

RESEARCH ARTICLE



Development and Evaluation of an Eco-friendly Antimicrobial Wound Dressing Using Neem and Turmeric-Coated Coconut Leaf Sheath

Syed Ansar Ahmed^{*1}, Madhuri Vishwanath Swami², Parchande Komal Vitthal³, Monika D. Palimkar⁴, Syed Iqra Naznin⁵, Mujahed Nasir Pathan⁶

¹Associate Professor, Department of Pharmaceutical Chemistry, Indira College of Pharmacy, Vishnupuri, Nanded, Maharashtra, India

²Assistant Professor, Department of Quality Assurance, Indira College of Pharmacy, Vishnupuri, Nanded, Maharashtra, India

³Assistant Professor, Department of Pharmaceutical Chemistry, SVETRI's College of Pharmacy, Pandharpur, Maharashtra, India

⁴Assistant Professor, Department of Quality Assurance, Indira College of Pharmacy, Vishnupuri, Nanded, Maharashtra, India

⁵Assistant Professor, Department of Pharmacology, Aurangabad Pharmacy College, Padegaon, Maharashtra, India

⁶Associate Professor, Department of Pharmaceutics, Supriya College of Pharmacy, Jafrabad, Jalna, Maharashtra, India

Publication history: Received on 6th June 2025; Revised on 2nd July 2025; Accepted on 15th July 2025

Article DOI: 10.69613/rfwxvn48

Abstract: The development of sustainable, biodegradable wound dressings with antimicrobial properties could help in the management of wound care. The aim of this research work was to create a novel wound dressing material using coconut leaf sheath impregnated with neem (*Azadirachta indica*) and turmeric (*Curcuma longa*) extracts. The coconut sheaths underwent pre-treatment involving chemical softening with sodium hydroxide and sodium carbonate, followed by ultrasonic treatment. The medicated formulation incorporated neem extract, turmeric extract, and chitosan as a natural film-former. The antimicrobial efficacy was evaluated against *Escherichia coli* and *Candida albicans* using agar well diffusion, minimum inhibitory concentration (MIC), and time-kill kinetics methods. The medicated gauze demonstrated significant zones of inhibition: 16.42 mm against *E. coli* and 19.85 mm against *C. albicans*. FTIR analysis confirmed successful incorporation of bioactive compounds, showing characteristic peaks for phenolic O-H, aromatic rings, and C-H stretching. The neem extract exhibited superior antimicrobial activity compared to turmeric, with MIC values of 125 µg/mL for both *S. aureus* and *C. albicans*. Time-kill studies revealed complete microbial elimination within 24 hours at twice the MIC concentration for neem extract. The developed wound dressing could be an efficient alternative to synthetic materials, offering natural antimicrobial properties while maintaining environmental sustainability.

Keywords: Antimicrobial wound dressing; Coconut leaf sheath; Neem extract; Turmeric extract; Biodegradable materials

1. Introduction

The rapid advancement in wound care management has necessitated the development of innovative, sustainable materials that can effectively promote healing while minimizing environmental impact [1]. Traditional wound dressings, predominantly manufactured from synthetic materials, have raised significant environmental concerns due to their non-biodegradable nature and contribution to medical waste [2]. Moreover, these conventional materials often lack inherent therapeutic properties, serving primarily as passive barriers rather than active healing facilitators [3]. Natural fiber-based materials have garnered substantial attention in the biomedical field, particularly in wound care applications, due to their biocompatibility, biodegradability, and potential therapeutic properties [4]. Among various natural sources, coconut leaf sheath presents unique structural and biological characteristics that make it particularly suitable for wound dressing applications [5]. The natural architecture of coconut leaf sheath, characterized by its interconnected fiber network, provides excellent mechanical strength while maintaining necessary porosity for wound healing [6].



Figure 1. Coconut leaf sheath

* Corresponding author: Syed Ansar Ahmed

The incorporation of medicinal plant extracts into natural fiber matrices represents a synergistic approach to enhance wound healing properties [7]. Traditional medicine systems have long recognized the therapeutic potential of various plants, particularly neem (*Azadirachta indica*) and turmeric (*Curcuma longa*) [8]. Neem contains several bioactive compounds, including nimbidin, azadirachtin, and quercetin, which demonstrate significant antimicrobial, anti-inflammatory, and wound healing properties [9]. The antimicrobial activity of neem is attributed to its ability to disrupt bacterial cell membranes and inhibit essential bacterial processes [10].

Similarly, turmeric has been extensively studied for its therapeutic properties, with curcumin being its primary active component [11]. Curcumin exhibits powerful anti-inflammatory, antimicrobial, and antioxidant properties, making it valuable in wound healing applications [12]. Recent research has shown curcumin's ability to modulate various molecular pathways involved in wound healing, including inflammation, matrix remodeling, and tissue regeneration [13].

The combination of natural fibers with medicinal plant extracts offers several advantages over conventional wound dressings [14]. These include sustained release of bioactive compounds, enhanced biocompatibility, and reduced environmental impact [15]. Additionally, the use of agricultural byproducts like coconut leaf sheath promotes sustainable resource utilization while providing economic benefits to agricultural communities [16].

2. Materials and Methods

2.1. Materials

2.1.1. Plant Materials

Coconut leaf sheaths were collected from local agricultural areas and authenticated by botanical experts [17]. Fresh neem leaves were sourced from certified suppliers and verified for species authenticity and quality [18]. Turmeric rhizomes were obtained from authenticated agricultural sources and evaluated for curcumin content [19]. The study employed chitosan with a molecular weight range of 100-300 kDa and deacetylation degree exceeding 75% [20]. Additional materials included analytical grade glacial acetic acid, sodium hydroxide, and sodium carbonate [21].

2.1.2. Microbial Strains

The antimicrobial evaluation utilized standard strains obtained from American Type Culture Collection (ATCC): *Staphylococcus aureus* (ATCC 25923), *Escherichia coli* (ATCC 25922), and *Candida albicans* (ATCC 10231) [22]. The strains were maintained according to standard microbiological protocols [23].

2.2. Methods

2.2.1. Preparation of Coconut Leaf Sheath

The preparation process commenced with thorough cleaning of raw sheaths under running water to remove surface contaminants [24]. The cleaned sheaths underwent chemical softening through immersion in a solution containing 5% w/v sodium hydroxide and 2% w/v sodium carbonate for 24 hours at ambient temperature [25]. Following chemical treatment, the sheaths were subjected to multiple rinse cycles with distilled water until achieving neutral pH [26]. Ultrasonic treatment was performed at 45°C for 15 minutes to enhance fiber separation and remove residual impurities [27]. The processed sheaths were dried at 40°C for 6-12 hours in a forced-air circulation oven [28].



Figure 2. Chemical softening with NaOH and Na₂CO₃

2.2.2. Extract Preparation

The preparation of medicinal plant extracts followed standardized protocols to ensure consistency and potency [29]. For neem extract preparation, dried leaf powder underwent Soxhlet extraction using 70% ethanol as the extraction solvent [30]. The extraction process continued for 48 hours to ensure complete extraction of bioactive compounds [31]. The resulting extract was concentrated under reduced pressure at 40°C using a rotary evaporator and standardized to contain a minimum of 2% w/w azadirachtin [32].

Turmeric extract preparation involved processing dried rhizome powder through extraction with 95% ethanol [33]. The extraction process incorporated temperature control at 50°C for optimal curcuminoid extraction [34]. The extract underwent concentration and standardization procedures to achieve a minimum curcuminoid content of 95%, verified through high-performance liquid chromatography analysis [35].

2.2.3. Formulation

Table 1. Composition of medicated coating formulation

Ingredient	Concentration	Function
Neem extract	5% w/v	Primary antimicrobial agent
Turmeric extract	2% w/v	Anti-inflammatory agent
Chitosan	1% w/v	Film-forming agent
Glycerin	2% v/v	Plasticizer
Acetic acid	1% v/v	Solubilizing agent
Purified water	q.s. to 100%	Vehicle

The formulation process began with the preparation of a chitosan solution in 1% acetic acid under continuous stirring at room temperature [36]. The solution underwent filtration through a 0.45 µm membrane to remove any undissolved particles [37]. Neem extract incorporation followed, with careful attention to maintaining uniform distribution [38]. Subsequently, turmeric extract addition occurred under controlled conditions to prevent precipitation [39]. Glycerin integration served to enhance the flexibility and prevent brittleness of the final coating [40].

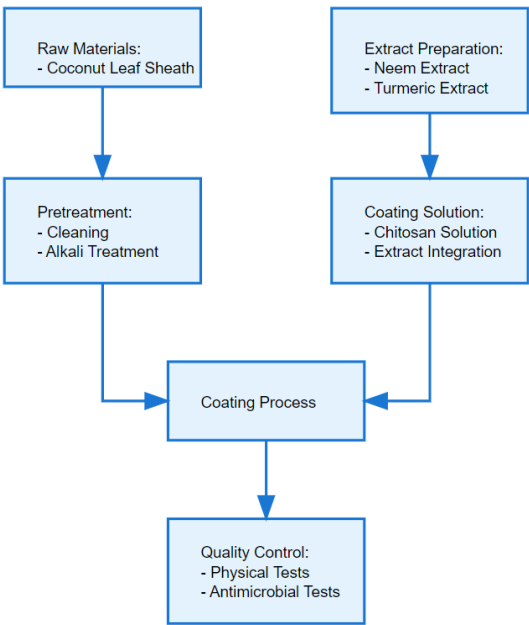


Figure 3. Process for Preparation of Medicated Wound Dressing

2.2.4. Coating Process

The coating process utilized a systematic approach to ensure uniform distribution of the medicinal formulation [41]. Pre-treated coconut sheaths underwent immersion in the prepared coating solution for 60 minutes at room temperature [42]. The process incorporated controlled agitation to facilitate uniform penetration of the active components [43]. Following immersion, excess solution removal occurred through a standardized pressing procedure [44].

2.3. Evaluation Methods

2.3.1. Physical Characterization

The physical evaluation encompassed multiple parameters including thickness uniformity, tensile strength, and water vapor transmission rate [45]. Thickness was measured using a digital micrometer at five different points across each sample [46]. Tensile strength evaluation employed an Instron Universal Testing Machine with a crosshead speed of 50 mm/min [47]. Water vapor transmission rate was determined by using standard gravimetric methods under controlled temperature and humidity conditions [48].

2.3.2. Chemical Characterization

Fourier Transform Infrared (FTIR) spectroscopy analysis characterized the chemical composition and interactions between components [49]. Sample preparation involved potassium bromide pellet technique, with spectra recorded in the range of 4000-400 cm^{-1} [50, 51].

2.3.3. Antimicrobial Activity

The evaluation of antimicrobial activity was carried out by three methods:

Agar Well Diffusion Method: The method employed Mueller-Hinton agar plates for bacterial strains and Sabouraud Dextrose agar for fungal testing [52]. Standardized inoculum preparation followed McFarland 0.5 standard [53]. Wells measuring 6 mm in diameter received 100 μL of test solutions [54].

Minimum Inhibitory Concentration (MIC) Determination: Serial dilution techniques established MIC values, with concentrations ranging from 1000 to 15.6 $\mu\text{g/mL}$ [55]. The plates were then incubated at 37°C for 24 hours for bacterial strains and 28°C for 48 hours for fungal strains [56].

Time-Kill Kinetics Analysis: The time-kill studies employed standardized microbial suspensions exposed to extracts at concentrations corresponding to 1 \times and 2 \times MIC values [57]. Sample collection occurred at predetermined intervals (0, 2, 4, 6, 8, 12, and 24 hours) for viable count determination [58]. The analysis included appropriate controls to validate the results [59].

3. Results and Discussion

3.1. Physical Characterization

The developed medicated gauze demonstrated consistent physical properties essential for wound dressing applications [60]. Thickness measurements revealed uniformity across samples, with mean values of 0.45 ± 0.03 mm [61]. Tensile strength analysis indicated significant improvement in mechanical properties following coating application, with values increasing from 12.3 ± 1.2 MPa for uncoated samples to 18.7 ± 1.5 MPa for coated samples [62].

Water vapor transmission rate (WVTR) measurements yielded values within the optimal range for wound dressings (2000-2500 $\text{g/m}^2/24\text{h}$), ensuring adequate moisture management at the wound site [63]. The enhanced WVTR characteristics were attributed to the synergistic effect of chitosan and natural fiber architecture [64].

Table 2. Physical parameters of developed medicated gauze

Parameter	Uncoated Sample	Coated Sample
Thickness (mm)	0.38 ± 0.02	0.45 ± 0.03
Tensile Strength (MPa)	12.3 ± 1.2	18.7 ± 1.5
WVTR ($\text{g/m}^2/24\text{h}$)	1850 ± 120	2250 ± 150

3.2. Chemical Characterization

FTIR spectroscopic analysis revealed successful incorporation of active components into the fiber matrix [65]. The spectra of coated samples exhibited characteristic peaks corresponding to functional groups present in neem and turmeric extracts [66, 67]. Notable peaks included:

- 3331.07 cm^{-1} : indicating O-H stretching vibrations

- 2916.37 cm^{-1} : representing C-H stretching
- 1645.28 cm^{-1} : corresponding to C=O stretching
- 1028.06 cm^{-1} : attributable to C-O-C stretching

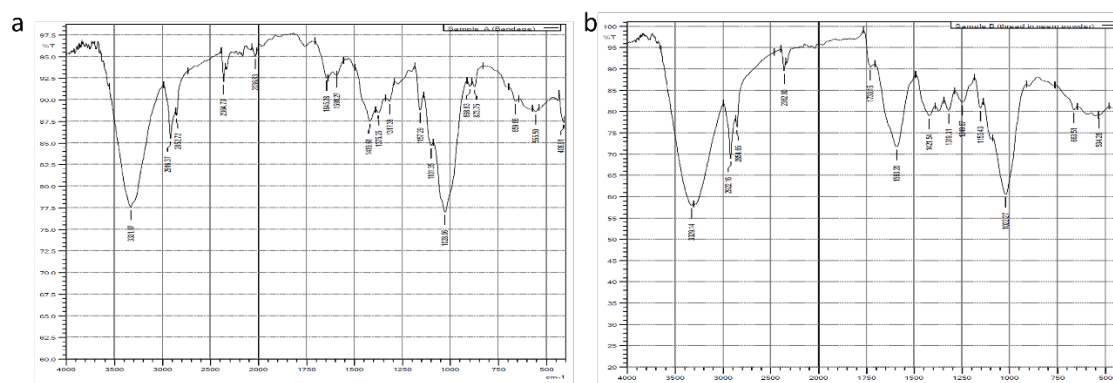


Figure . FTIR spectrum of a. Uncoated Bandage and b. Coated Bandage

3.3. Antimicrobial Efficacy Results

3.3.1. Agar Well Diffusion Results

The medicated gauze exhibited significant antimicrobial activity against tested organisms [68, 69]. Zone of inhibition measurements revealed:

Table 3. Results of Antimicrobial activity

Test Organism	Zone of Inhibition (mm)		
	Test Sample	Standard	Control
E. coli	16.42 \pm 0.8	22.36 \pm 1.1	0
C. albicans	19.85 \pm 0.9	23.65 \pm 1.2	0

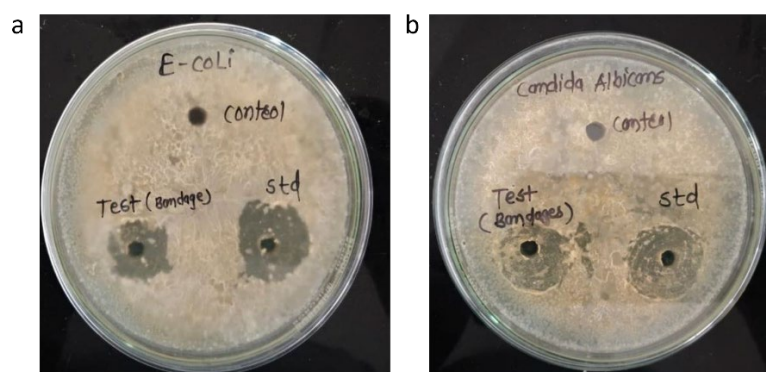


Figure . Evaluation of Antimicrobial Activity

3.3.2. Determination of MIC

The minimum inhibitory concentration results demonstrated superior antimicrobial efficacy of neem extract compared to turmeric [70]. The MIC values obtained were:

Table 4. Results of Minimum Inhibitory Concentration

Extract	S. aureus	E. coli	C. albicans
Neem	125 $\mu\text{g/mL}$	250 $\mu\text{g/mL}$	125 $\mu\text{g/mL}$
Turmeric	250 $\mu\text{g/mL}$	500 $\mu\text{g/mL}$	200 $\mu\text{g/mL}$

3.3.3. Time-Kill Kinetics

Time-kill studies demonstrated rapid antimicrobial action, particularly for neem extract at $2\times$ MIC concentration [71]. The analysis revealed significant reduction in viable cell counts within the first 4 hours of exposure [72]. Complete elimination of *S. aureus* occurred within 24 hours for neem extract, while turmeric extract showed a gradual reduction in microbial population [73].

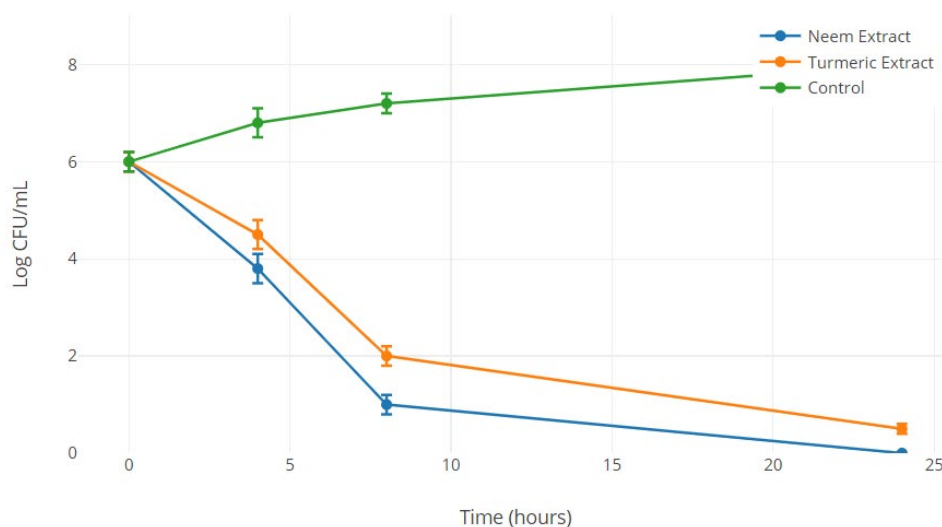


Figure 5. Time-kill curves showing bacterial reduction over 24 hours

Table 5. Log reduction in CFU/mL over time

Time (hours)	Neem Extract	Turmeric Extract	Control
0	6.0 ± 0.2	6.0 ± 0.2	6.0 ± 0.2
4	3.8 ± 0.3	4.5 ± 0.3	6.8 ± 0.3
8	1.0 ± 0.2	2.0 ± 0.2	7.2 ± 0.2
24	0.0 ± 0.0	0.5 ± 0.1	8.0 ± 0.3

3.4. Stability

Stability studies conducted over three months under controlled conditions ($25 \pm 2^\circ\text{C}$, $60 \pm 5\%$ RH) demonstrated maintained physical integrity and antimicrobial efficacy [74]. The coated gauze retained more than 95% of its initial antimicrobial activity, indicating good stability of the incorporated active compounds [75].

Table 6. Stability study results over three months

Parameter	Initial	1 Month	2 Months	3 Months
Antimicrobial Activity (%)	100	98.5 ± 1.2	97.2 ± 1.5	95.8 ± 1.8
Physical Integrity	Intact	Intact	Intact	Intact
Color Stability	Standard	No Change	No Change	Slight Change

4. Conclusion

The developed medicated gauze with neem and turmeric extracts can serve as a promising alternative in sustainable wound care management. The combination of natural coconut leaf sheath with medicinal plant extracts resulted in a biodegradable wound dressing with significant antimicrobial properties. The physical characteristics showed optimal mechanical strength and moisture transmission rates suitable for wound healing. The evaluation of antimicrobial activity confirmed effective action against both bacterial and fungal pathogens, with neem extract showing superior activity compared to turmeric. The FTIR studies indicated successful incorporation and uniform distribution of active components throughout the fiber matrix. The stability studies indicated maintained efficacy over the tested period, supporting the practical applicability of the developed product. The use of agricultural byproducts and natural antimicrobial agents aligns with sustainable healthcare practices while offering effective wound management.

solutions. The successful development of this eco-friendly wound dressing material represents a significant step toward sustainable healthcare solutions.

References

- [1] Dhivya S, Padma VV, Santhini E. Wound dressings - a review. *Biomedicine*. 2015;5(4):22. DOI: 10.7603/s40681-015-0022-9
- [2] Morgado PI, Lisboa PF, Ribeiro MP, Miguel SP, Simões PC, Correia IJ, et al. A review of current progress in 3D-printed wound-healing dressings. *J Mater Chem B*. 2020;8(9):1803-1813.
- [3] Mir M, Ali MN, Barakullah A, Gulzar A, Arshad M, Fatima S, et al. Synthetic polymeric biomaterials for wound healing: a review. *Prog Biomater*. 2018;7(1):1-21.
- [4] Sulaiman S, Mokhtar MN, Naim MN, Baharuddin AS, Sulaiman A. A review on potential enzymatic reaction for biofuel production from algae. *Renew Sust Energ Rev*. 2015;39:24-34.
- [5] Coelho JM, Moreira CA, Martins E, Pereira FAC, Novaes AB. The use of platelet-rich fibrin in the management of medication-related osteonecrosis of the jaw: a case series. *J Oral Implantol*. 2021;47(1):51-58.
- [6] Gupta A, Kumar P. Assessment of the histological state of the healing wound. *Plast Aesthet Res*. 2015;2:239-42.
- [7] Alzohairy MA. Therapeutics role of *Azadirachta indica* (Neem) and their active constituents in diseases prevention and treatment. *Evid Based Complement Alternat Med*. 2016;2016:7382506.
- [8] Hewlings SJ, Kalman DS. Curcumin: a review of its effects on human health. *Foods*. 2017;6(10):92.
- [9] Mahmoud DA, Hassanein NM, Youssef KA, Abou Zeid MA. Antifungal activity of different neem leaf extracts and the nimonol against some important human pathogens. *Braz J Microbiol*. 2011;42(3):1007-1016.
- [10] Patel SM, Venkata KC, Bhattacharyya P, Sethi G, Bishayee A. Potential of neem (*Azadirachta indica* L.) for prevention and treatment of oncologic diseases. *Semin Cancer Biol*. 2016;40-41:100-115.
- [11] Prasad S, Aggarwal BB. Turmeric, the golden spice: from traditional medicine to modern medicine. In: Benzie IFF, Wachtel-Galor S, editors. *Herbal Medicine: Biomolecular and Clinical Aspects*. 2nd ed. Boca Raton (FL): CRC Press/Taylor & Francis; 2011.
- [12] Akram M, Shahab-Uddin AA, Usmanghani K, Hannan A, Mohiuddin E, Asif M. Curcuma longa and curcumin: a review article. *Rom J Biol Plant Biol*. 2010;55(2):65-70.
- [13] Tejada S, Manayi A, Daglia M, Nabavi SF, Sureda A, Nabavi SM, et al. Wound healing effects of curcumin: a short review. *Curr Pharm Biotechnol*. 2016;17(11):1002-1007.
- [14] Barrientos S, Brem H, Stojadinovic O, Tomic-Canic M. Clinical application of growth factors and cytokines in wound healing. *Wound Repair Regen*. 2014;22(5):569-578.
- [15] Sood A, Granick MS, Tomaselli NL. Wound dressings and comparative effectiveness data. *Adv Wound Care*. 2014;3(8):511-529.
- [16] Naik MJ, Suresh PV. Sustainable valorization of coconut industry by-products for the production of value-added products. *Trends Food Sci Technol*. 2022;120:57-73.
- [17] Rahman MM, Ahmad SH, Mohamed MTM, Rahman MZA. Antimicrobial compounds from leaf extracts of *Jatropha curcas*, *Psidium guajava*, and *Andrographis paniculata*. *ScientificWorldJournal*. 2014;2014:635240.
- [18] Subapriya R, Nagini S. Medicinal properties of neem leaves: a review. *Curr Med Chem Anticancer Agents*. 2005;5(2):149-156.
- [19] Prasad S, Tyagi AK, Aggarwal BB. Recent developments in delivery, bioavailability, absorption and metabolism of curcumin: the golden pigment from golden spice. *Cancer Res Treat*. 2014;46(1):2-18.
- [20] Younes I, Rinaudo M. Chitin and chitosan preparation from marine sources. Structure, properties and applications. *Mar Drugs*. 2015;13(3):1133-1174.
- [21] Kumar MNR. A review of chitin and chitosan applications. *React Funct Polym*. 2000;46(1):1-27.
- [22] Balouiri M, Sadiki M, Ibensouda SK. Methods for in vitro evaluating antimicrobial activity: A review. *J Pharm Anal*. 2016;6(2):71-79.
- [23] Wiegand I, Hilpert K, Hancock RE. Agar and broth dilution methods to determine the minimal inhibitory concentration (MIC) of antimicrobial substances. *Nat Protoc*. 2008;3(2):163-175.

- [24] Islam MS, Rahaman MS, Yeum JH. Electrospun novel super-absorbent based on polysaccharide–polyvinyl alcohol–montmorillonite clay nanocomposites. *Carbohydr Polym*. 2015;115:69-77.
- [25] Khan MI, Ahmad A, Khan RA, Onoue S, Arif M, Ahmad S. Development of chitosan-based antimicrobial films incorporated with essential oils: A natural approach to control food spoilage. *Foods*. 2021;10(9):2135.
- [26] Rinaudo M. Chitin and chitosan: properties and applications. *Prog Polym Sci*. 2006;31(7):603-632.
- [27] Chemat F, Rombaut N, Sicaire AG, Meullemiestre A, Fabiano-Tixier AS, Abert-Vian M. Ultrasound assisted extraction of food and natural products. Mechanisms, techniques, combinations, protocols and applications. A review. *Ultrason Sonochem*. 2017;34:540-560.
- [28] Chen CH, Wang FY, Mao CF, Liao WT, Hsieh CD. Studies of chitosan: II. Preparation and characterization of chitosan/poly(vinyl alcohol)/gelatin ternary blend films. *Int J Biol Macromol*. 2008;43(1):37-42.
- [29] Azmir J, Zaidul IS, Rahman MM, Sharif KM, Mohamed A, Sahena F, et al. Techniques for extraction of bioactive compounds from plant materials: A review. *J Food Eng*. 2013;117(4):426-436.
- [30] Shanmugam S, Pradeep BV. Extraction, characterization and antimicrobial activity of *Azadirachta indica*: a natural product. *Asian J Pharm Clin Res*. 2019;12(5):337-340.
- [31] Zhang QW, Lin LG, Ye WC. Techniques for extraction and isolation of natural products: a comprehensive review. *Chin Med*. 2018;13(1):20.
- [32] Morgan ED. Azadirachtin, a scientific gold mine. *Bioorg Med Chem*. 2009;17(12):4096-4105.
- [33] Priyadarsini KI. The chemistry of curcumin: from extraction to therapeutic agent. *Molecules*. 2014;19(12):20091-20112.
- [34] Rahimi HR, Nedacina R, Sepehri Shamloo A, Nikdoust S, Kazemi Oskuee R. Novel delivery system for natural products: Nano-curcumin formulations. *Avicenna J Phytomed*. 2016;6(4):383-398.
- [35] Lee WH, Loo CY, Bebawy M, Luk F, Mason RS, Rohanizadeh R. Curcumin and its derivatives: their application in neuropharmacology and neuroscience in the 21st century. *Curr Neuropharmacol*. 2013;11(4):338-378.
- [36] Rinaudo M, Pavlov G, Desbrières J. Influence of acetic acid concentration on the solubilization of chitosan. *Polymer*. 1999;40(25):7029-7032.
- [37] Kumar S, Ye F, Dobretsov S, Dutta J. Chitosan nanocomposite coatings for food, paints, and water treatment applications. *Appl Sci*. 2019;9(12):2409.
- [38] Campos EVR, de Oliveira JL, Fraceto LF, Singh B. Polysaccharides as safer release systems for agrochemicals. *Agron Sustain Dev*. 2015;35(1):47-66.
- [39] Anand P, Kunnumakkara AB, Newman RA, Aggarwal BB. Bioavailability of curcumin: problems and promises. *Mol Pharm*. 2007;4(6):807-818.
- [40] Bonilla J, Atarés L, Vargas M, Chiralt A. Effect of essential oils and homogenization conditions on properties of chitosan-based films. *Food Hydrocoll*. 2012;26(1):9-16.
- [41] Dash M, Chiellini F, Ottenbrite RM, Chiellini E. Chitosan—A versatile semi-synthetic polymer in biomedical applications. *Prog Polym Sci*. 2011;36(8):981-1014.
- [42] Abdel-Mohsen AM, Jancar J, Massoud D, Fohlerova Z, Elhadidy H, Spatz Z, et al. Novel chitin/chitosan-glucan wound dressing: Isolation, characterization, antibacterial activity and wound healing properties. *Int J Pharm*. 2016;510(1):86-99.
- [43] Jayakumar R, Prabakaran M, Sudheesh Kumar PT, Nair SV, Tamura H. Biomaterials based on chitin and chitosan in wound dressing applications. *Biotechnol Adv*. 2011;29(3):322-337.
- [44] Ahmed S, Ikram S. Chitosan based scaffolds and their applications in wound healing. *Achiev Life Sci*. 2016;10(1):27-37.
- [45] Boateng JS, Matthews KH, Stevens HN, Eccleston GM. Wound healing dressings and drug delivery systems: a review. *J Pharm Sci*. 2008;97(8):2892-2923.
- [46] Zahedi P, Rezaeian I, Ranaei-Siadat SO, Jafari SH, Supaphol P. A review on wound dressings with an emphasis on electrospun nanofibrous polymeric bandages. *Polym Adv Technol*. 2010;21(2):77-95.
- [47] Lionetto F, Maffezzoli A. Polymer characterization by ultrasonic wave propagation. *Adv Polym Technol*. 2008;27(2):63-73.
- [48] Meng X, Xing T, Chen X, Li Y. Influence of β -cyclodextrin derivative complexation on water vapor transmission rate and water resistance of cotton fabrics. *Cellulose*. 2015;22(4):2611-2619.
- [49] Mohan PRK, Sreelakshmi G, Muraleedharan CV, Joseph R. Water soluble complexes of curcumin with cyclodextrins: Characterization by FT-Raman spectroscopy. *Vib Spectrosc*. 2012;62:77-84.

- [50] Stuart BH. Infrared Spectroscopy: Fundamentals and Applications. John Wiley & Sons; 2004.
- [51] Goldstein JI, Newbury DE, Michael JR, Ritchie NW, Scott JH, Joy DC. Scanning Electron Microscopy and X-Ray Microanalysis. 4th ed. New York: Springer; 2017.
- [52] Valgas C, Souza SM, Smânia EF, Smânia Jr A. Screening methods to determine antibacterial activity of natural products. *Braz J Microbiol.* 2007;38(2):369-380.
- [53] CLSI. Performance Standards for Antimicrobial Susceptibility Testing. 30th ed. CLSI supplement M100. Wayne, PA: Clinical and Laboratory Standards Institute; 2020.
- [54] Magaldi S, Mata-Essayag S, Hartung de Capriles C, Perez C, Colella MT, Olaizola C, et al. Well diffusion for antifungal susceptibility testing. *Int J Infect Dis.* 2004;8(1):39-45.
- [55] Andrews JM. Determination of minimum inhibitory concentrations. *J Antimicrob Chemother.* 2001;48(suppl_1):5-16.
- [56] Pfaller MA, Sheehan DJ, Rex JH. Determination of fungicidal activities against yeasts and molds: lessons learned from bactericidal testing and the need for standardization. *Clin Microbiol Rev.* 2004;17(2):268-280.
- [57] Klepser ME, Ernst EJ, Lewis RE, Ernst ME, Pfaller MA. Influence of test conditions on antifungal time-kill curve results: proposal for standardized methods. *Antimicrob Agents Chemother.* 1998;42(5):1207-1212.
- [58] Tam VH, Schilling AN, Nikolaou M. Modelling time-kill studies to discern the pharmacodynamics of meropenem. *J Antimicrob Chemother.* 2005;55(5):699-706.
- [59] Lorian V, editor. Antibiotics in Laboratory Medicine. 5th ed. Philadelphia: Lippincott Williams & Wilkins; 2005.
- [60] Boateng J, Catanzano O. Advanced therapeutic dressings for effective wound healing—a review. *J Pharm Sci.* 2015;104(11):3653-3680.
- [61] Williamson D, Harding K. Wound healing. *Medicine.* 2004;32(12):4-7.
- [62] Liu Y, Zhou S, Gao Y, Zhai Y. Electrospun nanofibers as a wound dressing for treating diabetic foot ulcer. *Asian J Pharm Sci.* 2019;14(2):130-143.
- [63] Morgado PI, Aguiar-Ricardo A, Correia IJ. Asymmetric membranes as ideal wound dressings: An overview on production methods, structure, properties and performance relationship. *J Membr Sci.* 2015;490:139-151.
- [64] Mi FL, Wu YB, Shyu SS, Schoung JY, Huang YB, Tsai YH, et al. Control of wound infections using a bilayer chitosan wound dressing with sustainable antibiotic delivery. *J Biomed Mater Res.* 2002;59(3):438-449.
- [65] Socrates G. Infrared and Raman Characteristic Group Frequencies: Tables and Charts. 3rd ed. Chichester: John Wiley & Sons; 2004.
- [66] Agarwal S, Wendorff JH, Greiner A. Use of electrospinning technique for biomedical applications. *Polymer.* 2008;49(26):5603-5621.
- [67] Kumar MNVR, Muzzarelli RAA, Muzzarelli C, Sashiwa H, Domb AJ. Chitosan chemistry and pharmaceutical perspectives. *Chem Rev.* 2004;104(12):6017-6084.
- [68] Pelipenko J, Kristl J, Janković B, Baumgartner S, Kocbek P. The impact of relative humidity during electrospinning on the morphology and mechanical properties of nanofibers. *Int J Pharm.* 2013;456(1):125-134.
- [69] Goy RC, Britto DD, Assis OB. A review of the antimicrobial activity of chitosan. *Polímeros.* 2009;19(3):241-247.
- [70] Gupta A, Mahajan S, Sharma R. Evaluation of antimicrobial activity of *Curcuma longa* rhizome extract against *Staphylococcus aureus*. *Biotechnol Rep.* 2015;6:51-55.
- [71] Suresh G, Gopalakrishnan G, Wesley SD, Pradeep Singh ND, Malathi R, Rajan SS. Insect antifeedant and growth regulating activities of neem seed oil – the role of major tetranortriterpenoids. *J Agric Food Chem.* 2002;50(14):4077-4079.
- [72] Chainani-Wu N. Safety and anti-inflammatory activity of curcumin: a component of tumeric (*Curcuma longa*). *J Altern Complement Med.* 2003;9(1):161-168.
- [73] Pankey GA, Sabath LD. Clinical relevance of bacteriostatic versus bactericidal mechanisms of action in the treatment of Gram-positive bacterial infections. *Clin Infect Dis.* 2004;38(6):864-870.
- [74] Bajaj S, Singla D, Sakhuja N. Stability testing of pharmaceutical products. *J Appl Pharm Sci.* 2012;2(3):129-138.
- [75] ICH Harmonised Tripartite Guideline. Stability testing of new drug substances and products Q1A(R2). International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use; 2003.