SUGU FABRICS & LINING







SUGU FABRICS & LINING

Sugu Fabrics and Linings, established in 2006, is a trusted manufacturer and supplier of needle felt made of FR, Aramid, M-Aramid, Preox, HPA, CPA, PTFE, Polyester, PP, Viscose, ultra-high molecular weight polyethylene fibers, garment accessories such as shoulder pads, under-collar felts, antimicrobial wiping fabrics, orthopedic cast padding, water-soluble film, and an array of other engineered fabrics.

Our factory is located in village Prithla, Faridabad, 30 km away from New Delhi International Airport, while our office is located in South Delhi's Okhla Industrial area."

Quality first, service-oriented, good faith, truth-speaking innovation is our company's purpose. We would like to cooperate with all friends to achieve a win-win profit. Potential buyers are welcome to contact us.







NEEDLE PUNCH TECHNOLOGY-ADVANCED MATERIAL

Flammability test: ISO 6941

1	Meta-aramid/Preox (60/40)	No sign of melting, dripping. No after flame and no afterglow	Level 4
2	FR viscose / Meta-aramid (60/40)	No sign of melting, dripping. No after flame and no afterglow	Level 4
3	FR viscose/Para-aramid (60/40)	No sign of melting, dripping. No after flame and no afterglow	Level 4
4	FR viscose/Para-aramid/low- melt (50/34/16)	No sign of melting, dripping. No after flame and no afterglow	Level 4

Contact heat: ISO 12127-part 1@ 100°C

Meta-ara	mid/Preox	FR viscos	se /Meta-	FR visco	se/Para-	FR viscos	e/Para-
(50/40)		aramid (60/40)	aramid	(50/40)	aramid/l	ow-melt
						(50/34/1	6)
GSM	Performance	GSM	Performance	GSM	Performance	GSM	Performance
(g/m2)	level (s)	(g/m2)	level (s)	(g/m2)	level (s)	(g/m2)	level (s)
100	32.8	100	22.5				
120	38.1	115	32.9				
125	43.8	140	36.5	90	22.9	110	35.0
130	47.2	160	40.9	90	22.9	110	25.9
140	55	180	43.1				
170	64						

All samples passed level 1 as Threshold time was ≥ 15 s at 100°C

Radiant heat: ISO 6942 @ 20 KW/m2

	Salam near 150 0512 & 20 KHz/III2								
Meta-ara (60/40)	amid/Preax	FR visco aramid	se /Meta- (60/40)	FR visco aramid	se/Para- (60/40)	FR viscose aramid/lo (50/34/16	w-melt		
GSM (g/m2)	Performance level (s)	GSM (g/m2)	Performance level (s)	GSM (g/m2)	Performance level (s)	GSM (g/m2)	Performance level (s)		
100	67	100	60						
140	78	140	72	90	73	110	85		
170	89	180	93						

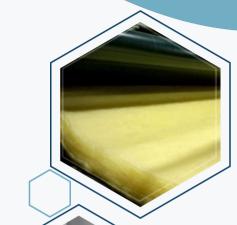
All samples passed level 3 as heat transmission index time ≥50 s and <95 s

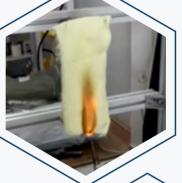
After Treatment Capabilities

- Calendaring
- Padding & Oven
- Lamination-With Nonwoven/Fabric, With release paper, Membrane, Aluminized film.

Final Converting

- Short rolls converting-25 running meters
- Individual width assembly-40 mm-3200 mm
- Different Shapes: CNC Cutter, Hydraulic Press











THERMO ACOUSTIC INSULATION PANELS





Thermal Insulation

Fabris code	OSM	Thickness - Mentioned (max)	Thickness - Tested of 2 Ma pressure (with)	Thermal Resistance (deg C. rid (W)	Thermal Insulation (190)
1	1100	- 50	34.3	0.0178	5.6
2	960	50	23.62	0.9897	6.2
3	790	12	9.52	0.4058	2.6
- 4	1280	- 26	22.37	0.6748	4.2
	1100	28	21.2	0.617	1.9
	1080	26	21.36	0.6718	4.3
1	1680	50	52.22	1.0288	6.6
	1780	50	49.32	1,3168	1.4
9	2280	56	52 66	0.7418	4.7
10	2580		53.81	-1,4738	
- 11	3000	50	54.7	0.7624	4.9

Acoustic

Fabric code	GSM	Thickness - Mentioned (mm)	Sound absorption coeff. (SAC, 6300 Hz)	Noise reduction coeff. (NRC)
1	1100	50	0.911	0.31
2	950	50	0.881	0.2
3	750	12	0.822	0.16
4	1200	25	0.943	0.39
5	1100	25	0.897	0.28
6	1000	25	0.846	0.21
7	1660	50	0.977	0.43
8	1700	50	0.815	0.5
9	2200	50	0.972	0.58
10	2550	50	0.984	0.61
11	2000	50	0.958	0.54

Note: Air permeability experiments for samples with thickness of 50 mm are not feasible in physical testing laboratory because the testing instruments are not suitable materials with thickness greater than 10 mm.

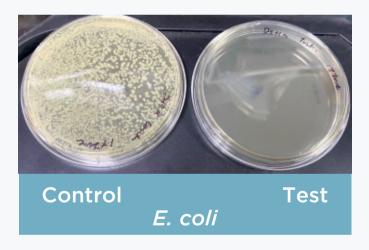
Sound Absorption Test

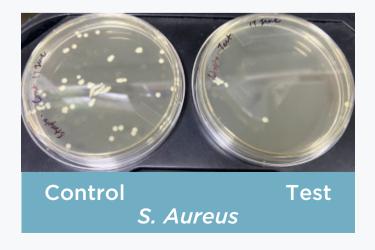
			Sound	absorptio	n coefficie	nt (SAC)					
GSM	1100	950	750	1200	1100	1000	1650	1700	2200	2550	2000
Thickness mentioned(mm)	50	50	12	25	25	25	50	50	50	50	50
Frequency (Hz)											
63	0.131	0.09	-0.057	0.09	0.056	0.053	0.197	0.098	0.019	0.188	0.455
80	0.115	0.03	0.047	0.03	0.036	0.179	0.26	0.12	0.124	0.265	0.37
100	0.109	0.072	0.028	0.072	0.059	0.21	0.255	0.095	0.187	0.242	0.346
125	0.116	0.082	0.05	0.082	0.071	0.194	0.237	0.104	0.206	0.235	0.342
200	0.147	0.115	0.04	0.115	0.077	0.193	0.25	0.168	0.244	0.262	0.342
250	0.159	0.132	0.06	0.132	0.094	0.185	0.248	0.199	0.275	0.292	0.339
315	0.202	0.165	0.068	0.165	0.115	0.19	0.224	0.242	0.308	0.311	0.377
400	0.242	0.17	0.073	0.17	0.135	0.194	0.29	0.295	0.376	0.384	0.425
500	0.274	0.205	0.084	0.205	0.153	0.201	0.386	0.347	0.443	0.458	0.484
630	0.333	0.261	0.117	0.261	0.196	0.23	0.47	0.432	0.537	0.561	0.572
800	0.372	0.33	0.124	0.33	0.233	0.263	0.568	0.51	0.65	0.677	0.684
1000	0.438	0.4	0.154	0.4	0.295	0.309	0.669	0.599	0.752	0.79	0.784
1250	0.5	0.438	0.176	0.438	0.359	0.34	0.742	0.658	0.792	0.864	0.834
1600	0.557	0.546	0.207	0.546	0.451	0.374	0.811	0.709	0.812	0.91	0.861
2000	0.607	0.638	0.251	0.638	0.56	0.457	0.831	0.747	0.825	0.904	0.869
2500	0.623	0.712	0.308	0.712	0.676	0.555	0.797	0.742	0.804	0.862	0.842
3150	0.613	0.735	0.39	0.735	0.782	0.654	0.755	0.725	0.798	0.848	0.834
4000	0.655	0.719	0.494	0.719	0.844	0.716	0.805	0.775	0.863	0.929	0.903
5000	0.714	0.713	0.602	0.713	0.851	0.729	0.889	0.831	0.898	0.957	0.93
6300	0.738	0.774	0.716	0.774	0.825	0.697	0.86	0.815	0.881	0.919	0.9
NRC	0.37	0.35	0.14	0.35	0.28	0.29	0.52	0.47	0.57	0.61	0.61
	Acoustics-C Specimen i And the oth Characteris Results bel	s tested un er paramet tic impedar	der Temper er, Density nce of Air: 4	of Air: 1.29 04.183Pa*	°C Humidi cg/m*3 Ve s/m	ty: 50.0% locity of So	Atmospher und: 345.62	ic Pressure		0Pa	



ANTI-BACTERIAL CLEANING!

Innovative technology meets antibacterial cleaning with our DRY wipes







INHIBIT BACTERIAL GROWTH >95%

Sustainable choice for your health and the environment

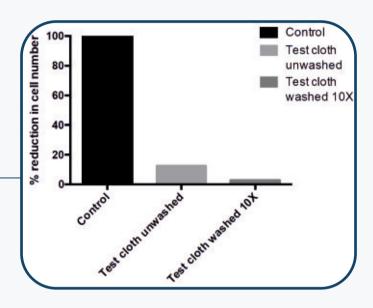
Our antibacterial wipes do the trick!

A healtheir you and a healthier planet all with our antibacterial and reusable wipes. It's the ultimate win-win



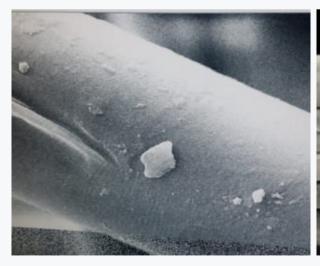
ANTIBACTERIAL NEEDLE PUNCH FABRIC

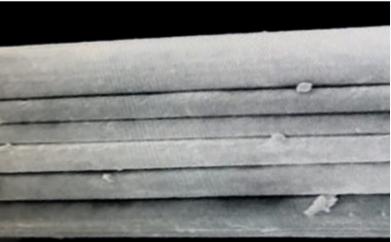
Our fabric exhibits antibacterial properties even after repeated washing. The results shown in the above graph validate our claim



How do we do it?

Through our proprietary method, we enrich our fabric with nanoparticles, as visualized through SEM (scanning electron microscope).





Our antibacterial fabric is versatile and can be converted into dry wipes that are ideal for a range of indoor and outdoor cleaning purposes. These wipes are perfect for cleaning kitchen surfaces, furry pets, children's toys, school furniture, high-touch surface areas such as lifts, handrails, and door knobs, assistive devices, bathroom furniture and fixtures.

Additionally, this fabric can be converted into bed sheets for patient care, and laminated with TPU membrane to make it waterproof. The fabric texture can be enhanced with different prints and embossing patterns, and it is available in six colors to suit We offer fabric cutting and packing



services to produce wipes in a 4-fold form, with embossed sizes of $12cm \times 16cm$, $13cm \times 16cm$, $14cm \times 16cm$, $15cm \times 16cm$, and $16cm \times 16cm$ different needs

We offer perforation options in the Roll form at intervals of 15cm, 20cm, 25cm, and 30cm. The perforation distance is also customizable based on client requirements.

The fabric texture can be enhanced with different prints and embossing patterns, and it is available in six colors to suit different needs





SKIN SENTIVITY TEST



TEST REPORT

LAB NO.: 2003232/1 DATE: 11/09/2020

NAME OF CUSTOMER : M/S. SUVI EXPORTS LLP

ADDRESS : X-48, Okhla Industrial Area Phase-II, New Delhi-110020

REFERENCE : Letter Ref. No. dated September 05, 2020

Kind Attention: Manish Gulati

DATE OF RECEIPT : 05/092020

DATE OF INITIATION : 05/09/2020

DATE OF COMPLETION : 11/09/2020

SAMPLE DESCRIPTION : TEST SAMPLE LABELED AS: -

Sr.No	Description
1.	Non Woven Fabric *VF 100*



Page 1 of 4

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Name of Test and Test Standard:

Invitro Cytotoxicity test

Test Standard:

- 1. ISO 10993-5:2009 (E)- Biological evaluation of medical devices; Tests for in vitro cytotoxicity
- 2. ISO 10993-12:2004 (E) Biological evaluation of medical devices; Sample preparation and reference materials.

Scope of test:

Test for cytotoxicity are designed to determine the biological response of mammalian cells to the test material/ Extract of test material. At the end of the exposure time, the evaluation of the presence and the extent of Cytotoxic effect is assessed. It signifies Biological compatibility of the test material and its potential to cause cell damage.

Cells line and Experimental details:

Cell line L929 - Mouse Connective tissue;

Used for assay for the below stated reasons

Low maintenance

high correlation with specific animal assay

First cell types that attach to implanted medical devices.

Better reproducibility and accuracy of the cytotoxic response.

Passage No. : Cells from PN 168

Cell Culture Medium : Complete MEM medium with 10% FBS

Positive Control : 0.001% SDS (Sodium Dodecyl sulphate) solution

Medium Control / Blank : Complete MEM medium with 10% FBS

: Complete MEM medium

Concentration used : 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% and 100% (neat)

Incubation Condition:

37°C with 5% Carbon dioxide atmosphere

Sample Preparation:

Representative portion of the supplied test sample was used for the assay.

2003232/1 Page 2 of 4

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Sample extraction:

1gm sample was sterilized at 121°C for 15mins to which 10 ml Complete MEM medium was added (0.01g/ml) and incubated at 37°C for 24 hrs.

Assay Principle:

MTT (3-4, 5 dimethylthiazol-2-yl)-2,5 diphenyl tetrazolium bromide Cytotoxicity assay. Test procedure is based on measurement of viability of cells via metabolic activity. Yellow water soluble MTT is metabolically reduced in viable cells to a blue violet insoluble Formazan. The number of viable cells co-relates to the colour intensity determined by photometric measurement after dissolving the formazan in DMSO.

Assay Procedure:

L929 cells seeded in 96 well plates at a concentration of 10,000 cells per 100 µl of MEM culture medium per well were maintained in culture for 24 hours to form a semi confluent layer and were exposed to the test material over a range of concentration. After 24 hours exposure, Formazan formation is determined for each treatment concentration and compared to that determined in growth control.

For each treatment the percentage inhibition of growth is calculated by Viability of cells as per formula -

Viability Percentage = 100 x O. D. 570 nm for extract
O. D. 570 nm for blank

Evaluation criteria:

The lower the viability percentage value, the higher the cytotoxic potential.

The percentage viability of 100% test sample is < 70%, it has cytotoxic potential.

The percentage viability of 100% test sample is ≥ 70%, it is non cytotoxic.

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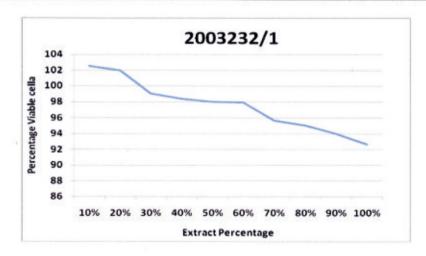
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Results:

2003232/1	Neg. control	Pos. control	Growth control	10%	20%	30%	40%	50%	60%	70%	80%	90%	100%
% cell viability		0.900	100	102.5	101.9	99.02	98.37	97.99	97.97	95.68	95.07	94.01	92.67
P value				0.98	0.95	0.92	0.90	0.85	0.84	0.71	0.70	0.69	0.43



INTERPRETATION:

- 1. For the assay, a concentration range from 10%- 100% was maintained.
- 2. At all concentration set in the assay the sample was found to be Non toxic to the cells
- 3. The values obtained were statistically significant with a p-value <0.05.

CONCLUSION:

Test product labeled as Non Woven Fabric "VF 100" under the extract testing conditions is found to be Non toxic for the cells of cellular culture.

Disclaimer

Any cytotoxic effect can be of concern. However, it is primarily an indication of potential for invivo toxicity and test material cannot necessarily be considered unsuitable for a given clinical application based solely on cytotoxicity data.

For BIOTECH TESTING SERVICES



Dr Shilpa U. Nair Quality Manager (Authorized Signatory)

> 2003232/1 Page 4 of 4

ANTIMICROBIAL EFFICACY OF TOWLETTE SAMPLE



BIOTECH TESTING SERVICES

TEST REPORT

LAB NO.: 2003204/ 1

DATE: 11/09/2020

NAME OF CUSTOMER

: M/S. SUVI EXPORTS LLP

ADDRESS

: X-48, Okhla Industrial Area Phase-II, New Delhi-110020

REFERENCE

: Letter Ref. No. dated September 05, 2020

Kind Attention: Manish Gulati

DATE OF RECEIPT

: 05/092020

DATE OF INITIATION

: 05/09/2020

DATE OF COMPLETION

: 11/09/2020

SAMPLE DESCRIPTION

: TEST SAMPLE LABELED AS: -

Sr. No.	Description	
1.	NON WOVEN FABRIC -APIT	



Page 1 of 7

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Name of Test:

Customer specified Method

US EPA Standard Operating Procedure for Disinfectant Towelette Test: Testing of Staphylococcus aureus, Pseudomonas aeruginosa and Salmonella enterica. The test is based on AOAC Method 961.02 (Germicidal Spray Products as Disinfectants)

Objective:

To study the Antimicrobial efficacy of Towellette samples

Scope:

Method describes the methodology used to determine the efficacy of Towelette-based disinfectants against Staphylococcus aureus, Pseudomonas aeruginosa, and Salmonella enterica on hard surfaces

Test Organisms:

- Staphylococcus aureus (ATCC No. 6538)
- 2. Pseudomonas aeruginosa (ATCC No. 15442)
- 3. Salmonella enterica (ATCC No. 10708)

Additional Test Information:

- 1. Sample size: 18×18 cm
- Towelette with 5 folds like Paper fan; The area of the towelette used for wiping is folded and rotated so as to expose a new surface of the towelette for each carrier.
- 1. No. of wipes used for 10 slides: 1 in 5 folds (front and Back)
- 2. Carrier: Glass Slide Carriers, 25 mm × 75 mm
- 3. Method of Sterilization of sample: UV Sterilization
- 4. Inoculum Carrier: Phosphate Buffered water
- 5. Neutralizer: DE Broth
- 6. Media: Nutrient broth for culture transfer, Trypticase soya agar for Plating and Carrier Enumeration
- Method of wiping: Back and Forth 3 times
- Contact time for wiping carriers (glass slide) surface: 10 seconds
- Contact time given to wipes after use: 15 minutes

Culture Preparation:

Test cultures maintained in Nutrient Broth. Inoculate 10 ml Nutrient broth with 10µl of 24 hr old Culture, vortexed, incubated for 48-54 hours, at 37°C.

2003204/1 Page 2 of 7

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Carrier Inoculation:

Inoculate 80 carriers. 60 carriers for Testing 3 cultures in duplicate. Rest for Viability and Sterility Control. Middle portion of broth culture was diluted, OD adjusted at 650 nm for obtaining 10⁷ CFU/ ml. (1 part culture + 1 part Saline broth). Diluted culture was used for Carrier within 30 minutes.

10 μ l culture was spread on Test carrier placed in Petridish. Culture was placed at the end of the slide and spread with Nichome wire. Dry the carrier in Incubator at 36 \pm 1 $^{\circ}$ C for 30 minutes. Test should be performed within 2 hours of drying.

Two set of 3 carriers are evaluated for Viable count, one immediately prior to conducting Efficacy test and one set Immediately following the test. For determining count, dry carrier is placed in 50 ml conical tube containing 20 ml Letheen broth. Vortexed for 60 seconds for Pseudomonas aeruginosa, 120 seconds for Staphylococcus aureus and Salmonella enterica. After vortexing, serial 10 fold dilutions done in Phosphate buffered Distilled water, 0.1 ml aliquots plated on Trypticase Soya agar. Alternatively Broth can be pooled for 3 carriers and plated.

Test Procedure:

Aseptically removed Towelette, folded lengthwise. Beginning with bottom, folded upwards the top 5 times. This was used to perform the wiping procedure.

Petriplate having carrier is removed. Carrier is visible, wipe the slide back and forth 3 times lengthwise with Towelette for total of 6 passes (3 back and 3 forth). Wiping should be done within 5 seconds. Maintain carrier in Vertical position.

Wiping procedure is repeated with each folding section to expose new surface for wiping each slide. After 5th slide, unfold vertical fold and reverse the towelette so that the used surface of the towelette face inward. Continue wiping on additional 5 slide folding towelette between each slide to expose a new surface.

After the last slide of the test (10th one), Transfer each slide into Primary subculture tube containing neutralizer within 5 seconds. Drain the excess disinfectant from each slide before the transfer. While placing wiped end of the slide should be dropped into neutralizer and diluted serially in duplicates on SCDA plates. Incubate plates for 48 hours at 37°C.

Viability Control:

Dried untreated carriers are separately added to Neutralizer subculture broth. Incubate tubes along with Efficacy test.

Sterility Control:

Sterile untreated carriers before and after drying are placed into separate tubes of Neutralizing broth to perform serial dilution and enumeration

2003204/1 Page 3 of 7

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Controls:

1. Viable counts of Staphylococcus aureus on carrier glass slides before and after drying of inoculum

Sample Description	Count Obtained (CFU)	Log value
VC1 (Before drying)	6.00 x 10 ⁵	5.78
VC2 (Before drying)	6.50 x 10 ⁵	5.81
VC3 (Before drying)	6.30 x 10 ⁵	5.80
VC4 (Before drying)	6.10 x 10 ⁵	5.79
VC5 (Before drying)	6.00 x 10 ⁵	5.78
VC6 (After 30 minutes of drying)	5.90 x 10 ⁵	5.77
VC7 (After 30 minutes of drying)	6.80 x 10 ⁴	4.83
VC8 (After 30 minutes of drying)	6.70 x 10 ⁴	4.83
VC9 (After 30 minutes of drying)	5.20 x 10 ⁵	5.72
VC10 (After 30 minutes of drying)	5.90 x 10 ⁵	5.77

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2. Viable counts of Pseudomonas aeruginosa on carrier glass slides before and after drying of inoculum

Sample Description	Count Obtained (CFU)	Log value
VC1 (Before drying)	5.20 x 10 ⁵	5.71
VC2 (Before drying)	5.80 x 10 ⁵	5.76
VC3 (Before drying)	5.40 x 10 ⁵	5.73
VC4 (Before drying)	5.10 x 10 ⁵	5.70
VC5 (Before drying)	5.60 x 10 ⁵	5.74
VC6 (After 30 minutes of drying)	5.20 x 10 ⁵	5.71
VC7 (After 30 minutes of drying)	4.80 x 10 ⁴	4.68
VC8 (After 30 minutes of drying)	6.10 x 10 ⁴	4.78
VC9 (After 30 minutes of drying)	5.10 x 10 ⁵	5.70
VC10 (After 30 minutes of drying)	4.60 x 10 ⁵	4.66

3. Viable counts of Salmonella enterica on carrier glass slides before and after drying of inoculum

Sample Description	Count Obtained (CFU)	Log value
VC1 (Before drying)	4.80 x 10 ⁵	4.68
VC2 (Before drying)	5.20 x 10 ⁵	4.71
VC3 (Before drying)	5.50 x 10 ⁵	4.74
VC4 (Before drying)	5.80 x 10 ⁵	4.76
VC5 (Before drying)	5.10 x 10 ⁵	4.70
VC6 (After 30 minutes of drying)	5.70 x 10 ⁵	4.75
VC7 (After 30 minutes of drying)	4.10 x 10 ⁴	4.61
VC8 (After 30 minutes of drying)	4.00 x 10 ⁴	4.60
VC9 (After 30 minutes of drying)	4.70 x 10 ⁵	4.67
VC10 (After 30 minutes of drying)	4.90 x 10 ⁵	4.69

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BIOTECH TESTING SERVICES

Results:

1. Viable counts of Pseudomonas aeruginosa on carrier glass slides wiped with Fabric

Sample Description	Count Obtained (CFU)	Log value
T ₁ ,	< 10	<1
T ₂	< 10	<1
T ₃	< 10	<1
T ₄	< 10	<1
T ₅	< 10	<1
T ₆	< 10	<1
T ₇	< 10	<1
T ₈	< 10	< 1
Tg	< 10	<1
T ₁₀	< 10	<1

2. Viable counts of Staphylococcus aureus on carrier glass slides wiped with Fabric

Sample Description	Count Obtained (CFU)	Log value
T ₁	< 10	< 1
T ₂	< 10	< 1
T ₃	< 10	<1
T ₄	1.32 x 10 ²	2.12
T ₅	< 10	<1
Ts	< 10	<1
Т7	< 10	<1
T ₈	< 10	<1
Tg	< 10	< 1
T ₁₀	< 10	< 1

3. Viable counts of Salmonella enterica on carrier glass slides wiped with Fabric

Sample Description	Count Obtained (CFU)	Log value
T ₁	< 10	< 1
T ₂	· < 10	< 1
T ₃	. < 10	< 1
T ₄	< 10	< 1
T ₅	< 10	<1
T ₆	< 10	<1
T ₇	< 10	<1
* T ₈	< 10	. <1
Ta	< 10	<1
T ₁₀	< 10	< 1

2003204/1 Page 6 of 7

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INTERPRETATION:

Sample labeled as NON WOVEN FABRIC -APIT has shown Good Disinfection property towards Staphylococcus aureus, Pseudomonas aeruginosa and Salmonella enterica when tested as per Customer specified US EPA Standard Operating Procedure for Disinfectant Towelettes.

For BIOTECH TESTING SERVICES

BTS SEPTING SE

Dr Shilpa U. Nair Quality Manager (Authorized Signatory)

> 2003204/1 Page 7 of 7



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Customized prints with low moq of 100 MTR

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PRINTED UNDER COLLAR FELT

"GIVING YOUR JACKET'S COLLAR THE SUPPORT IT NEEDS TO STAND UP TO THE COMPETITION."

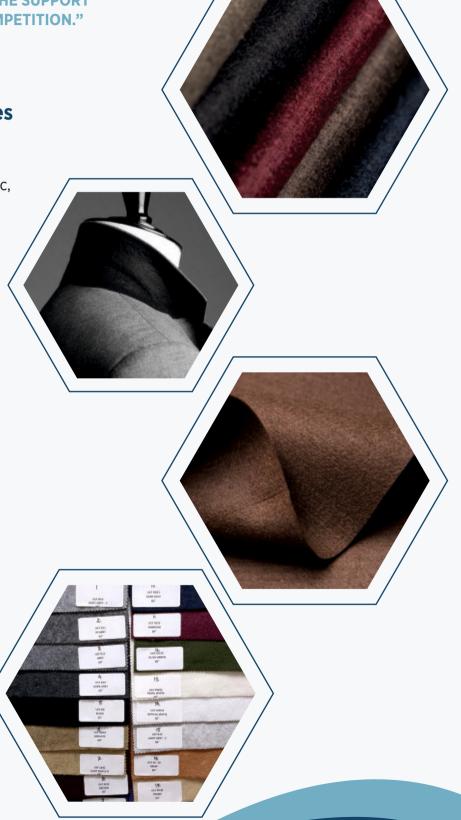
After Treatment Capabilities

- Calendaring
- Padding & Oven
- Lamination-With Nonwoven/Fabric, With release paper, Membrane, Aluminized film.

Final Converting

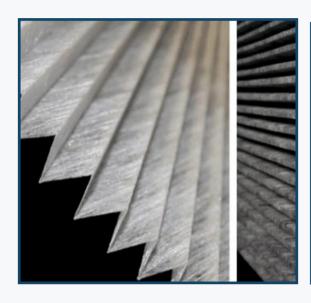
- Non-Toxic, Eco-friendly
- Thickness 10mm-50 mm
- Gsm 500-2000
- Sac .9 (2000 gsm)
- Nrc .6 (2000 gsm)
- Thermal insulation (clo) 4.91
- Maximum Size 72 x 48 inches

Product Name - UCF Width - 60" Length - 30 mtr Weight -180-220 gsm Low shrinkage 20 colors





FILTRATION-FOR DRY FILTRATION



For Ambient Air Filtration, we have engineered media to allow high air flow with the least differential pressure. They are used in air intake filter systems for big compressors, HVAC applications, engine air intake filters and as pre-filters for gas turbines. It's converted into panel, pocket or pleated form to suit the application and system in question.

Model No.: 6535 Color:White

Basic Weight: 110g/m2±10% Thickness: 0.45mm±0.1

Air pressure:1070 (200 pa/dm2/min)

Filter Efficiency for PM2.5:>80% and PM1.0(up to 72%) Application:Mainly Used for Automotive Cabin Air Filters

Model No.: 65354cb

Application: element air cleaner filter

Material: Polyester Process : Needle Punch Area Weight (ASTMD6242) Thickness (150 9073-2)

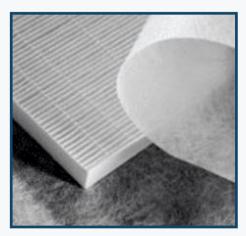
Air Permeability (ASTMD737@200Pa)

Tensile Strength (ASTMD5035)

Bursting Strength(ASTMD3786)

Application

gm/m2 275±10%
mm 2.5±10%
l/dm2/min 700±20%
MD(Kgf/50mm) >=25
co(Kgf/SOmm) >=15
Kg/cm2 >=15
Engine Intake Air Filter









Water Soluble Film/Pouch

Chemical Composition:-

The Product is composed of 65% to 80% of Polyvinyl Alcohol and remaining 20% to 35% of compound plasticizers and organic additive.

All the Substances are Water Soluble and Biodegradable Physical and Chemical Properties:-

Appearance- Solid and translucent, Colour: Natural, red,

yellow, blue, etc., and

Smell: slightly Fermented Odor,

Melting Point — 130°C to 135°C

Flammability — Burn at about the rate of paper

Effect of organic solvent — Resistant to most solvent

Decomposition temperature, More than 183°C

Explosion Danger —No

PH value — 6 to 8 (4wt% water soluble)

Specific Weight -1.2 - 1.3g/cm3

Moisture content — Varies according to the temperature and humidity Solubility in water — 25°C, dissolves within 60 seconds with stirring

Equilibrium moisture content (RH 50— 60) -6.8%

Embroiderry,Chemical

s Packing, Mold releasing agent

Thickness 35 um,45 um,55 um,65 um SIZE 100MM-1400MM

HYGIENIC, CUSTOMIZABLE, ECO-FRIENDLY, HEAT SEALING









For embroiderry

For Flush Tank

HYGIENIC, CUSTOMIZABLE, ECO-FRIENDLY, HEAT SEALING





How do we do it?

Production is professionally managed as per industry 4.0 standards. The human resource is also placed out as per industry 4.0 standards starting from the managers, the shift in charge, machine operators and workers assisted by a maintenance team.





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