

Design and Development of Surgical Gown using Red Seaweed Extract for Hygienic Textiles

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ABSTRACT

The drastic growth in the area of technical textiles and their end-use in healthcare and hygienic textiles create numerous <mark>oppo</mark>rtunities for the application of nove<mark>l fin</mark>ishing agents. Novel finishing agents with improved functionality act as natural antimicrobial agents will help to reduce the adverse effects caused by microbes and it is used for several end applications particularly in the barrier material and infection control. The present study was to develop the surgical gown and investigate the presence of bioactive compounds present in the Acanthophora Spicifera seaweed and analysis the antioxidant and antimicrobial activity of the extract and the treated fabrics using DPPH scavenging activity, disc diffusion, and ENISO20645 test methods. The extractobtained from red seaweed is applied to the bamboo fabric using the pad-dry-cure method for making surgical gown material. The test result shows that the maximum antibacterial activity of the 28mm inhibition zone was observed and the maximum antioxidant inhibition percentage of 77 ±0.17% for both extract and treated fabrics. The presence of bioactive compounds such as flavonoids, tannins, phenols, saponins, and Anthocyanins in the herbal extract was analyzed by using Thin Layer Chromatography (TLC) and the functional group present in the herbal extract was analyzed by Fourier Transform Infrared Radiation Spectroscopy (FTIR). The physical and comfort properties like air permeability, tensile strength, tearing strength, and wicking behaviour of the treated fabrics were decreases compared to untreated fabrics. Thetreated fabrics are used for a variety of medical applications such as gloves, surgical drapes, wound healing, operation room table covers, and face masks for hygienic and healthcare textiles.

KEYWORDS:*Red seaweed, Antioxidant activity, Antibacterial activity, Bioactive compounds, Comfort properties.*

I. INTRODUCTION

Medical textiles have a numerous opportunities for many textile manufacturers, and the textile materials made of knitted, woven, non-woven and composite have been used for the different health care and medical applications [1]. Medical applications can be categorized in to implantable and non-implantable applications. Implantable materials such as surgical suture, artificial skin, artificial ligament, and artificial cartilage. Non-implantable materials such as gauze, pressure bandages, surgical gowns, surgical masks, bed covers, upholstery fabrics, nappies, and face masks have to attain the highest standard properties [2]. Hygienic textiles help to prevent infection control and preserve health for human

life. Inorder to facilitate hygienic products made of textile materials can be used for personal life and healthcare properties.An numerous growth in medical textiles helps to get better comfort nature for patients and end-users [3]. Most of the hygiene products are disposable in nature due to environment friendly and the important prerequisites for successful healthcare products are durable, non-toxicity, mechanical properties, elasticity, non-allergic and biocompatibility. Nowadays, most of the textile products fall in the medical and health care sector is disposable and remaining textile products can be reused after suitablesterilization/cleaning.

Nonwovens are most widely used in the field of non-implantable devices and hygiene products. The development of updating technology and novelty fibres helps to produce compatible, cheaper, pioneering and biodegradable products. Most predominantly used fibres such as cotton, polypropylene, polysaccharides, viscose, polyester, hollow fibre, and elastomeric fibres are used inmedical textiles[4].

Seaweed is a marine plant that is found in every sea or ocean and may belong to one of several groups of multicellular algae. Seaweeds consist of three different groups based on thallus colours such as red, green, and brown seaweeds. Marine macroalgae have a brilliant source of bioactive compounds with various functional activities such as antifungal, anticoagulant, antiviral, antitumor, antibacterial, and anti-inflammatory properties. Seaweeds are used as medicine in older daysand eaten by people in coastal areas. Trace minerals found in seaweeds are regularly used as herbal medicine to conquer inflammation and curing of various infectious diseases like allergy, cancer, influenza, common cold, and tuberculosis. The extracts derived from the seaweed scan be used as a good nutrient for food supplements and act as an anti-inflammatory agent to cure immunologic disorders and allergic diseases in the pharmaceutical industry[5].

Red algae have various bioactive compounds such as phenols, terpenoids, ketones, alkanes, amino acids, phlorotannins, acrylic acid, and saponins [6,7]. The Acanthophora Spicifera species have excellent anti-microbial, anti-oxidant, anti-fouling, ultraviolet-screening anti-tumor agents, and anti-allergy properties compared to other seaweeds. All seaweeds have a phenolic compound in its functional group can act as an excellent antioxidant and antimicrobial activities. Largest phenolic compounds in seaweeds constitute flavonoids that identify the wide band of biological activities including anti-oxidant, anti-viral, and anti-fouling properties[8]. Steroids, terpenoids, and saponins present in the red seaweed have higher antioxidant activities and also the secondary metabolites present in all seaweeds have а higher potential for pharmaceutical application[9].In this study, the extract obtained from red seaweed applied to the bamboo fabric used for making a surgical gown for hygienic textiles.

II. MATERIALS AND METHODS

The sample of Acanthophora Spicifera of edible red seaweed hasa good amount of nutrients, anti-oxidant, and antibacterial activities. They were freshly collected from thonidurai coast of mandapam and were rinsed in seawater and packed in aseptic bags and brought to the laboratory for further processing. The bamboo fabric(4 meters) was purchased fromSri Karthikeyan Textiles, Karur. The fabric parameters are listed as shown in Table 2.1.The chemicals such as Methanol, pectinase and cellulase enzyme, sodium phosphate buffer solution, citric acid,2, 2-Diphenyl-1- picrylhydrazyl, chloramphenicol, ascorbic acidand nutrient agar of laboratorygrade have purchased from the Sri Mahalakshmi Scientific Company, Coimbatore.

TABLE 2.1 FABRICPARAMETERS

Parameters	Bamboo Fabric			
Structure	Plain			
Picks per Inch	88			
Ends Per Inch	100			
Yarn Count	40Ne			
GSM	140			
	ParametersStructurePicks per InchEnds Per InchYarn Count			

Extraction of Red Seaweed

A 30g of powdered seaweed sample was filtered in a Whatman No.1 filter paper thimble was placed into an extraction chamber. The extraction chamber was then connected to a flask containing 300ml of methanol with a ratio of 1:5 (Seaweed: Methanol), then stirred with a magnetic mixer for 3 hours, and the extraction was filtered using Whatman No.1 filter paper. The constant heat source was supplied for this procedure (50°C) for 24 hours until the was clear. All the extracts extract were concentrated under reduced pressure using a rotary evaporator at a temperature of 60°C. The final concentrate after extraction was dissolved in methanol and stored at 4°C for further process as shown in Fig. 2.1[10].



Fig 2.1: Herbal extract of Acanthophora Spicifera(Red seaweed)

Enzyme Scouring

The fabric was immersed in the pectinase solution with 0.05M sodium phosphate buffer at 55° C,M:L ratio of 1:50 for 1 hour. After treatment, the temperature was raised to 100°C for 10 min to stop the enzyme activity. The fabric was washed with hot water followed by cold water and dried[11].

Enzyme Bleaching

The scoured bamboo fabric was taken and placed in a 0.1 M sodium phosphate buffer solution (pH 7.0). Add cellulase to it and mix well and incubate at 50°C for3hours. Inactivate the enzyme by boiling water for 5 min. The fabric was washed thoroughly in tap water and then in distilled water, dried in air for 1hour[12].

Finishing Process Wet Dip Method

The method of dipping is done by immersing the fabric material in the treatment bath containing the *Acanthophora Spicifera* extract for 5 minutes. The fabric was removed from the bath and squeezed gently and dried in air. The fabric is finished using a wet dip method as a continuous process. The extract made from *Acanthophora Spicifera* is applied to the fabric for finishing and citric acid is used as a binder for fixation of the extract to the bamboo fabric. The fabric is dried for 30 minutes at 60°C[13].

Assessment of bioactive compounds using Thin Layer Chromatography(TLC)

The TLC technique was used to separate non-volatile mixtures. The investigationwas conducted using a sheet of aluminum foil coated with a thin layer of silica gel. The activate TLC plates were applied to*Acanthophora Spicifera* extract with the help of capillary tube at a 1/2 inch apart from the lower edge of TLC plate, and the TLC plate was kept in a developing chamber containing suitable solvent system for a specific time until the developing solvent reaches the top of the upper edge of TLC plate. The TLC plate was taken out of the developing chamber, dried and the solvent front is marked by lead pencil [14]. The compounds have a higher affinitythat will move slowly towards the stationary phase and the compounds with lower affinitywill travel fast. In the separation process, the bioactive compounds present in the extract appear as spots at respective levels on the TLC chromatoplate by using suitable spraying reagent for the presence of specific compounds. The visualized spots of the components in the chromatoplate are marked and the Rf value of each spot is calculated by the formula:

 R_f =Distance travelled by sample (cm) (1) Distance travelled by solvent (cm)

Investigate the functional group of red seaweed extract using Fourier Transform Infrared Radiation Spectroscopy (FTIR)

The functional groups present in the red seaweed extract shows the presence of bioactive compounds isrecognized using FTIR spectroscopy. A peak value arisesin the form of spectral curves at various wavelengths with the help of infrared radiation was measured. FTIR absorption spectra can be varied according to the extract solution and the spectral curve formed with respect to different wavelengths from 400-4000cm⁻¹ was recorded using a Thermo ScientificNicolet iS10 FTIR Spectrometer, USA. Take 3g of powdered seaweed extract added with18g of potassium bromide was filled using mortar with the ratio of 1:6. The large crystal present in the mixtures is broken down in to smaller ones using pestle are grinded thoroughly. Then place the mixture in the pellet diesefficiently and pressure is applied to form pellets. The pellet is placed in the FTIR sample pellet holder. The changesin the functional group existing in the red seaweed extract were revealed during this analysis [15].

Assessment of the AntioxidantActivityDPPH radical scavenging activity

Free radical scavenging activity was measured by 2,2-Diphenyl-1-picryl hydrazyl (DPPH) according to the method of Yen and Chen (1995)[16]. Briefly, a 4.0ml aliquot of the test sample was added to 4.0ml of 0.16ml DPPH solution. The mixture was shaken vigorously, then left to stand at room temperature for 30 min in darkness. Changes in the absorbance of the samples were measured at 517 NM using a UV spectrophotometer. The ability

to scavenge the DPPH radical was evaluated using the following equation:

Scavenging effect (%)

=1- $(A_{\text{Sample}}-A_{\text{Sampleblank}})(2)$

A $_{\rm control}$

Where:

A_{control} is the absorbance of the control (DPPH solution without sample),

A_{sample} is the absorbance of the test sample (DPPH solution plus test sample),

A $_{\rm sample\ blank}$ is the absorbance of the sample only (sample without any DPPH solution).

Ascorbic acid was used as a positive control.

Assessment of Antibacterial Activity Disc diffusion method for extract

The antibacterial activity of the extract was evaluated using disc diffusion method. Briefly, 6mm paper disks were impregnated with two different concentrations 100µland 200µl using methanol of seaweed extract and left dried on the disc. Then the discs were applied to the agar plates that were inoculated previously with the test organisms (Staphylococcus Aureus and Escherichia Coli) and incubated for 1 hour. The bacterial plates were incubated at 35°C for 28hours. The chloramphenicol was used as a positive control for two different concentrations (100 and 200 μ g/disc). After incubation, all plates were observed in the zone of inhibition and the diameter of the zones was measured in millimeter. All tests were carried out three times under sterile conditions[17].

ENISO 20645 anti-bacterial method for treated fabrics

The nutrient agar plates were prepared by pouring 15ml of agar media into sterile petri plates. The plates were allowed to solidify for 5min and the test bacterial culture (*Staphylococcus aureus* and *Escherichia coli*) was inoculated uniformly and dry for 10 minutes. The control and treated fabric were cut into 2.5-3cm size and placed over the inoculated test bacterial cultures, separately. The plates were kept inincubation at 39°C for 24hours. After incubation,the inhibition zone present around the fabric was calculated in millimeters and recorded[18].

Evaluation of fabric properties

The physical and comfort properties of fabric were carried outusing the ASTM standards, and AATCC test methods. The standard atmospheric condition was maintained with a temperature of 27° C, and $65\pm 2\%$ RH. The fabric tensile strength was evaluated according to standard ASTM D5034-95.

The fabric tearing strength was measured according to standardASTM D1424-96. The quantity of air passes per second through one square cm of the fabric was calculated according to standard ASTM D737-04. The degree of colour change and colour staining was evaluated according to standard AATCC61-13. The extraction has a pale red colour and the quantity of colour removed from the treated fabric both in wet and dry conditions was calculated according to standard AATCC TM08-96. The vertical wicking behaviour of the fabrics was evaluated according to standard AATCC 197-11[26].

Development of Woven Surgical Gown

The surgical gown is made on the treated bamboo fabric.The pattern is made according to the standard specificationsgiven for surgical gown. The treated fabric was cut according to the pattern and stitch the cut fabric as the surgical gownis shown in Fig. 2.2.



Fig 2.2: Developed Surgical Gown

A low density seam is given for the gown to attain lower bulkiness. The yoke part is provided in front of the surgical gown.Ropes are made on the back pattern for making the knot in the surgical gown.Comfortably, the surgical gown was developed for hygienic applications.

III. RESULT AND DISCUSSIONS

Assessment of bioactive compounds in Acanthophora Spicifera extractusing TLC technique

The dissolving solvent act as a mobile phase with the help of a suitable spraying agent was used to identify the bioactive compounds present in the *Acanthophora Spicifera* extract are presented in Table 3.1. The precise spot colour and its R_f values produced in seaweed extract are shown in Fig.3.1 and 3.2[19].



Fig 3.1: Analysis of bioactive compounds in Acanthophora Spicifera extract

Bioactive compounds	Mobile phase	Spraying reagent	Spot colour	Rf Valu e
Flavonoids	Ethyl acetate: Butanol: Formic acid (2:2:1)	Aluminiu m chloride	Orange	0.92
Phenols	Chloroform: Methanol (25:1)	Ferric chloride reagent	Blue	0.89
Tannin	Methanol: Water (6:4)	Ferric chloride r <mark>eagent</mark>	Browni sh grey	0.86
Saponins	M <mark>ethan</mark> ol: Water (7:3)	Ferric chloride	Light Orange	0.62
Anthocyanin s	Acetonitrile: Water: Formic acid (2:2:1)	Acetic acid	Pink colour	0.30

Table 3.1 Assessment of bioactive compounds using TLC Technique



Fig 3.2: Presence of (a) Flavonoids (b) Phenols (c) Tannins (d) Saponins (e) Anthocyanins in Acanthophora Spicifera extract

The figure 3.1 and 3.2 shows that the *Acanthophora Spicifera*extract has the presence of five major bioactive compounds such as flavonoids, phenol, saponins, anthocyanins and tannin attributing for their anti-bacterial, antiviral, antitumor, anti-cancer and anti-inflammatory properties[19].

Assessment of functional groups in Acanthophora Spicifera extract using FTIR Spectroscopy

FTIR spectrum was used to identify the functional groups present in the seaweed extract based on the peak value in the region of the infrared spectrum as shown in Fig. 3.3. The prominent peaks occur in 3354.65cm⁻¹, 2919.59cm⁻¹,2364.78cm⁻¹, and 2135.57cm⁻¹ represents the major functional groups such as phenols, amines, alkane and aldehydes present in the seaweed extract[20].The prominent peaks occur at 1757.48cm⁻¹, 1639.74cm⁻¹, and 1417.60cm⁻¹ represents the major functional groups such as carboxylic acid and alkene present in the seaweed extract. The prominent peaks occur in1155.89cm⁻¹, 1000cm⁻¹, and 895.64 cm⁻¹ represents the major functional groups such as ketone, anhydride and alkene present in the seaweed extract as shown in Table 3.2.

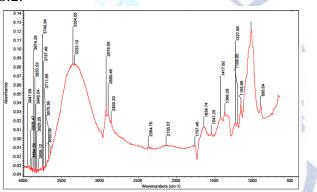


Fig 3.3:FTIR Spectroscopy for Acanthophora Spiciferaextract

Table 3.2FTIR analysis for	Acanthophora
Spiciferaextract with respec	ct to wavelength

			Atra-	
S.	Wavelength	Bond /	Intensity	Functional
No.	(cm ^{- 1})	Stretching	9	Group
1.	3354.65	N-H Stretch	Medium	Phenols and Amines
2.	2919.59	C-H stretching	Medium	Alkane
3.	2364.78 & 2135.57	C-H stretch	Medium	Aldehyde
4.	1757.48	C=O stretching	Strong	Carboxylic acid
5.	1639.74	C=C stretch	Weak	Alkene
6.	1417.60	O-H bending	Strong	Carboxylic acid
7.	1155.89	C=O	Very strong	Ketone
8.	1000	CO-O-CO stretching	Very strong	Anhydride
9.	895.64	C=C bending	Strong	Alkene
10.	694	C=C bending	Strong	Alkene

The peak present in the spectrum has confirmed the presence of functional groups such as flavonoids, anthocyanins, phenols, tannin,and saponins compounds present in the treated fabric. The functional groups present in the seaweed extract are responsible for antibacterial, antiviral, antitumor, anti-cancer,and anti-inflammatory properties.

Evaluation of Antioxidant Activity

The antioxidant activity of *Acanthophora Spicifera* was measured using the DPPH method. The result shows that the antioxidant activity increases with increasing the concentration of the extract and treated fabric sample[21].

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Seaweed Extract

The antioxidant property of *Acanthophora Spicifera* extract was analyzed by the DPPH radical scavenging method. The percentage of scavenging activity present in the red seaweed extract is shown in Table 3.3.

Table 3.3 Antioxidantactivity of the methanol extract

S.No.	Seaweeds		PPH Free Radical Scavenging Activity % Inhibition		
1000	Name	Ascorbic Acid	Aqueous Extract		
1.	inthophora Spicifera	77 ± 0.17	24. <mark>85 <u>+</u> 2.</mark> 87		

The results indicate that the percentage of radical scavenging activity increases, with an increase the concentration of ascorbic acid and aqueous extract. The antioxidant property of the seaweed extract will help to reduce the number of free radicals present in the oxygen species and fast growth of cell membrane present in the skin.

Treated Fabric

The percentage of free radical scavenging activity of treated fabric was analyzed by the DPPH method. The test results are shown in Table 3.4.

Table 3.4 Antioxidant activity of the treatedfabric

S.No.	Treated fabric	DPPH Free Radical Scavengin Activity % Inhibition Ascorbic acid Aqueous Extrac	
1.	Acanthophora Spicifera	76 ± 0.26	23.95 <u>+</u> 1.12

The test result of the treated fabric shows that the percentage of radical scavenging activity was more or less the same for seaweed extract and treated fabric. The antioxidant activity helps to trap the free radial oxygen species and inhibits the cell damage and develops the cell growth present in the skin.

Evaluation of Antibacterial Activity

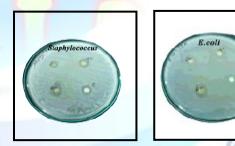
The antibacterial activities of seaweed extract and treated fabric are tested on the standard of theDisc diffusion method and ENISO 20645 test methods. It was determined mainly by two bacterial species such as *Staphylococcus aureus* and *Escherichia coli*. The size of the sampleistaken in the range of 1.5×10^{8} CFU/ml. In both test methods, the result shows that the excellent antibacterial activity was attained against *Staphylococcus Aureus* compared to *Escherichia Coli*.

Disc diffusion method for extract

The test result shows that a maximum of 28 ± 1.5 mm of inhibition zone was occurred at $200\mu g/ml$ for *Staphylococcus Aureus* compared to *Escherichia Coli* as shown in Table 3.5 and Fig. 3.4.The higher inhibition zone was attained due to the presence of bioactive compounds such as phenols, tannins, and saponins attributed to the excellentantibacterial activity of the seaweed extract[17].

Table 3.5 Disc diffusion method of extract

S.No.	Seaweed	Extraction	Concent -ration (µg/ml)	coccus Aureus	Escheric hia Coli(mm)
1.	Acanthophora	Methanol	100	17 <u>+</u> 3.9	13 <u>+</u> 1.3
1.	Spicifera	Methanol	200	28 + 1.5	23 + 2.6



(a)Staphylococcus Aureus (b) Escherichia Coli Fig 3.4: Antibacterial activity of Acanthophora Spicifera extract

ENISO 20645 method for treated fabric

The test result shows that the maximum of 28mm of the inhibition zone was occurring against Staphylococcus Aureus compared to Escherichia Coli as shown in Table 3.6 and Fig.3.5[18]. The treated fabric has a property to inhibit the growth of bacteria to a maximum extent was mainly due to the presence of bioactive compounds such as phenols, tannins, and saponinsattributed to theexcellent antibacterial property.

Table 3.6 Antibacterial activity of treated fabric

G	S. Fabric Seaweed		Zone of Inhibi	ition (mm)
	. Sample		Staphylococcus Aureus	Escherichia Coli
1.	100%Ba mboo	Acanthophora Spicifera	28	25



 (a) Staphylococcus Aureus(b) EscherichiaColi
 Fig 3.5: Antibacterial activity of Acanthophora Spicifera extract treated fabric

The above figure shows that the zone of inhibition was clearly seen on the sides of the treated fabric against *Staphylococcus Aureus Aureus Coli*.

ENISO 20645 method of treated fabricafter washing

The test result shows that after repeated washing cycle of 20 times the treated fabric had an excellent antibacterial activity shown in Table 3.7.

Table 3.7Antibacterialactivity of treated fabric after washing

s.	Fabric	Seaweed	Zone of Inhibition (mm)	
	Sample		Staphylococ cusAureus	Escherichia Coli
1.	100%Ba mboo	Acan <mark>tho</mark> phora Spicifera	26	23

(a) Staphylococcus aureus(b) Escherichia coli

Fig 3.6:Antibacterial activity of Acanthophora Spicifera treated fabric after washing

The detergents used in washing will not affect the treated fabric, because the extract is evenly coated on the interstices of the fabrics and the affinity against detergents is lower and the better inhibition against pathogenic bacteria after repeated washing is shown in Fig.3.6[18].

Analysis thetensile strengthof treated and untreated fabrics

The tensile strength test of both treated and untreated fabrics is tested according to the ASTM D5034-95 standard. The results obtained from the tensile strength test arepresented inTable 3.8[22].

Table 3.8 Tensile strength of the treated anduntreated fabric

Untreated Fabric	Treated Fabric				
Warp way sample	Weft way sample				

S.No.	Strength	Elongation	Strength	Elongation
	(kg)	(mm)	(kg)	(mm)
1.	30.1	14.3	21.3	10.4
2.	32.4	15.4	19.6	10.0
3.	26.1	12.5	19.5	10.2
4.	31.5	14.6	23.1	10.7
5.	28.2	13.5	20.1	10.0
Mean	29.66	14.06	20.72	10.26

The result shows that the tensile strength increases with a decrease in elongation in the untreated fabric compared to treated fabric. In the case of treated fabric, the tensile strength decreases with an increase in elongation of the fabric was mainly due to bio-scoured, bio-bleached, and finishing condition or evenly coated on the surface of the fabric.

Analysis the tearing strength of treated and untreated fabrics

The tearing strength test of both treated and untreated fabrics is tested according to the ASTM D1424-96 standard. The results obtained from the tearing strength test arepresented inTable 3.9[23].

Table 3.9 Tearing strength of the treated and untreated fabric

S.No.	Untreate	d Fabric	Treated	d Fabric	
1	Warp (kgf)	Weft (kgf)	Warp (kgf)	Weft (kgf)	
1.	2.92	3.24	2.79	3.20	
2.	2.86	3.03	2.76	2.89	
3.	2.84	3.52	2.68	3.15	
4.	2.85	3.59	2.83	3.53	
5.	2.72	3.17	2.65	2.78	
Mean	2.83	3.31	2.74	3.11	

The



resultshows that the tearing strength increases for untreated fabric compared to treated fabric. In the case of treated fabric, the bio-scoured, bio-bleached, and finishing treatment deteriorates the tear strength of the fabric from both warp and weft side was mainly due to the carboxylic acidic grouppresent in the extracts that weakens the fibres. The acidic nature of the extracts hydrolyzes the cellulose present in the bamboo fabric more quickly will reduce the tearing strength of the treated fabric compared to untreated fabric.

Assessmentofair permeabilityproperties

The permeability of air is asignificant feature in the performance of treated and untreated fabrics were tested according to ASTM D737-96 standard. The results obtained from the fabric air permeability test are shown in Table 3.10[24].

Table 5.10 All Termeability test			
S.No.	Untreated fabric	Treated fabric]
1.	50.9	49.3	
2.	51.2	49.7	4
3.	52.7	50.5	
4.	53.5	51.9	-
5.	53.4	52.8	-
Mean	52.34	50.84	

Table 3.10 Air Permeability test

The test result shows that the air permeability increases for the untreated fabric compared to treated fabric. In the case of treated fabric due to the absorption of the extract on the interstices between the yarns present in the fabric will decrease the fabric cover factor and pore size will reduce the air permeability. Whereas in untreated fabric, the fabric cover factor and pore size remain the same will increase the permeability of air passed through the fabric.

Assessment of washing fastness oftreated fabric

It is used to provide an indication of the removal of extract present on the surface of the fabric and it will affect the performance and aesthetic value of the fabric. The treated fabric washing with different cycles (5, 10, 15,and20) will have a very good wash fastness after repeated washing. At each stage of washing take place, the grade change for the white and colour fabric was in the range of 3-4 as shown in Table 3.11[25].

Tables. I I washing fastness test			
Cycle of Washing	and the second s	Evaluate the Colour	Evaluate the Staining
	- 7 P * 2	Change Grade	Grade
5	(4	3
10	Treated	4	3
15	fabric	4	3
20		4	3

Table3.11Washing fastness test

The result clearly proved that even absorption of extract takes place in between the intermolecular structure of the yarn present in the treated fabric. The amount of extract absorbed in the fabric was not cleavage between the fibre present in the yarn. It also forms astrong bond between the fibre will lead to uniform uptake of the extract and minimum amount of the extract are removed at the surface of fabric during repeated washing after 20 cycles.

Assessment ofrubbing fastness properties of treated fabric

It is used to provide an indication of the removal of the extract on the surface of the fabric during rubbing on both dry and wet samples. The result obtained from the fabric rubbing fastness test is shown in Table 3.12[25].

Table	3.12	Rubbing	fastness	test
I abic	0.14	Rubbing	Idociic 33	COSC

S.No.	Particulars	Rating
1.	The numerical rating for change	5
	in colour of the dry specimen	
2.	The numerical rating for change	4
۷.	in colour of the wet specimen	4
_	The numerical rating for change	
3.	in colour of the dry - wet	4
1	specimen	
1	The numerical rating for change	
4.	in colour of the wet - dry	4-5
	specimen	2

The result shows that for the dry samples, the rubbing fastness properties are good compared to wet samples. In the wet state of the treated fabric, the extract applied on the surface is slightly removed after repeated rubbing as per the ASTM (08) standard and the grade result in the range of Good to Excellent.

In the dry state of treated fabric shows good fastness property, because of the strong fixation of seaweed extract on the intermolecular structure of the fabric, will increase the affinity between the extract and the fabric pore size.

Assessment of wicking behavior oftreated and untreated fabrics

It is used to find the water absorbency on the strip of the fabric against time in seconds. The results obtained from wickability test of the treated and untreated fabrics are shown in Table 3.13.

	S.No.	Time	Untreatedf	Treated
5.NO.	(Sec)	abric	fabric	
	1.	5	3.6	2.9
	2.	10	3.5	2.8
	3.	15	3.5	2.8
	4.	20	3.4	2.7
	5.	25	3.4	2.6

Table 3.13 Wickability test

The result shows that the treated fabric is absorbed little less compared to untreated fabric because the seaweed extract are absorbed in the intermolecular structure between the yarns will increase the weight of the treated fabric[26].

IV. CONCLUSION

In this study, *Acanthophora Spicifera* seaweed was collected from different sites that were screened for bioactive compounds, antioxidant, and antibacterial activity. The Acanthophora Spicifera seaweed extraction process wascarried out in a Soxhlet apparatus using methanol as a solvent for finishingbamboo fabric inhealthcare and hygienic applications. The bioactive compounds present in seaweed were confirmed by usingthe TLC technique and the functional group present in the seaweed extract was confirmed using FTIR spectroscopy analysis. Antioxidant properties of the red seaweed extract and the treated fabric was confirmed by using DPPH free radical scavenging activity. The results indicate that the presence of phenol, flavonoids, and anthocyanins compounds should be major contributors toexcellent antioxidant activity.

Antibacterial activity of the seaweed extract and the treated fabric was confirmed using disc diffusion and ENISO 20645 test methods. The results indicate that Acanthophora Spicifera extracts exhibit a promising antibacterial activity against Staphylococcus aureus and Escherichia Coli. The maximum zone of inhibition was around28mm was achieved in both extract and treated fabricscompared to untreated fabrics. The treated fabric has better durability after repeated washing and the antimicrobial property remains withstand after 20 washing cycles. The physical and comfort properties like tensile strength, tearing strength, air permeability, and wicking behaviour of the treated fabrics were decreases compared to untreated fabrics. The treated fabrics are used for a variety of medical applications such gloves, surgical drapes, wound healing, as operation room table covers, and face masks for hygienic and healthcare textiles.

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