



PONGAMIA PINNATA: AN HEIRLOOM HERBAL MEDICINE

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Abstract

Pongamia pinnata, or Karanja or Indian Beech has been among the pillars of ancient Indian medicine and invaluable ecological resource. It is a popular Ayurvedic, Siddha, and Unani medicine, an indigenous Indian sub-continental plant with centuries of human experience of its vast curing power. A rich constitution of bioactive compounds, which are flavonoids, alkaloids, terpenoids, steroids and fatty acids, are found in the leaves, seeds and oil and basically characterizes its medicinal image. More recent scientific findings support these old claims and demonstrate that P. pinnata is a good antioxidant, anti-microbial agent, anti-inflammatory, anti-glycemia and anti-wound-healing agent. Furthermore, its application is significantly broader since farmers apply it to improve the soil, environmentalists to restore the land and engineers to make the oil a possibly promising source of clean biofuel. The current paper will build a progressive argument of how P. pinnata has evolved into a modern medicine and environmental resource as well as a traditional medicine and how intergenerational ethnomedicinal knowledge is connected to the scientific paradigm of drug development.

Keywords: Pangamia, Herbal medicine, Drug development, Bioactive compound

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Introduction

The recent shift in health paradigms in the world during the last several decades was towards natural and plant-based products as a result of the necessity to possess the holistic health. It is not a trend, but a legitimate response to the factual clinical problems of the negative side effects of synthetic drugs, and the rising epidemic of

antibiotic resistance. Patients and medical practitioners are also in the hunt to pursue less prone to obsolescence and safer and sustainable therapeutic options even as most modern day pharmaceuticals are highly characterized by obsolescence. *Pongamia pinnata* is a widely growing and culturally significant plant that has developed into a highly fascinating research

topic among all the numerous healing plants. The medicinal folk practitioners had long surpassed scientists and pharmacologists who have been able to slowly catch up with the comprehensive and well-developed therapeutic profile of the plant, and validate it scientifically.

Pongamia pinnata is a plant that is very much embedded in the life and culture of rural and urban regions of the Indian sub continent. It is also commonly referred to as Karanja in most regions and continues to be stock medicine in most households medicine cabinets like it was the case with the previous generations. It is a medium and fast growing evergreen tree with a broad canopy, smooth grayish and pinkish-white fragrant flowers. The most outstanding botanical characteristic about it is that it has immense ecological power that makes it thrive in extremely hostile environments such as saline soils, waterlogged and even the wastelands that are so much degraded. This longevity not only offers the availability of the raw medicinal products at all times but also makes the plant an ideal candidate to the enormous afforestation and sustainable agriculture programs.

Despite its wide application in the past and the ecological availability of *pongamia pinnata*, there exist gaps in the procedure of complete utilization of the plant given a variety of factors.

First, there are no standard extraction and formulation methods, which leads to large-scale variability of phytochemical yield on a batch-to-batch basis, which severely affects clinical reproducibility and subsequent regulatory acceptance. Second, the recent technologies of multi-omics and stringent algorithmic classification are not necessarily included in the traditional research methods and the complex synergistic mechanisms of the different bioactive mixtures of the plant are largely unexplored and are not associated with the pharmacological needs of the time. Such methodological flaws do not allow utilizing the full therapeutic potential of the plant and restrict its implementation into mainstream allopathic medical services.

In order to address these prevailing obstacles and to improve the position of this heirloom plant, this paper suggests a contemporary scientific method. Particularly, our work makes the following principal contributions:

To actively mediate between the traditional ethnomedicinal use of *Pongamia pinnata* and modern scientific information, a detailed taxonomic, phytochemical, and pharmacological characterization of this plant are presented in this paper.

We propose systematic methodology of extraction, stabilization, evaluation, and hypothetical clinical validation of the bioactive compounds of this natural product, which is

grounded on the recent advances in the field of network pharmacology and metabolomics.

Related Work

The scientific evaluation of traditional herbal medicines has recently benefited from interdisciplinary methodologies that range from advanced biomaterial stabilization to predictive computational modeling. We categorize the related literature into three primary domains that inform our modern approach to *Pongamia pinnata*: carrier-based formulation strategies, network pharmacology for target identification, and advanced spectral classification techniques.

Delivery and Stabilization of Bioactive Compounds

The preservation of the stability and bioavailability of complex phytochemicals including phenolic compounds and pigments is one of the greatest challenges in herbal medicine. The new trends have demonstrated the usefulness of the use of edible polysaccharides such as starch, maltodextrin, alginate, pectin, and chitosan as food and nutraceutical stabilizers and carriers (Oliveira et al., 2025). These biopolymers make sure that active substances are not degraded when storing them or passing through the gastrointestinal tract, which increases the bio accessibility of the substance (Oliveira et al., 2025). The key

benefit of this technique is its biocompatibility and scalability, and one of its potential weaknesses is that the wall material must be optimized to provide the required hydrophobicity of the encapsulated extract. The evaluation of these polysaccharide encapsulation protocols as compared to our work is a highly encouraging procedure of stabilizing the lipid-rich, flavonoid-heavy extracts of *P. pinnata*, to ensure the reliability of therapeutical delivery.

Network Pharmacology and Target Prediction

Network pharmacology and quantitative structure-activity relationship (QSAR) modeling are in silico approaches that are increasingly applied to elucidate the complicated multi-target mechanisms of herbal compounds. A case in point is that recent studies have been in a position to utilize network pharmacology and graph deep learning to uncover the molecular pathways of xanthonones of *Garcinia cowa* against cancer cells (Son et al., 2025). On the same note, the computational models have been utilized to unravel the active compounds and target pathways (like HSPA1A/BST2) of the traditional Chinese medicine capsules to intervene in leukopenia (Zhang et al., 2025). Network pharmacology is the most robust since it is able to rapidly screen the complicated herbal matrices in addition to preempting synergistic biological pathways without

exposing the preliminary test to the intensive in vitro testing (Zhang et al., 2025)(Son et al., 2025). Nevertheless, these calculational learnings are always theoretical in nature and need to be strictly experimented. This is exactly what network pharmacology entails that we are going to integrate to map the molecular interactivity of *P. pinnata* flavonoids, such as karanjin and pongamol.

Advanced Analytical and Spectral Classification

The geographic origin and minor variations in the chemical profiles also require very precise methods of classification to verify the standardization of herbal medicines. Electronic nose technologies, which are accompanied by sophisticated machine learning algorithms like conformal prediction with shrunken centroids (CPSC), have been highly successful in estimating the trustworthiness of prediction and overcoming the issue of dimensionality of features in the classification of herbals (Liu et al., 2021). Furthermore, the conformal prediction methods alongside data augmentation have considerably improved the discrimination error of unlabelled data of herbal medicines (Liu et al., 2021). Techniques that extract both time- and spectral-based data of electronic noses are also known to be more successful when it comes to determining the specific origins of therapeutic plants (Liu et al., 2021). At the same time, Nuclear Magnetic Resonance (NMR) based

pharmaco-metabolomics is becoming a flexible method to disclose the altered metabolic markers in the response to conventional herbal treatment, and it can be used to supplement genomics and proteomics (Kumar et al., 2016). Although these sophisticated analysis tools consume a lot of computing power and special equipment, they offer an unparalleled amount of quality control which we implement in our assessment of *P. pinnata*. In addition, the bacteriostatic activity of herbal preparations has been assessed using the methodology that demonstrated that the preparation procedure has a significant impact on the pathogen inhibition of *E. coli* and other pathogens (Ma et al., 2017), which is essential to our pharmacological analysis phase.

Method/Approach

To transition *Pongamia pinnata* from a traditionally utilized heirloom remedy into a rigorously validated modern pharmaceutical asset, we outline a structured methodological framework. This framework systematically addresses the core challenges of standardization, phytochemical degradation, and mechanistic ambiguity. Our approach combines targeted extraction, computational target prediction, polysaccharide-based stabilization, and a comprehensive biological evaluation plan.

Step 1: Botanical Collection and Standardized Extraction

The initial procedure is the accurate geographical tagging and harvesting of *P. pinnata* parts, namely the seeds, leaves, and bark, and then a solvent-mediated extraction procedure. Since the therapeutic efficiency of natural isolates may greatly rely on the structural resistance to degradation, it is essential to apply optimized methods of breakdown and extraction, a concept also reported in the context of RNA isolation of recalcitrant natural isolates (Guez et al., 2020). We suggest that a sequential maceration and Soxhlet extraction procedure with solvents of different polarities (hexane, ethyl acetate, and methanol) be used to systematically extract the fixed oils and the flavonoids and alkaloids. The extraction conditions such as temperature, time and amount of solvent-solid ratio should be closely observed to reduce the exposure of heat sensitive triterpenes and glycosides to thermal degradation.

Step 2: Phytochemical Profiling and Network Pharmacology

Once the standardized extracts are obtained, the second phase focuses on identifying the active constituents and their theoretical physiological targets. High-Performance Liquid Chromatography coupled with Mass Spectrometry (HPLC-MS) is employed to quantify marker compounds such as karanjin,

pongamol, and pongapin. Following the quantification, we employ an *in silico* network pharmacology approach to map these identified phytoconstituents against known human disease pathways. By utilizing databases to cross-reference *P. pinnata* flavonoids with inflammatory and oxidative stress targets, we can construct a compound-protein-disease interaction network. This design choice dramatically accelerates the discovery phase by narrowing down the probable mechanisms of action before entering resource-intensive biological assays.

Step 3: Polysaccharide-Based Formulation

To address the poor aqueous solubility and potential oxidative degradation of Karanja oil and its isolated bioactive fractions, we propose a microencapsulation strategy. Utilizing edible polysaccharides such as alginate and chitosan as wall materials, the lipid-rich extracts are formulated into stable nano-emulsions (Oliveira et al., 2025). The rationale behind this design choice is twofold: it significantly enhances the gastrointestinal survivability of the flavonoids and masks the inherently astringent taste and odor of the raw oil. These polysaccharide carriers act as a robust delivery vehicle, ensuring that the therapeutic payload maintains its structural integrity from the point of administration to the physiological target site (Oliveira et al., 2025).

Step 4: Hypothetical Evaluation Plan

We outline a rigorous, hypothetical evaluation plan to validate the efficacy and safety of the engineered formulations across *in vitro* and *in vivo* models. First, the antioxidant capacity of the encapsulated extracts will be assessed using standard DPPH and ABTS radical scavenging assays. Next, the broad-spectrum antimicrobial activity will be tested using plate culture methods to analyze bacteriostatic effects against multidrug-resistant strains, akin to protocols used for evaluating traditional herbal medicines against *E. coli* (Ma et al., 2017). Furthermore, an *in vivo* hypothetical murine model will be established to evaluate anti-inflammatory and wound healing properties, utilizing topical applications of the stabilized Karanja formulations to measure tissue regeneration rates and the suppression of pro-inflammatory cytokines.

Pseudo-Algorithm for Extract Processing

To ensure maximum reproducibility in industrial applications, the sequential processing of *P. pinnata* can be summarized in the following algorithmic pipeline:

Initialize batch raw material (\$R\$) from verified geographical origins.

Subject \$R\$ to sequential solvent extraction to yield crude fractions (\$C_1, C_2, \dots, C_n\$).

Perform HPLC-MS on all fractions to output phytochemical vector \$V\$.

If the concentration of primary marker (karanjin) in \$V < \$ threshold, reject batch.

Else, proceed to encapsulate fraction using polysaccharide matrix to generate final formulation \$F\$.

Submit \$F\$ to *in vitro* quality and stability control benchmarks.

Discussion

The practical implications of successfully standardizing and formulating *Pongamia pinnata* extracts are vast, bridging critical gaps in both modern healthcare and environmental sustainability. From a clinical deployment perspective, introducing a reliable, highly bioavailable herbal formulation derived from *P. pinnata* offers an alternative mechanism to manage chronic inflammatory conditions and superficial microbial infections, potentially reducing the over-reliance on synthetic topical antibiotics. Ecologically and industrially, incentivizing the cultivation of this resilient tree for pharmaceutical-grade extracts inherently supports large-scale wasteland reclamation and provides communities with supplementary economic avenues, including the conversion of the residual seed biomass into biodiesel. Thus, the integration of this plant into the global

supply chain represents a multifaceted victory for agro-economics and green healthcare.

Despite these promising implications, the proposed methodologies and general utilization of *P. pinnata* face several notable limitations and potential failure modes. First, the inherent variability in phytochemical composition driven by disparate environmental factors—such as soil salinity, rainfall, and temperature—can make batch-to-batch standardization extremely difficult, even with advanced extraction pipelines. Second, there are documented toxicological concerns; while generally safe at therapeutic doses, the excessive consumption or improper processing of Karanja seed oil can induce mild to moderate toxicity, necessitating stringent dose control. Third, the transition from successful laboratory-scale nano-emulsions to large-scale industrial pharmaceutical manufacturing poses significant engineering and economic challenges, particularly regarding the cost of sophisticated polysaccharide matrices and multi-omics profiling tools.

Furthermore, integrating heirloom traditional medicines into global commercial markets raises important ethical considerations and potential risks. Primary among these is the risk of biopiracy, wherein indigenous knowledge regarding the therapeutic applications of *P.*

pinnata is exploited by corporate entities without providing equitable biological or financial compensation to the native communities who safeguarded this knowledge. Additionally, there is a distinct ecological risk: if the plant becomes overly lucrative for its biofuel and pharmaceutical yields, it could inadvertently encourage large-scale monoculture farming, which subsequently threatens local biodiversity and disrupts pre-existing agricultural ecosystems. Establishing strict regulatory frameworks for equitable benefit-sharing and sustainable agroforestry is essential to mitigate these profound ethical risks.

Looking ahead, future work must focus on actively addressing these existing limitations to ensure the safe and widespread adoption of *P. pinnata*. One critical area for future research is the execution of comprehensive, large-scale human clinical trials to definitively establish the long-term safety profile and precise pharmacokinetic behavior of the isolated active constituents. Another vital direction involves the exploration of advanced, targeted nanotechnology-based drug delivery systems—beyond basic polysaccharide encapsulation—that can deliver specific flavonoids like pongamol directly to inflamed joints or compromised cellular tissues. By pursuing these future research avenues, the scientific community can cement the plant's role as a staple of modern integrative medicine.

Conclusion

Pongamia pinnata represents a remarkable confluence of traditional ethnomedicinal wisdom and cutting-edge scientific innovation. As an heirloom herbal medicine, its diverse phytochemical backbone—rich in flavonoids and essential fatty acids—has historically provided holistic relief for a wide spectrum of skin, digestive, and inflammatory ailments. By applying modern analytical rigor, advanced biomaterial stabilization, and computational pharmacology, we are now uniquely positioned to validate and elevate these ancient remedies. The structured methodologies proposed in this paper offer a tangible pathway to overcome historical barriers of toxicity and standard variability.

Ultimately, the successful integration of *P. pinnata* into modern pharmacopeia does more than just expand our medical repertoire; it actively promotes environmental sustainability and socio-economic resilience. The plant's ability to concurrently serve as a therapeutic agent, a natural soil enhancer, and a sustainable biofuel source highlights its unparalleled versatility. With continued scientific exploration, ethical commercialization, and rigorous clinical validation, *Pongamia pinnata* will undoubtedly play a significant and enduring role in addressing the dual global challenges of healthcare equity and ecological preservation. Furthermore, technological

advances such as electronic nose systems now enable more precise characterization and quality control of herbal medicines, including *P.*

References

- Oliveira, Liliane Siqueira de, Silva, Davi Vieira Teixeira da, Trindade, Lucileno Rodrigues da, Baião, Diego dos Santos, Almeida, Cristine Couto de, Ferreira, Vitor Francisco, & Paschoalin, Vania Margaret Flosi (2025). *edible polysaccharides as stabilizers and carriers for the delivery of phenolic compounds and pigments in food formulations*.
<https://arxiv.org/pdf/2511.07264v1>
<https://arxiv.org/pdf/2511.07264v1>
- Son, Nguyen Manh, Vang, Pham Huu, Dung, Nguyen Thi, Thao, Nguyen Manh Ha. Ta Thi, Thuy, Tran Thi Thu, & Giang, Phan Minh (2025). *In silico study on the cytotoxicity against Hela cancer cells of xanthenes bioactive compounds from Garcinia cowa: QSAR based on Graph Deep Learning, Network Pharmacology, and Molecular Docking*.
<https://arxiv.org/pdf/2508.10117v1>
<https://arxiv.org/pdf/2508.10117v1>
- Zhang, Dingfan, Huang, Congshu, Zhou, Lei, Wang, Boyang, Zhou, Wei, Xia, Tiantian, Shen, Pan, Li, Shao, & Gao, Yue (2025). *Network Pharmacology Reveals HSPA1A/BST2 as Potential Targets of Ci Bai Capsule's Active Compounds Intervening in*

Leukopenia.

<https://arxiv.org/pdf/2506.12107v1>

<https://arxiv.org/pdf/2506.12107v1>

- Liu, Li, Zhan, Xianghao, Yang, Xikai, Guan, Xiaoqing, Wu, Rumeng, Wang, Zhan, Luo, Zhiyuan, Wang, You, & Li, Guang (2021). *CPSC: Conformal prediction with shrunken centroids for efficient prediction reliability quantification and data augmentation, a case in alternative herbal medicine classification with electronic nose.* <https://arxiv.org/pdf/2108.00777v1>
<https://arxiv.org/pdf/2108.00777v1>
- Liu, Li, Zhan, Xianghao, Wu, Rumeng, Guan, Xiaoqing, Wang, Zhan, Zhang, Wei, Pilanci, Mert, Wang, You, Luo, Zhiyuan, & Li, Guang (2021). *Boost AI Power: Data Augmentation Strategies with unlabelled Data and Conformal Prediction, a Case in Alternative Herbal Medicine Discrimination with Electronic Nose.* <https://arxiv.org/pdf/2102.03088v3>
<https://arxiv.org/pdf/2102.03088v3>
- Liu, Li, Zhan, Xianghao, Duan, Ziheng, Wu, Yi, Wu, Rumeng, Guan, Xiaoqing, Wang, Zhan, Wang, You, & Li, Guang (2021). *Classifying herbal medicine origins by temporal and spectral data mining of electronic nose.* <https://arxiv.org/pdf/2104.06640v3>
<https://arxiv.org/pdf/2104.06640v3>
- Kumar, Dinesh, Rawat, Atul, Dubey, Durgesh, Kumar, Umesh, Keshari, Amit K, Saha, Sudipta, & Guleria, Anupam (2016). *NMR based Pharmaco-metabolomics: An efficient and agile tool for therapeutic evaluation of Traditional Herbal Medicines.* <https://arxiv.org/pdf/1602.02492v1>
<https://arxiv.org/pdf/1602.02492v1>
- Ma, Li, Chen, Shuangjie, & Yang, Yongguang (2017). *Analysis of Bacteriostatic Effect of Chinese Herbal Medicine Against E.coli.* <https://arxiv.org/pdf/1706.06868v1>
<https://arxiv.org/pdf/1706.06868v1>
- Guez, Jean-Sebastien, Coutte, François, Drucbert, Anne-Sophie, Danzé, Pierre-Marie, & Jacques, Philippe (2020). *Resistance of the cell wall to degradation is a critical parameter for isolation of high quality RNA from natural isolates of Bacillus subtilis.* Archives of Microbiology, Springer Verlag, 2009, 191, pp.669 - 673. <https://doi.org/10.1007/s00203-009-0487-6>
<https://doi.org/10.1007/s00203-009-0487-6>