rate: 15 second (99%)
SARS-CoV-2			2.22E+03				
(COVID-19)		Rigid	2.35E+03			3.22	
	30 Min	Sample-1	1.90E+04	3.68E+05	3.94		
		(Test)	7.27E+05	3.002,00	0.51		
			7.20E+03				
			2.62E+03				

\* TCIDso Tissue Culture Infectivity Dose at the 50% Endpoint

\* Logio and Percent Reductions for Control sample at 30Min calculated relative to Control sample immediately upon inoculation (Time Zero)

\* Logio and Percent Reduction for the three Test sample at 30 Min calculated relative to Control sample mean viral titer at 30 Min.

\* SAMPLE SIZE: DIAMETER = 4.80 +/= 0.1CM.\* PRE INCUBATION C: SARBOURAUD GLUCOSE AGAR.\* PRE INCUBATION D: SARBOURAUD GLUCOSE BROTH

DILUTE AGENT FOR INOCULATION: PHYSIOLOGIC SALT SOLUTION. INCUBATION: 30 DEG C. SPECIMEN PREPARATION BATION 15 MIN @ 134 DEG C

\*\*End of Report\*\*

Authorized Signatory by Rayi Kar

The results listed refer only to tested sample & applicable parameters endorsement of product is neither

Total liability of our lab is limited to the invoiced amount & sample will be destroyed after 90 days from the unless specified otherwise.

The test report is not to be reproduced wholly or in part & cannot be used as evidence in the court of law & should not be used in any advertising media without our special permission in writing.

\* The test report refers to the sample submitted to us & not drawn by Himway Test House Pvt. Ltd. unless mentioned otherwise.

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**Report No.** (6720)308-0582 **Page No.** 04 of 04

### TEST RESULTS

TESTED AGAINST: Influenza A Virus (H1N1): ATCC VR-1469

Sample: Fabric – Brown	Log Value
Reference Specimen Immediately after inoculation	5.2
Reference Specimen after 2 Hrs. contact time (of inoculation)	0.8
Antiviral Activity Log (Avg. Reduction)	4.4
Total Viral Activity Reduction (%)	99.997 %

-----END OF TEST REPORT-----



### THE SOUTH INDIA TEXTILE RESEARCH ASSOCIATION

### **CENTRE OF EXCELLENCE FOR MEDICAL TEXTILES**

### Physical, Chemical & Biological Testing Laboratories

(ISO/IEC 17025 NABL ACCREDITED)

13/37, Avinashi Road, Aerodrome Post, Coimbatore - 641 014, INDIA

Test Report No : M2001990 Report Date : 19-09-2020

Ref: dt 05.09.2020

99,99999%

7 Log Reduction

<u>Test Name</u>: Antibacterial activity - Quantitative - ISO 20743

**Test Condition:** 

Test Organisms Used : Staphylococcus aureus ATCC 6538, Klebsiella pneumoniae ATCC 4352

 $\begin{array}{lll} \text{Sample size / Volume} & : 0.4 \pm 0.05 \text{ g} \\ \text{Method of sterilization} & : \text{Steam (Autoclave)} \\ \text{Media used} & : \text{Nutrient agar} \\ \text{Dilution medium used} & : \text{SCDLP medium} \\ \text{Method of plating} & : \text{Spread plate method} \\ \end{array}$ 

 $\begin{array}{ll} \hbox{Inoculum / plate} & : 0.1 \ \hbox{mL} \\ \hbox{Incubation conditions} & : 37^0 \hbox{C for 24 h} \\ \end{array}$ 

Observation:

	M2001990-1 Described by the customer: Solid: NMAF01-0409-102 Sample		
	Staphylococcus aureus ATCC 6538	Klebsiella pneumoniae ATCC 4352	
Inoculated bacterial concentration (CFU / mL)	2.5 x 10 <sup>5</sup>	2.4 x 10 <sup>5</sup>	
Difference of extremes for three control specimens initial (Criteria = Not more than 1.0)	0.09	0.08	
Difference of extremes for three control specimens final (Criteria = Not more than 1.0)	0.01	0.125	
Difference of extremes for three antibacterial test specimens initial (Criteria = Not more than 1.0)	0.017	0.019	
Difference of extremes for three antibacterial test specimens final (Criteria = Not more than 1.0)	0	0	
Growth value (F)	2.084	2.088	
Growth value of (G)	-5.391	-5.373	
Antibacterial activity value	7.475	7.461	
Measuring method of bacterial concentration	Absorption Method	Absorption Method	

**Result:** The Test sample showed 7.475 antibacterial activity against *Staphylococcus aureus* ATCC 6538 and 7.461 antibacterial activity against *Klebsiella pneumoniae* ATCC 4352 when tested for ISO 20743 - Absorption Method

Verified by Authorized Signatory



Report No.: CEG/GA/20-21/03638

CEG Tower, B - 11 (G), Malviya Industrial Area Jaipur - 302017, Rajasthan, INDIA Tel.: 91-141-4046599, Fax: 91-141-2751806 info@cegtesthouse.com | www.cegtesthouse.com CIN: U73100RJ2005PTC020304

Date: 18/08/2020

### RESULTS

				E.coli	S. aureus						
Sr. No.	Size of Sample (cm²)	Contact Duration (Mins)	Number of bacteria inoculated at 0 min (cfu)	bacteria noculated at 0 min (cfu)  bacteria observed after contact duration  Bacterial Reduction		Number of bacteria inoculated at	Number of bacteria observed after contact	Bacterial Reduction			
				(B)	(cfu) (A)	Log		0 min (cfu) (B)	duration (cfu) (A)	Log	%
01.	.01	5	230000	Nil	5	99.999	150000	12000	1	92.00	
02.	01	10	230000	Nil	5	99.999	150000	Nil	5	99.999	
03.	01	30	230000	Nil	5	99.999	150000	Nil	5	99,999	

CFU=Colony Forming Unit | Percentage Reduction = (B - A/B)×100

Remark: The above tested product having antibacterial efficacy against gram-positive & gram-negative bacteria.

\*\*End of the Report\*\*

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Pay 18/103/2020 Checked by

#### Note:

- Total liability of this laboratory is limited to the invoiced amount.
- The results listed refer only to the tested sample and applicable parameters. Endorsement of Product is neither inferred nor implied.
- This Test Report shall not be reproduced wholly or in part and can not be used as an evidence in the court of law without written approval of M/S CEG TH & RC
- From the date of issue of test report, the sample shall be stored, for 1 month in case of non perishable items, upto 1 year for pharma sample, unless otherwise specified in applicable standards/regulatory requirement.
- Sample(s) not drawn by M/S CEG TH & RC, unless specified in the report.





method.

### THE SOUTH INDIA TEXTILE RESEARCH ASSOCIATION

### CENTRE OF EXCELLENCE FOR MEDICAL TEXTILES

### Physical, Chemical & Biological Testing Laboratories

(ISO/IEC 17025 NABL ACCREDITED)

13/37, Avinashi Road, Aerodrome Post, Coimbatore - 641 014, INDIA

Test Report No : O2000605 Report Date : 10-08-2020

Ref: DT 01.08.2020

Bacterial filtration efficiency - ASTM F 2101	O2000605-1 Sample Particulars: Sample ID- NMAF01-0409-056 -MP
Test Condition[s]	· ·
Test organisms used	Staphylococcus aureus ATCC 6538
Inoculum size	5 x 10 <sup>5</sup> CFU/mL
Media used	Tryptic soya agar
Dilution medium used	Peptone water
Incubation conditions	37 <sup>0</sup> C for 24 h
Results	2
Area of test specimen exposed	Fabric 10 x 10 cm
Sample exposure side	Face side
Flow rate of aerosol	28.5 L / min
Mean particle size of challenging aerosol	$3.0 \pm 0.3$ micron
Average plate count of positive control	2091
Average plate count of negative control	0
BFE of test specimen (%)	99.4
BFE of test specimen (%)	99.4  aphylococcus aureus ATCC 6538 when tested according to ASTM F 2

- End of Report -

(This is a computer generated report, hence does not require signature.)



### Anti-viral testing Report

Sample Membrane used for preparing mask

Chemical Nature of the material Not informed

Sample Code No provided by Client NMAF01-0409-078

Sample Code given by Seagull BioSolutions SBPL-AV-001

Tests performed 1. Antiviral Test

### **Sample Preparation**

Membrane was cut into one piece of 0.5 sq inch and another piece of 1 sq inch. Both these pieces were dipped in about 1 mL of animal cell culture medium (DMEM) and observed periodically for its ability to soak the medium. It was observed that the membrane soaked the culture medium after 1 mins and 20 sec exposure. At 2 mins, the 90% of the culture medium was recovered from the membrane. This was used for both the tests.

To ensure that all the antiviral component was extracted from the membrane, the membrane was soaked in 1 mL DMSO and incubated for 2 mins. The DMSO component was recovered and used for further testing.

Results: Total number of viable viruses present after exposure to membrane was determined.

No	Description of sample	No.of viruses (TCID <sub>50</sub> )/mL	% killing of the virus
1	MV virus treated with 0.5*0.5 inch membrane and used for infection on Vero cells	15800	99.4
2	MV virus treated with 1*1 inch membrane and used for infection on Vero cells	8890	99.7
3	MV Virus infected vero cells	2810000	850

#### Method

- Actively growing Vero cells (NCCS, Pune) were split into 96 well plates at a density of 10,000 cells/well and allowed to settle overnight in complete medium (DMEM containing 10% Fetal calf serum).
- 2. Membrane was exposed to 1 mL of Measles Vaccine virus in DMEM for 2 mins and the acqueous extract collected. This was diluted 10 folds serially and used to infect animal cells (96 well plate) in quadruplicate (4 wells per dilution).



### Creating Innovative Healthcare Technologies

- 3. Virus Control: 1 mL of Measles Vaccine virus in DMEM was diluted 10 folds serially and used to infect 4 wells/dilution in 96 well plate.
- 4. The plates were incubated for 7 days and observed every 24 hrs for development of Measles virus cytopathic effect.
- 5. At the end of the 7 days, the number of viruses present in the Virus Control and the Measles virus exposed to the membrane was determined using TCID50 method.

### Conclusion

• The membrane appears to reduce the number of viruses by more than 99% following 2 mins exposure.

### Disclaimer:

1. Seagull is not accredited by any regulatory authority for testing materials for anti-viral activity.