

The Journal of Phytopharmacology

(Pharmacognosy and phytomedicine Research)



Review Article

ISSN 2320-480X

JPHYTO 2025; 14(4): 291-297

July- August

Received: 24-06-2025

Accepted: 28-08-2025

Published: 30-09-2025

©2025, All rights reserved

doi: 10.31254/phyto.2025.14410

Nisha B. Suryawanshi

1. Department of Botany, K.R.T. Arts, B.H. Commerce, and A.M. Science College, Nashik- 422002, Maharashtra, India

2. Department of Botany, Mahant Jamanadas Maharaj Arts, Commerce and Science College, Nashik- 422208, Maharashtra, India

Arati C. Sutar

Department of Botany, Yashwantrao Chavan Institute of Science, Karmaveer Bhaurao Patil University, Satara-415001, Maharashtra, India

Sumaiya S. Shaikh

Department of Botany, Yashwantrao Chavan Institute of Science, Karmaveer Bhaurao Patil University, Satara-415001, Maharashtra, India

Nitin T. Gore

Department of Botany, Yashwantrao Chavan Institute of Science, Karmaveer Bhaurao Patil University, Satara-415001, Maharashtra, India

Avinash S. Jondhale

Department of Botany, Mahant Jamanadas Maharaj Arts, Commerce and Science College, Nashik- 422208, Maharashtra, India

Smita P. Chavan

Department of Botany, Mahant Jamanadas Maharaj Arts, Commerce and Science College, Nashik- 422208, Maharashtra, India

Mahendra L. Ahire

Department of Botany, Yashwantrao Chavan Institute of Science, Karmaveer Bhaurao Patil University, Satara-415001, Maharashtra, India

Correspondence:

Dr. Mahendra Laxman Ahire

Department of Botany, Yashwantrao Chavan Institute of Science, Karmaveer Bhaurao Patil University, Satara-415001, Maharashtra, India
Email: mlahire@gmail.com

Malvastrum coromandelianum (L.) Garcke: A versatile weed with promising pharmacological applications

Nisha B. Suryawanshi, Arati C. Sutar, Sumaiya S. Shaikh, Nitin T. Gore, Avinash S. Jondhale, Smita P. Chavan, Mahendra L. Ahire

ABSTRACT

Malvastrum coromandelianum (L.) Garcke, commonly known as false mallow, belongs to the family Malvaceae. This invasive alien weed is widely distributed throughout tropical and subtropical regions, where it has become well established due to its resilience and adaptability. Although often regarded as a weed, *M. coromandelianum* possesses significant pharmacological potential, particularly in traditional medicine systems where it has been used for various therapeutic purposes. The plant is a rich source of bioactive compounds such as tannins, flavonoids, alkaloids, and phenolics, all known for their medicinal properties. Recent studies have identified novel bioactive molecules in *M. coromandelianum*, revealing antimicrobial, anti-inflammatory, antioxidant, and anticancer activities. These findings underscore its promise as a source of natural therapeutic agents. However, several challenges hinder its practical application, including variability in phytochemical content, lack of standardization in herbal formulations, and the need to ensure safety and efficacy in clinical settings. Further research is required to elucidate its mechanisms of action, optimize its therapeutic use, and integrate it into modern pharmacological practices. This review aims to consolidate current knowledge, highlight its medicinal potential, and provide direction for future research.

Keywords: Antioxidant activity, Bioactive compounds, Flavonoids, Phenolics, Phytochemical content.

INTRODUCTION

Malvastrum coromandelianum (L.) Garcke, commonly known as Coromandel False Mallow, is an annual herb widely used in traditional medicine, particularly for treating wounds, inflammation, and digestive disorders [1, 2]. Also referred to as *Malva coromandelica*, it has a long-standing role in Indian and Sri Lankan ethnomedicine. Its wound-healing efficacy has been substantiated by recent studies [3, 4]. Modern research has begun to validate these traditional claims by exploring the plant's phytochemical and pharmacological properties. Integrating traditional knowledge with scientific evidence enhances the credibility of natural remedies in modern medicine. Ecologically, *M. coromandelianum* is adaptable and found in tropical Africa, across Asia (notably India and Southeast Asia), and parts of the Americas. Despite its medicinal value, it can be invasive in certain regions, necessitating ecological monitoring [4]. Phytochemical investigations reveal that *M. coromandelianum* contains flavonoids, alkaloids, and other bioactive compounds, contributing to its anti-inflammatory, analgesic, and gastroprotective effects [5]. Its antioxidant properties suggest potential roles in managing oxidative stress-related disorders [6, 7].

Traditionally, the plant has been used for treating rashes, respiratory conditions, and diabetes [8]. Notably, zinc nanoparticles synthesized from its extracts have demonstrated hepatoprotective effects by modulating antioxidant enzymes and reducing oxidative and inflammatory damage [9]. To validate its efficacy and ensure safety for contemporary use, traditional applications must be supported by systematic scientific studies [10, 11].

This review aims to identify and characterize the phytochemicals in *M. coromandelianum*, evaluate its therapeutic efficacy with a focus on anti-inflammatory and antioxidant activities, and emphasize the integration of traditional knowledge with modern scientific methods in drug discovery. These insights are particularly relevant considering the global burden of inflammation and oxidative stress-related diseases. The fusion of traditional knowledge with modern approaches such as metabolomics, genomics, and bioinformatics is reshaping medicinal plant research. This trend not only helps preserve traditional practices but also opens new avenues for drug development.

Phytochemical Composition

Phytochemical analysis of *M. coromandelianum* leaf extracts using solvents of varying polarity reveals significant differences in phenolic and flavonoid contents (Table 1). Solvent polarity plays a vital role, with aqueous methanol and methanol being particularly effective [12]. Aqueous acetone also yields good

results [13], while non-polar solvents like chloroform and petroleum ether are less effective. Thus, polar solvents are preferred for isolating antioxidant constituents for medicinal purposes.

Pharmacological Activities

M. coromandelianum demonstrates a wide range of biological activities, including anti-inflammatory, antimicrobial, antioxidant, analgesic, hepatoprotective, and pharmacognostic effects (Table 2; Figure 1). Aqueous and ethyl acetate extracts have shown anti-inflammatory and antibacterial activities in experimental models such as carrageenan-induced paw edema and the formalin-induced pain test [13, 14].

Phytochemical screening has identified compounds such as β -sitosterol, flavonoids, tannins, and phenolic constituents, which are linked to its antioxidant and antimicrobial effects. GC-MS analysis has revealed 29 bioactive compounds, including terpenoids, fatty acids, vitamins, and phenols, which support the plant's traditional applications.

Green-synthesized zinc oxide nanoparticles from *M. coromandelianum* have exhibited hepatoprotective and antioxidant properties by mitigating oxidative stress in animal models [13]. A polysaccharide extract from the plant has also demonstrated potent antioxidant activity [15].

Network pharmacology and chemoinformatic analyses further support the plant's potential in anti-aging, anti-inflammatory, and antioxidant therapies [14]. These findings collectively highlight the therapeutic promise of this plant in the development of natural drugs.

Bioactive Compounds and Therapeutic Effects

The therapeutic potential of *M. coromandelianum* is attributed to a diverse array of phytochemicals (Table 3; Figure 2). β -sitosterol,

abundant in the leaves, is associated with analgesic, anti-inflammatory, and anticancer activities [16 - 20]. Apigenin derivatives offer neuroprotective, cardioprotective, and antimicrobial benefits [21 - 24].

Fatty acids such as 9,12,15-octadecatrienoic acid possess pronounced anti-inflammatory and antioxidant properties [25 - 28]. Diosgenin has been recognized for its anticancer, neuroprotective, and hypoglycemic effects [29 - 32]. Other bioactive compounds, including squalene, vitamin E, and phytol, contribute to antioxidative, anti-aging, and antimicrobial actions [33 - 49]. Altogether, these phytochemicals affirm the pharmacological potential of *M. coromandelianum* as a valuable source of natural bioactive agents.

Nanoparticle Biosynthesis

Green synthesis of nanoparticles using *M. coromandelianum* provides an eco-friendly and sustainable alternative to conventional methods. Plant extracts from various parts have been employed to synthesize silver, zinc oxide, and magnesium oxide nanoparticles. These extracts serve both as reducing and stabilizing agents due to their rich phytochemical content.

Advanced characterization techniques such as UV-Vis, FTIR, XRD, SEM, TEM, and DLS confirm their nanoscale properties. These nanoparticles display strong antimicrobial activity against *Escherichia coli* and *Staphylococcus aureus* [50]. They also show antioxidant and hepatoprotective effects in CCl₄-induced liver injury models [9] and exhibit antiviral and mosquitocidal activity against the dengue virus and its vectors [51].

Given their wide range of bioactivities, these nanoparticles hold promise for applications in biomedicine, drug delivery, catalysis, and environmental remediation [52]. However, further research is necessary to fully assess their biocompatibility, cytotoxicity, and long-term safety.

Table 1: Phytochemical composition in leaves of *M. coromandelianum* (L) Garcke

S. No.	Solvent	Phytochemical analysis		References
		Total Phenolic Content (mg GAE/g or μ g GAE/mg extract)	Total Flavonoid Content (mg QE/g or equivalent)	
1.	Methanol	45.25 \pm 3.48	78.12 \pm 2.45	Sanghai et al. [12]
2.	Acetone	41.78 \pm 7.86	35.42 \pm 1.98	
3.	Chloroform	29.54 \pm 2.36	22.30 \pm 1.56	
4.	Petroleum ether	15.25 \pm 1.25	11.78 \pm 0.95	
5.	Aqua Methanol (AM)	88.45 \pm 1.72	54.60 \pm 1.12	Saxena and Rao [13]
6.	Aqua Acetone (AA)	67.50 \pm 1.58	43.35 \pm 0.98	

Table 2: Pharmacological activities of *M. coromandelianum* (L) Garcke

S. No.	Activity	Solvents	Plant Part used	Reference
1.	Ant inflammatory	Aqueous	Whole plant	Sanghai et al. [12]; Aderogba et al. [22]; Yadav and Mohite [14]
2.	Analgesic	Aqueous, Alcohol-water	Whole plant	Sanghai et al. [12]; Bhupesh et al. [53]
3.	Antinociceptive	Alcohol-water	Areal parts	Reddy et al. [54]; Sanghai et al. [12]
4.	Larvicidal and ovicidal	Methanol	Leaves	Saxena et al. [4]; Irudayaraj and Fabiola [55]
5.	Anthelmintic	Methanol	Leaves	Yadav and Mohite [56]; Saxena et al. [4]
6.	Anti-Mycobacterial	Methanol	Leaves	Aderogba et al. [12]
7.	Antiulcer	Ethanol	Whole plant	Balekar et al. [57]; Saxena et al. [4]
8.	Hemolytic	Methanol	Roots	Sheema et al. [5]

9.	Antidiabetic	Aqueous	Leaves	Andrade-Cetto and Heinrich ^[58] ; Deore et al. ^[8]
10.	Antidiarrheal	Ethanol	Leaves	Balekar et al. ^[59] ; Saxena et al. ^[4]
11.	Anticancer	Methanol	Whole plant	Yadav and Mohite ^[56]
12.	Antioxidant	Aqueous, Aqua methanol, Acetone	Leaves	Saxena and Rao ^[13] ; Yadav and Mohite ^[14]
13.	Antibacterial and antifungal	Ethyl acetate, Ethanol, Chloroform, Hexane, DMSO, Aqueous	Areal parts, Leaves	Yadav and Mohite ^[14] ; Sheema et al. ^[5]

Table 3: Bioactive compounds reported in *M. coromandelianum* (L) Garcke

S. No.	Name of compound	Chemical formula	Plant part	Biological activity	References
1.	β -sitosterol	C ₂₉ H ₅₀ O	Leaves	Analgesic, anthelmintic, antimutagenic, anti-inflammatory, antihepatotoxic, anti-cancer and antiviral	Sanghai et al. ^[12]
2.	Apigenin-7-O- β -6'' (p-coumaroyl)-glucopyranoside	C ₃₀ H ₂₆ O ₁₂	Leaves	Anti-inflammatory, antimycobacterial, neuroprotective, cardioprotective, and intestinal protection	Aderogba et al. ^[22]
3.	1,2-benzenedicarboxylic acid	C ₈ H ₆ O ₄	Leaves	Antimicrobial, neuroprotective, cytotoxic, antioxidant	Saxena and Rao ^[13]
4.	2-Methoxy-4-vinylphenol	C ₉ H ₁₀ O ₂	Leaves	Antimicrobial, anti-inflammatory, antitumor, antioxidant	Saxena and Rao ^[13]
5.	3-Bromocholest-5-ene	C ₃₀ H ₅₀ Br	Leaves	Antifungal and cytotoxic	Saxena and Rao ^[13]
6.	4H-pyran-4-one, 2,3-dihydro-3,5-dihydroxy-6-methyl	C ₆ H ₈ O ₄	Leaves	Antioxidant, anti-inflammatory, skin health benefits	Saxena and Rao ^[13]
7.	Diosgenin	C ₂₇ H ₄₂ O ₃	Leaves	Anticancer, anti-inflammatory, antioxidant, neuroprotective, hypolipidemic and hypoglycemic, antimicrobial and immunomodulatory, antidiabetic, cardioprotective, osteogenic, antithrombotic	Saxena and Rao ^[13]
8.	9,12,15-octadecatrienoic acid, (Z,Z,Z)	C ₁₈ H ₃₀ O ₂	Leaves	Anti-Inflammatory, antioxidant, antimicrobial, neuroprotective, antidepressant, antitumor and cardiovascular benefits	Saxena and Rao ^[13]
9.	9,12,15-octadecatrienoic acid, methyl ester, (Z,Z,Z)	C ₁₉ H ₃₂ O ₂	Leaves	Anti-Inflammatory, antioxidant and antimicrobial	Saxena and Rao ^[13]
10.	γ -sitosterol	C ₂₉ H ₅₀ O	Leaves	Antihyperglycemic, anticancer, anti-inflammatory	Saxena and Rao ^[13]
11.	Guanosine	C ₁₀ H ₁₃ N ₅ O ₅	Leaves	Neuroprotective, antiprotozoal, immunostimulatory, neuronal Differentiation and anticancer	Saxena and Rao ^[13]
12.	Hexadecanoic acid, methyl ester	C ₁₇ H ₃₄ O ₂	Leaves	Antioxidant, antihypocholestermic, anti-inflammatory and antibacterial	Saxena and Rao ^[13]
13.	Neophytadiene	C ₂₀ H ₃₈	Leaves	Anti-inflammatory, anxiolytic-like and anticonvulsant	Saxena and Rao ^[13]
14.	Octadecanoic acid	C ₁₈ H ₃₆ O ₂	Leaves	Antimicrobial, anti-inflammatory, antioxidant, anticancer and lipid metabolism and mitochondrial function	Saxena and Rao ^[13]
15.	Phenol, 2,4-Bis (1,1-dimethylethyl)	C ₁₇ H ₃₀ OSi	Leaves	Antimicrobial, anti-inflammatory, antioxidant, cytotoxic, anticancer, phytotoxic, insecticidal and nematocidal	Saxena and Rao ^[13]
16.	Phytol	C ₂₀ H ₄₀ O	Leaves	Antimicrobial, anti-quorum sensing, cytotoxic,	Saxena and Rao ^[13]

				immuno-adjuvants, antinociceptive, antitumor, anticonvulsant, anti-inflammatory, anxiolytic, anticandidal, antioxidant and antihyperpigmentation	
17.	Squalene	C ₃₀ H ₅₀	Leaves	Antioxidant, anti-inflammatory, antiviral and anticancer	Saxena and Rao ^[13]
18.	Vitamin E	C ₂₉ H ₅₀ O ₂	Leaves	Anti-inflammatory, antioxidative, cardiovascular health and anti-aging	Saxena and Rao ^[13]
19.	Sabinene	C ₁₀ H ₁₆	Leaves	Antimicrobial, antioxidant, angiostatic, antiangiogenic, anti-inflammatory, anticancer	Irudayaraj and Fabiola ^[55]
20.	Limonene	C ₁₀ H ₁₆	Leaves	Anticancer, anti-inflammatory, antioxidant, neuroprotective, antimicrobial, antileishmanial, anthelmintic and insecticidal	Irudayaraj and Fabiola ^[55]
21.	Caryophyllene	C ₁₅ H ₂₄	Leaves	Anti-inflammatory, antimicrobial, immunomodulatory, analgesic, anticancer, neuroprotective, antioxidant	Irudayaraj and Fabiola ^[55]
22.	Naphthalene	C ₁₀ H ₈	Leaves	Anti-inflammatory, antimicrobial, antioxidant, antiproliferative, anticancer, anticandidal, and antidiabetic	Irudayaraj and Fabiola ^[55]
23.	Bicyclogermacrene	C ₁₅ H ₂₄	Leaves	Antimicrobial, antioxidant, cytotoxic and anti-inflammatory,	Irudayaraj and Fabiola ^[55]
24.	Spathulenol	C ₁₅ H ₂₄ O	Leaves	Antioxidant, anti-inflammatory, antiproliferative, antimycobacterial, antimicrobial, anticholinesterase, antinociceptive, anti-hyperalgesic, insecticidal and antitumor	Irudayaraj and Fabiola ^[55]
25.	Ledol	C ₁₅ H ₂₆ O	Leaves	Antimicrobial, anti-inflammatory, insecticidal, antioxidant, cytotoxic and analgesic	Irudayaraj and Fabiola ^[55]
26.	2-Hydrazinopyridine	C ₅ H ₇ N ₃	Leaves	Anticancer, antibacterial and antioxidant	Irudayaraj and Fabiola ^[55]
27.	Tiliroside [kaempferol-3-O-β-d-(6"-E-p-coumaryl) glucopyranoside]	C ₃₀ H ₂₆ O ₁₃	Leaves	Antidiabetic, anti-inflammatory, antioxidant, anticarcinogenic, hepatoprotective	Devi and Kumar ^[60]
28.	Quercetrin (quercetin 3-O-α-l-rhamnopyranoside)	C ₂₁ H ₂₀ O ₁₁	Leaves	Antioxidant, antibacterial, wound healing, immunomodulatory and antiviral	Devi and Kumar ^[60]
29.	Propiophenone	C ₉ H ₁₀ O	Leaves	Antioxidant and anti-inflammatory	Irudayaraj and Fabiola ^[55]
30.	Nonanoic acid	C ₉ H ₁₈ O ₂	Root	Antialgal, antimicrobial and anticancer	Sheema et al. ^[5]
31.	Oleic Acid	C ₁₈ H ₃₄ O ₂	Root	Antioxidant, anti-inflammatory, antimicrobial, anticancer	Sheema et al. ^[5]
32.	Eicosane	C ₂₀ H ₄₂	Root	Antioxidant, antifungal, anti-inflammatory, neuroprotective, wound healing	Sheema et al. ^[5]
33.	Ar-turmerone	C ₁₅ H ₂₀ O	Root	Anti-malignant, anti-aging, anti-inflammatory, anticancer, antioxidant, antivenom, antifungal, insecticidal, hepatoprotective, hypoglycemic	Sheema et al. ^[5]

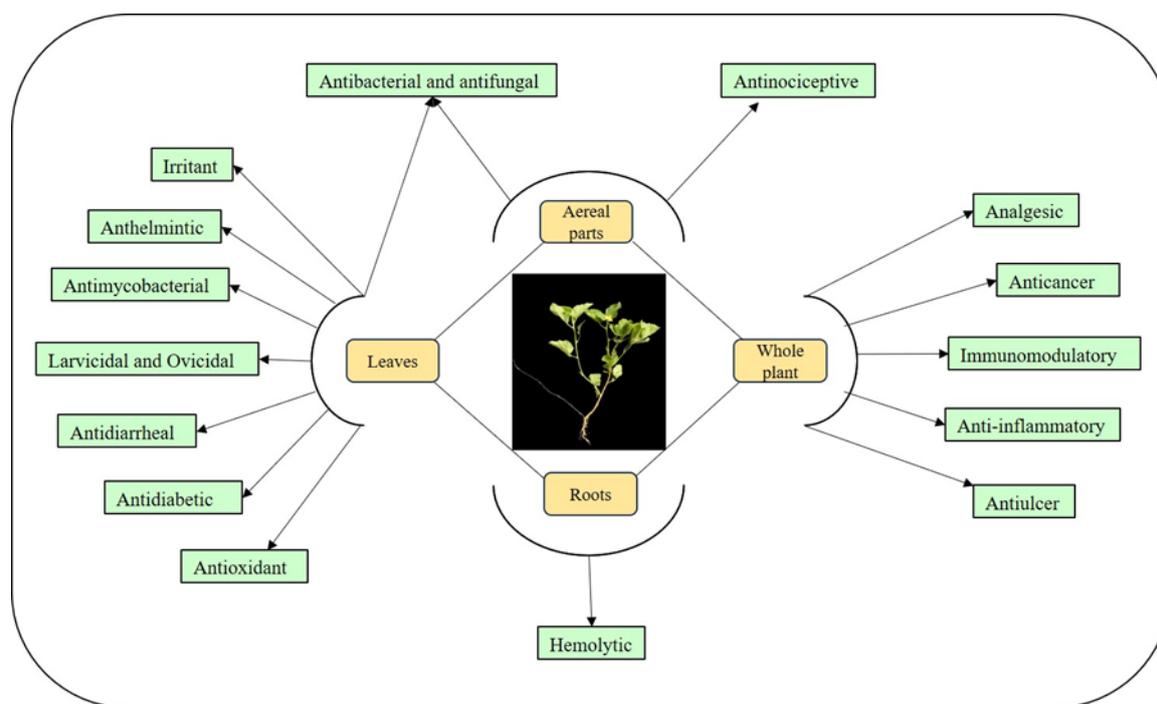


Figure 1: Pharmacological activities of *M. coromandelianum* (L) Garcke

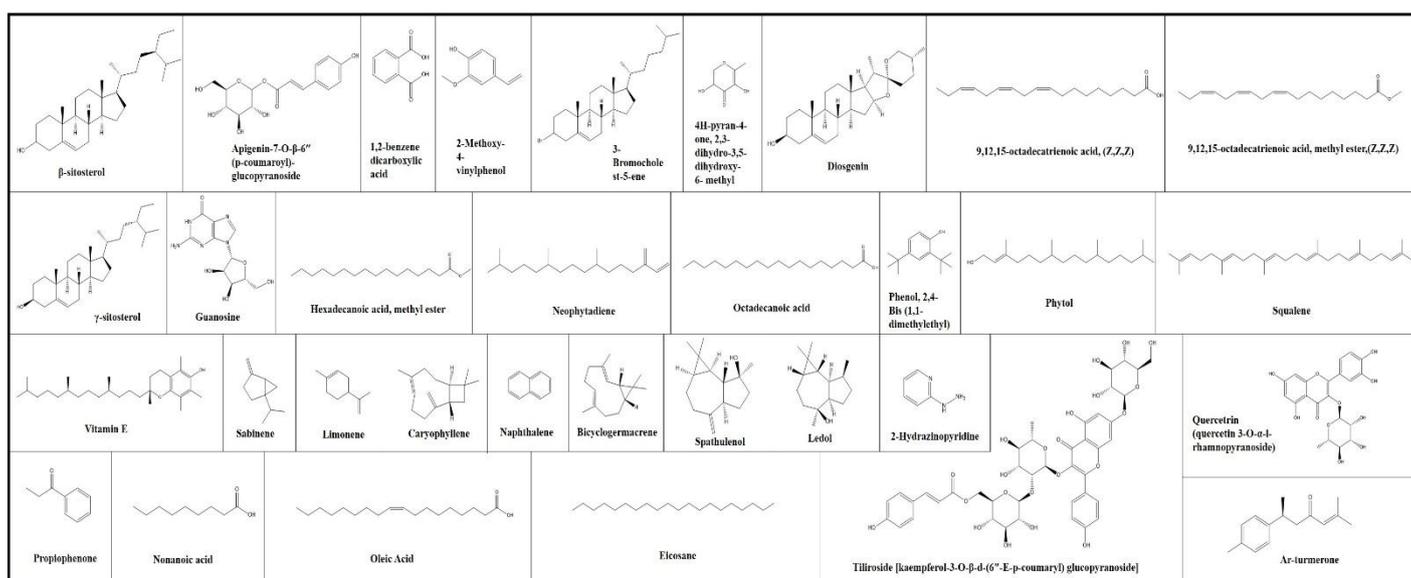


Figure 2: Chemical structures of reported compounds present in *M. coromandelianum* (L) Garcke

CONCLUSION

M. coromandelianum is a traditionally significant medicinal plant with a diverse phytochemical profile that substantiates its widespread ethnopharmacological uses. The presence of bioactive compounds such as flavonoids, β -sitosterol, fatty acids, and saponins underlies its broad spectrum of pharmacological activities, including anti-inflammatory, antioxidant, antimicrobial, and hepatoprotective effects. Recent advances in green nanotechnology using *M. coromandelianum* extracts have expanded its therapeutic potential, particularly in synthesizing biocompatible nanoparticles with promising biomedical applications. The integration of traditional knowledge with modern scientific validation techniques, including phytochemical profiling and network pharmacology, is critical for advancing *M. coromandelianum* from folk medicine to evidence-based therapeutic use. However, despite encouraging preliminary data, further comprehensive studies particularly *in vivo* and clinical trials are essential to confirm its safety, efficacy, and mechanistic pathways. Given the global rise in diseases linked to inflammation

and oxidative stress, *M. coromandelianum* holds considerable promise as a source of novel bioactive compounds and sustainable nanomaterials. Continued interdisciplinary research will not only deepen understanding of its pharmacological mechanisms but also promote the sustainable utilization of this versatile plant in modern healthcare and environmental applications.

Acknowledgments

The authors are also grateful to Yashavantrao Chavan Institute of Science, Satara, for the financial support under the self-funded Project for faculty and Rayat Institute of Research and Development, Satara, for providing necessary laboratory facilities. The authors sincerely acknowledge K. R. T. Arts, B. H. Commerce & A. M. Science College, Nashik, for their generous support and contributions to this study. Authors also extend sincere gratitude to Savitribai Phule Pune University (SPPU) for its steadfast dedication to academic excellence, which has significantly influenced the trajectory of this research.

Conflict of interest

The authors declared no conflict of interest.

Financial Support

None declared.

ORCID ID

Nisha B. Suryawanshi: <https://orcid.org/0009-0000-6802-6483>

Arati C. Sutar: <https://orcid.org/0009-0008-7875-5004>

Sumaiya S. Shaikh: <https://orcid.org/0009-0009-8999-5940>

Nitin T. Gore: <https://orcid.org/0000-0002-4927-6468>

Avinash S. Jondhale: <https://orcid.org/0000-0002-7621-0632>

Smita P. Chavan: <https://orcid.org/0000-0003-0084-0715>

Mahendra L. Ahire: <https://orcid.org/0000-0002-9690-9870>

REFERENCES

1. Chaudhary S, Kumar R. Folk medicinal plants in Ghaziabad district of Western Uttar Pradesh, India. *J Indian Bot Soc.* 2015;94:73-80.
2. Trinh XT, Long NV, Van Anh LT, Nga PT, Giang NN, Chien PN, *et al.* A comprehensive review of natural compounds for wound healing: targeting bioactivity perspective. *Int J Mol Sci.* 2022;23:9573.
3. Budovsky A, Yarmolinsky L, Ben-Shabat S. Effect of polyherbal preparations on wound healing. *Wound Repair Regen.* 2016;24:196-204.
4. Saxena S, Rawat DS, Rao PB. *Malvastrum coromandelianum* (L.) Garcke: an invasive weed with multiple ethnopharmacological properties. *Int J Pharm Phytochem Res.* 2020;12:16-22.
5. Sheema, Zafar S, Ullah N, Khan I, ud din G. Phytochemical and in vitro biological evaluation of roots of *Malvastrum coromandelianum* (L.) Garcke. *J Chem Soc Pak.* 2023;45:568-74.
6. Islam MA, Alam F, Solayman M, Khalil MI, Kamal MA, Gan SH. Dietary phytochemicals: natural swords combating inflammation and oxidation-mediated degenerative diseases. *Oxid Med Cell Longev.* 2016;2016:5137431.
7. Gaikwad S, Srivastava SK. Role of phytochemicals in perturbation of redox homeostasis in cancer. *Phytochem Rev.* 2021;20:1-20.
8. Deore AB, Chavan PN, Sapakal VD, Naikwade NS. Antidiabetic and antihyperlipidemic activities of *Malvastrum coromandelianum* Linn. leaves in alloxan-induced diabetic rats. *Int J PharmTech Res.* 2011;4:351-7.
9. Naseer S, Naseer A, Fatima M, Iqbal J, Kanwal S, Abbasi BA, *et al.* Ameliorative effect of green synthesized zinc oxide nanoparticles of *Malvastrum coromandelianum* Linn. on CCl₄-induced oxidative stress in rat liver. 2024.
10. Polito L, Bortolotti M, Maiello S, Battelli MG, Bolognesi A. Plants producing ribosome-inactivating proteins in traditional medicine. *Molecules.* 2016;21:1560.
11. Kirdeeva Y, Fedorova O, Daks A, Barlev N, Shuvalov O. How should the worldwide knowledge of traditional cancer healing be integrated with herbs and mushrooms into modern molecular pharmacology? *Pharmaceuticals.* 2022;15:868.
12. Sanghai DB, Kumar SV, Srinivasan KK, Aswatharam HN, Shreedhara CS. Pharmacognostic and phytochemical investigation of the leaves of *Malvastrum coromandelianum* (L.) Garcke. *Anc Sci Life.* 2013;33:39-44.
13. Saxena S, Rao PB. GC-MS screening of bioactive constituents and antioxidant profiling in an invasive weed, *Malvastrum coromandelianum* (L.) Garcke. *Pharma Innov J.* 2018;7:738-46.
14. Yadav A, Mohite S. Screening of *in vitro* anti-inflammatory and antibacterial assay of *Malvastrum coromandelianum*. *Int J Pharm Sci Res.* 2020;11:68-70.
15. Devi TS, Vijay K, Vidhyavathi RM, Kumar P, Govarthanam M, Kavitha T. Antifungal activity and molecular docking of phenol, 2,4-bis(1,1-dimethylethyl) produced by plant growth-promoting actinobacterium *Kutzneria* sp. strain TSII from mangrove sediments. *Arch Microbiol.* 2021;203:4051-64.
16. Villasenor IM, Angelada J, Canlas AP, Echegoyen D. Bioactivity studies on β -sitosterol and its glucoside. *Phytother Res.* 2002;16:417-21.
17. Prieto JM, Recio MC, Giner RM. Anti-inflammatory activity of β -sitosterol in a model of oxazolone-induced contact-delayed-type hypersensitivity. *Bol Latinoam Caribe Plantas Med Aromat.* 2006;5:57-62.
18. Gawade SP, Chandrashekar Rao MV. Antihepatotoxic activities of β -sitosterol isolated from fruits and leaves of *Coccinia indica*. *Indian J Pharm Educ Res.* 2012;46:4-8.
19. Shokry S, Hegazy A, Abbas AM, Mostafa I, Eissa IH, Metwaly AM, *et al.* Phytoestrogen β -sitosterol exhibits potent *in vitro* antiviral activity against influenza A viruses. *Vaccines.* 2023;11:228.
20. Nandi S, Nag A, Khatua S, Sen S, Chakraborty N, Naskar A, *et al.* Anticancer activity and other biomedical properties of β -sitosterol: bridging phytochemistry and current pharmacological evidence for future translational approaches. *Phytother Res.* 2024;38:592-619.
21. Cai M, Ma Y, Zhang W, Wang S, Wang Y, Tian L, *et al.* Apigenin-7-O- β -D-(6"-p-coumaroyl)-glucopyranoside treatment elicits neuroprotective effect against experimental ischemic stroke. *Int J Biol Sci.* 2016;12:42.
22. Aderogba MA, Madikizela B, McGaw LJ. Bioactive constituents from *Malvastrum coromandelianum* (L.) Garcke leaf extracts. *S Afr J Bot.* 2019;126:371-6.
23. Quan W, Ma S, Zhu Y, Shao Q, Hou J, Li X. Apigenin-7-O- β -D-(6"-p-coumaroyl)-glucopyranoside reduces myocardial ischaemia/reperfusion injury in an experimental model via regulating the inflammation response. *Pharm Biol.* 2020;58:80-8.
24. Feng YD, Ye W, Tian W, Meng JR, Zhang M, Sun Y, *et al.* Old targets, new strategy: Apigenin-7-O- β -D-(6"-p-coumaroyl)-glucopyranoside prevents endothelial ferroptosis and alleviates intestinal ischemia-reperfusion injury through HO-1 and MAO-B inhibition. *Free Radic Biol Med.* 2022;184:74-88.
25. Rahman MM, Ahmad SH, Mohamed MTM, Ab Rahman MZ. Antimicrobial compounds from leaf extracts of *Jatropha curcas*, *Psidium guajava*, and *Andrographis paniculata*. *Sci World J.* 2014;2014:635240.
26. Blondeau N, Lipsky RH, Bourourou M, Duncan MW, Gorelick PB, Marini AM. Alpha-linolenic acid: an omega-3 fatty acid with neuroprotective properties ready for use in the stroke clinic? *Biomed Res Int.* 2015;2015:519830.
27. Tian C, Gao X, Yang J, Guo Y, Wang H, Liu M. Chemical compositions, extraction technology, and antioxidant activity of petroleum ether extract from *Abutilon theophrasti* Medic leaves. *Int J Food Prop.* 2018;21:1789-99.
28. Cambiaggi L, Chakravarty A, Noureddine N, Hersberger M. The role of α -linolenic acid and its oxylipins in human cardiovascular diseases. *Int J Mol Sci.* 2023;24:6110.
29. Kim JK, Park SU. An update on the biological and pharmacological activities of diosgenin. *EXCLI J.* 2018;17:24-8.
30. Cai B, Zhang Y, Wang Z, Xu D, Jia Y, Guan Y, *et al.* Therapeutic potential of diosgenin and its major derivatives

- against neurological diseases: recent advances. *Oxid Med Cell Longev.* 2020;2020:3153082.
31. Huang N, Yu D, Wu J, Du X. Diosgenin: an important natural pharmaceutical active ingredient. *Food Sci Technol.* 2021;42:e94521.
 32. Ren QL, Wang Q, Zhang XQ, Wang M, Hu H, Tang JJ, *et al.* Anticancer activity of diosgenin and its molecular mechanism. *Chin J Integr Med.* 2023;29:738–49.
 33. Huang ZR, Lin YK, Fang JY. Biological and pharmacological activities of squalene and related compounds: potential uses in cosmetic dermatology. *Molecules.* 2009;14:540–54.
 34. Costa EV, Dutra LM, Nogueira PCDL, Moraes VRDS, Salvador MJ, Ribeiro LHG. Essential oil from the leaves of *Annona vepretorum*: chemical composition and bioactivity. *Nat Prod Commun.* 2012;7:265–6.
 35. Chowdhury RR, Fitch RW, Ghosh SK. Efficacy of phytol-derived diterpenoid immunoadjuvants over alum in shaping the murine host's immune response to *Staphylococcus aureus*. *Vaccine.* 2013;31:1178–86.
 36. Santos CCDMP, Salvadori MS, Mota VG, Costa LM, de Almeida AAC, de Oliveira GAL, *et al.* Antinociceptive and antioxidant activities of phytol *in vivo* and *in vitro* models. *Neurosci J.* 2013;2013:949452.
 37. Jiang Q. Natural forms of vitamin E: metabolism, antioxidant, and anti-inflammatory activities and their role in disease prevention and therapy. *Free Radic Biol Med.* 2014;72:76–90.
 38. Ghaneian MT, Ehrampoush MH, Jebali A, Hekmatimoghaddam S, Mahmoudi M. Antimicrobial activity, toxicity and stability of phytol as a novel surface disinfectant. *Environ Health Eng Manag.* 2015;2:13–6.
 39. Kim CW, Lee HJ, Jung JH, Kim YH, Jung DB, Sohn EJ, *et al.* Activation of caspase-9/3 and inhibition of epithelial mesenchymal transition are critically involved in antitumor effect of phytol in hepatocellular carcinoma cells. *Phytother Res.* 2015;29:1026–31.
 40. Pejin B, Ciric A, Glamoclija J, Nikolic M, Sokovic M. *In vitro* anti-quorum sensing activity of phytol. *Nat Prod Res.* 2015;29:374–7.
 41. Song Y, Cho SK. Phytol induces apoptosis and ROS-mediated protective autophagy in human gastric adenocarcinoma AGS cells. *Biochem Anal Biochem.* 2015;4:1.
 42. Islam MT, Ali ES, Uddin SJ, Shaw S, Islam MA, Ahmed MI, *et al.* Phytol: a review of biomedical activities. *Food Chem Toxicol.* 2018;121:82–94.
 43. Lima TL, Souza LB, Tavares-Pessoa LC, Santos-Silva AMD, Cavalcante RS, Araújo-Júnior RFD, *et al.* Phytol-loaded solid lipid nanoparticles as a novel anticandidal nanobiotechnological approach. *Pharmaceutics.* 2020;12:871.
 44. Claro-Cala CM, Grao-Cruces E, Toscano R, Millan-Linares MC, Montserrat-de la Paz S, Martin ME. Acyclic diterpene phytol from hemp seed oil (*Cannabis sativa* L.) exerts anti-inflammatory activity on primary human monocytes-macrophages. *Foods.* 2022;11:2366.
 45. Ebrahimi M, Farhadian N, Amiri AR, Hataminia F, Soflaei SS, Karimi M. Evaluating the efficacy of extracted squalene from seed oil in the form of microemulsion for the treatment of COVID-19: a clinical study. *J Med Virol.* 2022;94:119–30.
 46. Liao S, Omega SO, Börmel L, Kluge S, Schubert M, Wallert M, *et al.* Vitamin E and metabolic health: relevance of interactions with other micronutrients. *Antioxidants.* 2022;11:1785.
 47. Xiong Z, Liu L, Jian Z, Ma Y, Li H, Jin X, *et al.* Vitamin E and multiple health outcomes: an umbrella review of meta-analyses. *Nutrients.* 2023;15:3301.
 48. Du X, Ma X, Gao Y. The physiological function of squalene and its application prospects in animal husbandry. *Front Vet Sci.* 2024;10:1284500.
 49. Rosa GP, Seca AM, Pinto DC, Barreto MC. New phytol derivatives with increased cosmeceutical potential. *Molecules.* 2024;29:4917.
 50. Sheema, Jamal Q, Uddin M, Khan AR, Zafar S. *Malvastrum coromandelianum* mediated bio-fabrication of MgO:ZnO nanocomposites and their biological potential. *J Inorg Organomet Polym.* 2024;34:5636–50.
 51. Kovendan K, Fabiola M, Jebanesan A, Rajaganesh R. Green synthesis of *Malvastrum coromandelianum* fabricated AgNPs: anti-dengue and mosquitoicidal studies. *Inorg Chem Commun.* 2024;161:112067.
 52. Patra JK, Das G, Fraceto LF, Campos EVR, Rodriguez-Torres MP, Acosta-Torres LS, *et al.* Nano based drug delivery systems: recent developments and future prospects. *J Nanobiotechnol.* 2018;16:71.
 53. Bhupesh CS, Rohit B, Kalyani D. Analgesic activity of hydroalcoholic extract of aerial parts of *Malvastrum coromandelianum*. *Asian J Pharm Clin Res.* 2016;9:146–9.
 54. Reddy YSR, Venkatesh S, Suresh B. Antinociceptive activity of *Malvastrum coromandelianum*. *Fitoterapia.* 2001;72:278–80.
 55. Irudayaraj F, Fabiola J. Phytochemical analysis of *Malvastrum coromandelianum* and *Mimusops elengi*: mosquitoicidal properties of their leaf extract against dengue vector, *Aedes aegypti* (L.). *Int J Entomol Res.* 2022;7:171–80.
 56. Yadav A, Mohite S. Anticancer activity and in-silico ADMET analysis of *Malvastrum coromandelianum*. *Int J Pharma Sci Res.* 2020;11:71–3.
 57. Balekar N, Jain DK, Dixit PV, Bhadoriya SS. Evaluation of antiulcer activity of whole plant extract of *Malvastrum tricuspidatum* in experimental animals. *Iran J Pharmacol Therap.* 2012;11:53–9.
 58. Andrade-Cetto A, Heinrich M. Mexican plants with hypoglycemic effect used in the treatment of diabetes. *J Ethnopharmacol.* 2005;99:325–48.
 59. Balekar N, Parihar G, Jain DK, Gupta S. Antidiarrheal potential of ethanolic leaf extract of *Malvastrum tricuspidatum* in albino rats. *J Adv Pharm Edu Res.* 2014;4:233–9.
 60. Devi S, Kumar V. Comprehensive structural analysis of cis- and trans-tiliroside and quercetrin from *Malvastrum coromandelianum* and their antioxidant activities. *Arab J Chem.* 2020;13:1720–30.

HOW TO CITE THIS ARTICLE

Suryawanshi NB, Sutar AC, Shaikh SS, Gore NT, Jondhale AS, Chavan SP, *et al.* *Malvastrum coromandelianum* (L.) Garcke: A versatile weed with promising pharmacological applications. *J Phytopharmacol* 2025; 14(4):291-297. doi: 10.31254/phyto.2025.14410

Creative Commons (CC) License-

This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY 4.0) license. This license permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. (<http://creativecommons.org/licenses/by/4.0/>).