

Review Article**Curcuma caesia Roxb. (Black Turmeric): A brief Literature review****Ritesh Tiwari¹, Suresh Kumar Dev², Chetna Baregama², Ajay Kumar Shukla^{3*}, Akhil Mangal⁴, Rajesh Babu Vemula⁵, Yogesh Kumar Apurva², Ayush Garg², Mohammad Junaid Alam Mansoori², Vijay Kumar Bansal⁶, Mohini Vishwas⁵, Vaibhav Rathore⁷**¹Pharmaceutical Science, Madhyanchal Professional University, Bhopal, Madhya Pradesh, India²Venkateshwar Institute of Pharmacy, Sai Tirupati University, Udaipur, Rajasthan-313015, India³Institute of Pharmacy, Dr. Rammanohar Lohia Avadh University Uttar Pradesh, India⁴Bhai Gurudas College of Pharmacy, Sangrur, Punjab-148002, India⁵Faculty of Pharmacy, Pacific Academy of Higher Education and Research University, Udaipur, Rajasthan-313001, India⁶Lachoo Memorial College of Science & Technology "Pharmacy Wing" Jodhpur, Rajasthan- 342001, India⁷Teerthanker Mahaveer College of Pharmacy, Teerthanker Mahaveer University, Delhi Road, NH 9, Moradabad, Uttar Pradesh-244102, India

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Abstract

Curcuma caesia Roxb., commonly known as black turmeric, is a medicinally important rhizomatous herb belonging to the family Zingiberaceae and widely distributed in Northeast and Central India. Traditionally, the species has been valued in Ayurveda, Siddha, and folk medicine for its diverse therapeutic properties, including anti-inflammatory, analgesic, antimicrobial, antiasthmatic, wound healing, and neuroprotective actions. The article aims to bridge traditional knowledge with scientific evidence to encourage further exploration of black turmeric as a potential source for novel drug discovery and therapeutic development. This review provides a comprehensive overview of the botanical characteristics, ethnomedicinal relevance, phytochemistry, pharmacological activities, and future research prospects of *Curcuma caesia*. The plant is characterized by its distinctive bluish-black rhizome, rich aromatic profile, and the presence of bioactive constituents such as curcumin, camphor, ar-turmerone, germacrone, and various essential oils. In vitro and in vivo studies have demonstrated significant antioxidant, cytotoxic, antidiabetic, antiepileptic, adaptogenic, and anti-ulcer activities, suggesting promising applications in modern therapeutics. Despite its high medicinal value, *Curcuma caesia* remains underutilized and is categorized as a threatened species in several regions due to overharvesting and limited cultivation practices.

Keywords: Black turmeric; *Curcuma caesia*, phytochemistry, pharmacological activities, conservation strategies

Introduction

Plants play a vital role in the food chain, offering a wealth of nutrients and biochemical compounds beneficial to humans and other organisms. The use of herbal plant products is growing in many segments of the population (Dev et al., 2017). One such plant, *Curcuma caesia* Roxb., belonging to the Zingiberaceae family and commonly known as black turmeric, is extensively

utilized in Ayurvedic medicine and traditional remedies. Its leaves and rhizomes are rich in essential oils, including camphor, ar-turmerone, (Z)- β -ocimene, and others. Research indicates diverse biological activities of *C. caesia*, such as antioxidant, antimicrobial, and anti-inflammatory properties, attributed to its bioactive components. This plant's chemical makeup holds promise for various industries, including health, food, and cosmetics. Recent studies have focused on its in vitro and in vivo activities and ongoing clinical investigations. Summarizing these findings offers insights into potential health benefits and mechanisms, facilitating the development of drugs, functional foods, and cosmetics (Ibrahim, 2023).

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Used as functional food

The study investigated the health benefits of *Curcuma* spp. rhizomes, including *Curcuma caesia*, *Curcuma zedoaria*, and *Curcuma aeruginosa*, through antioxidant, anti-inflammatory, and anti-tumor cell proliferation assays. *C. caesia* (black turmeric) demonstrated the most significant biological activities, including inhibition of lipid peroxidation (LPO) and cyclooxygenase (COX-1 and COX-2) enzymes, as well as tumor cell growth. Hexane and methanolic extracts of *C. caesia* showed notable inhibition of LPO and COX-2 enzymes. Eleven terpenoids were identified from the extracts. The antioxidant assay revealed comparable activity to positive controls, and both extracts and isolates showed inhibition of COX enzymes. These findings support the traditional medicinal use of *C. caesia* rhizome (Liu, 2013).

Pharmacognostic parameters of *Curcuma caesia*

Ethnomedicinal practices often utilize *Curcuma* species for treating ailments, yet *Curcuma caesia* Roxb. remains largely unexplored. This study aims to establish pharmacognostic standards for evaluating *Curcuma caesia* Roxb. by examining its morphology, microscopy, physicochemical properties, and phytochemical profiles. Salient diagnostic features, major chemical constituents, extractive values, and other relevant characteristics are documented for comprehensive analysis (Paliwal, 2011).

Phytochemical Constituents of *Curcuma caesia* Roxb:

1. Pakkirisamy et al. (2017) studied phytochemical screening, GC-MS and FT-IR analysis of methanolic extract of *Curcuma caesia* Roxb (black turmeric) were identified compounds as: α -Santalol, retinal, ar-turmerone, alloaromadendrene, megastigma-3,7(E),9-triene, etc (Pakkirisamy et al., 2017).
2. Baghel et al. (2013) studied pharmacological activities of *Curcuma caesia* Roxb. a review of traditional uses and phytochemical study reported major constituents include camphor, ar-turmerone, (Z)-ocimene, ar-curcumene, 1,8-cineole, elemene, borneol, bornyl acetate, and curcumene.
3. Leela et al., (2024) *Curcuma caesia* Roxb. were updated phytochemicals and pharmacological properties. They reports major rhizome oil constituents: 1,8-cineole, camphor, ar-turmerone, linalool, ocimene, ar-curcumene, zingiberol, curzerenone and tropolone.
4. Lenka et al. (2025) unlocked terpenoid treasures of rhizome and leaf volatiles of *Curcuma caesia* Roxb through GC-GC TOFMS analysis. They identified 151 volatile constituents in rhizome and leaf oils, including curzerenone, eucalyptol, curzerene, epicurzerenone, camphor and ar-turmerone.

5. Jose et al. (2021) reported phytochemicals, antioxidants and antimicrobial components in leaf extracts of *Curcuma caesia* Roxb. They reported diverse phytochemicals in leaf extracts (phenolics, flavonoids, alkaloids, etc.).

6. Lawand et al. (2013) studied about comparison of *Curcuma caesia* Roxb. with other *Curcuma* species by HPTLC. They identified phenolics, alkaloids, tannins, and terpenoids; noted the presence of camphor in *C. caesia*.

7. Pandey et al. (2025) studied that medicinal potential of *Curcuma caesia* Roxb.: phytochemical composition and TLC profile of hydroalcoholic rhizome extract for identification of flavonoids, phenolics, alkaloids, terpenoids, saponins and glycosides through TLC profiling.

Analytical Advancements in *Curcuma* Species

Recent research has prioritized the development of precise methods to quantify active compounds in the *Curcuma* genus. A validated High-Performance Thin-Layer Chromatography (HPTLC) method demonstrated that curcumin levels vary significantly across species with Soxhlet extraction yielding higher concentrations than sonication (Gangal et al., 2025). Notably, while curcumin is abundant in *C. longa*, it is absent in *C. caesia*.

To address this, researchers have turned to 1H-NMR and GC \times GC-TOFMS to map the complex profile of *C. caesia*, identifying over 151 compounds, including thermolabile sesquiterpenes like curzerenone and zederone (Lenka et al., 2025; Mahanta, 2020).

Therapeutic Applications of *Curcuma caesia*

Curcuma caesia exhibits diverse pharmacological and preservative properties:

Anticancer Potential: Derivatives from black turmeric have shown promise in modulating **Mis-Match Repair (MMR)** pathways in hematologic malignancies (Bhattacharya, 2021). Furthermore, gold nanoparticles synthesized from *C. caesia* extracts demonstrate selective toxicity against aggressive breast cancer cells (MDA-MB-231) while remaining safe for healthy cells (Das et al., 2024).

Biotransformation: The endophytic fungus *Ovatospora brasiliensis* found in the plant can successfully convert curcumin into **calebin-A**, a bioactive compound found in mature rhizomes (Majeed, 2019).

Food Preservation: Essential oils from the rhizome serve as effective edible coatings for fruits like mangoes, reducing weight loss and extending shelf life by preserving firmness and antioxidant levels (Kotha et al., 2025).

The integration of advanced HPTLC and NMR techniques has standardized the quality assessment of *Curcuma* species, highlighting the chemical uniqueness of *C. caesia*. While it lacks curcumin, its rich sesquiterpene profile and the efficacy of its nanoparticle formulations position it as a potent candidate for targeted cancer therapies and natural food preservation.

Bhattacharya, (2021) reported computational analysis of Curcuma derivatives in MMR-pathway-driven cancers.

Das et al. (2024) prepared Gold nanoparticle formulations of *C. caesia* in breast cancer therapy.

Gangal et al. (2025) performed HPTLC technique for routine analysis of curcumin in *Curcuma* species.

Kotha et al. (2025) identified Tropolone-related compounds in *C. caesia* for mango shelf-life extension.

Lenka et al. (2025) investigated GC×GC-TOFMS profiling of *C. caesia* rhizomes and leaves.

Mahanta (2020) performed 1H-NMR identification of chemical markers in black turmeric essential oil.

Majeed (2019) reported Biotransformation of curcumin to calebin-A via endophytic fungi.

Rathi et al. (2024) performed pharmacognostical and elemental screening of *C. caesia* rhizomes

Genetic Diversity and Evolutionary Insights

Molecular characterization of Zingiberaceae species in eastern India has revealed high levels of genetic polymorphism. Using a combination of RAPD, ISSR, and SSR markers, researchers identified significant genetic distinction between wild and cultivated species, with *C. caesia* exhibiting the highest number of genetic loci (Mohanty et al., 2014).

Complementary meiotic analysis indicates that *Curcuma* species function as an allopolyploid complex. Species are typically categorized into two groups based on chromosome counts:

- **Group I ($2n = 42$):** Predominantly displays bivalent configurations, seen in species like *C. comosa* and *C. mangga*.
- **Group II ($2n = 63$):** Shows complex trivalents and quadrivalents, characteristic of *C. caesia* and *C. longa*.

The high basic chromosome number ($x = 21$) suggests that the genus likely evolved through hybridization between species with $2n = 24$ and 18 , forming dibasic amphidiploids (Lamo & Rao, 2017).

Agricultural Development and Stability

To address the endangered status of *C. caesia*, systematic selection trials have led to the identification of high-yielding, stable genotypes. **Jor Lab KH-2** emerged as a superior strain, maintaining high yield consistency across diverse environments

(Lal et al., 2022). This variety represents the first registered high-yielding cultivar of Black Turmeric, providing a vital resource for commercial cultivation and conservation (Lal et al., 2022).

Environmental Influence of UV-B Radiation

Ultraviolet-B (UV-B) radiation acts as both a stressor and a metabolic regulator. Exposure impacts carbon-nitrogen dynamics, often leading to:

- **Biomass Shifts:** Increased tuber biomass but reduced rhizome weight (Jaiswal & Agrawal, 2022).
- **Phytochemical Induction:** Enhanced production of essential oils and active compounds like D-camphor and 1,8-cineole.
- **Species-Specific Responses:** *C. caesia* typically shows higher enzymatic stimulation (e.g., PAL and CHI) and a greater increase in essential oil (16%) compared to *C. longa* (9%) under elevated UV-B (Jaiswal & Agrawal, 2021). The integration of molecular markers and cytogenetic data confirms that *Curcuma* is a genetically diverse, polyploid genus shaped by hybridization (Lamo & Rao, 2017; Mohanty et al., 2014). While environmental factors like UV-B radiation can be leveraged to boost secondary metabolites, the development of stable genotypes like Jor Lab KH-2 is essential for sustainable pharmaceutical production and species preservation (Jaiswal & Agrawal, 2022; Lal et al., 2022).

Therapeutic Potential of *Curcuma caesia* (Black Turmeric)

Bioactive compounds from the plant origin are considered more secure as they produce less harmful metabolites (Srivastava et al. 2022). *Curcuma caesia* Roxb., commonly known as **Black Turmeric**, is a rare medicinal herb of the Zingiberaceae family. Its distinct bluish-black rhizomes contain a high concentration of bioactive compounds, including curcuminoids, essential oils (camphor, eucalyptol), and sesquiterpenes, which contribute to its broad pharmacological profile (Borah et al., 2021; Chande et al., 2023).

Biological Activities

Antioxidant & Anti-inflammatory: Rhizome extracts exhibit potent free radical scavenging abilities. Methanol and chloroform extracts are particularly effective due to high phenolic and flavonoid content, which helps mitigate oxidative stress and inflammation (Ain Ibrahim, 2023; Reenu, 2015).

Antimicrobial & Antibacterial: The essential oil and extracts demonstrate significant activity against various

pathogens, including *B. subtilis*, *S. aureus*, and *M. smegmatis*. Supercritical fluid extraction (SFE) has been shown to enhance its antimycobacterial potential, validated through molecular docking studies (Chaturvedi, 2020; Paw et al., 2020).

Antidiabetic Potential: Recent studies highlight its ability to inhibit alpha-glucosidase and alpha-amylase enzymes, suggesting a role in managing postprandial glucose levels (Singh et al., 2025).

Genotoxicity: Studies indicate that *C. caesia* essential oil has minimal genotoxic effects, supporting its safety for potential commercial and pharmaceutical applications (Paw et al., 2020).

Skin Protection and Wound Healing

Traditionally used to treat cuts and bruises, *C. caesia* contains specific **serine proteases** that exhibit procoagulant and fibrinolytic properties. These enzymes facilitate hemostasis (halting bleeding) and initiate the healing process by promoting collagen deposition and tissue remodeling (Shivalingu, 2016). Additionally, its extracts are utilized in herbal sunscreens for their photoprotective qualities, improving skin hydration and viscoelasticity (Saraf, 2012).

Neuroprotective Roles

Research into brain function indicates that Zederone, a compound found in *C. caesia*, can enhance memory and reduce amyloid plaque formation in dementia models (Borah, 2022). Furthermore, the rhizome's essential oil and specific fractions have shown: **Anxiolytic effects** (reducing anxiety).

Antidepressant properties by modulating neuroinflammation and oxidative stress (Borah et al., 2021).

Curcuma caesia is a versatile therapeutic agent with validated antioxidant, antimicrobial, and neuroprotective properties. Its ability to aid in wound healing and metabolic regulation makes it a valuable candidate for future phytopharmaceutical development.

Anti-diabetic activity

Curcuma caesia has exhibited due to its potent antioxidant and anti-inflammatory properties, *C. caesia* has been found ability to alleviate the oxidative stress and neuronal inflammation that drive nerve damage in diabetic patients (Grover 2019).

As muscle relaxant

In respiratory health, hydroalcoholic extracts of *Curcuma caesia* act as an effective smooth muscle relaxant. The mechanism behind of this activity through the modulating calcium channels rather than interacting with specific biochemical receptors, suggesting a non-specific mechanism that could benefit asthma treatments (Arulmozhi 2006).

Anticancer property

The anticancer efficacy of *Curcuma caesia* nanoparticles was

assessed on HT-29 human colon cancer cells via the sulforhodamine B (SRB) assay. The yields for *C. longa*, *C. aromatica*, and *C. caesia* were 11.34g, 15.45g, and 12.67g, respectively, with resulting nanoparticles exhibiting smooth, spherical morphology. The nanoparticles demonstrated varied degrees of cytotoxicity against HT-29 cells, indicating potential as cancer therapeutics with low toxicity (Jain 2023). Nanoparticles are biodegradable, non-toxic lyotropic systems show promise for use in eco-friendly therapeutic, cosmetic, and advanced material applications (Gangal et al., 2025). It has been found that phytochemicals derived from *Aristolochia tagala* (AT) and *Curcuma caesia* (CC) possess significant therapeutic potential, specifically acting as anticancer agents by modulating inflammatory pathways (Hadem 2015).

Curcuma caesia Roxb., popularly known as Black Turmeric, is a medicinal plant native to India with a robust history in traditional medicine for treating tumors, asthma, and inflammation (Borah et al., 2020; Karmakar et al., 2013). Recent scientific evaluations have validated its therapeutic potential through various bioactive fractions and molecular mechanisms.

Researchers have identified several key chemical constituents through bioassay-guided fractionation. Hexane and chloroform extracts have shown efficacy against breast cancer cell lines (MCF-7 and MDA-MB-231). Specifically, compounds such as germacrone, zerumbone, and curcuzederone have been isolated; curcuzederone, in particular, significantly inhibits the migration of MDA-MB-231 cells, suggesting potential in preventing metastasis (Al-Amin, 2021).

Antioxidant and Antimutagenic Properties

C. caesia exhibits potent antioxidant activity, with ethanolic extracts reaching up to 86.91% scavenging efficiency in DPPH assays (Devi et al., 2015). These antioxidant properties contribute to its antimutagenic effects, protecting against DNA damage induced by mutagens like cyclophosphamide (Devi et al., 2015; Mukunthan et al., 2017, Karmakar et al., 2013, Hadem et al., 2014). Furthermore, in liver cancer models induced by diethylnitrosamine (DEN), *C. caesia* extracts lowered serum cancer markers (AST, ALT, ALP, and AChE) while boosting endogenous antioxidant enzymes like SOD and Catalase (Hadem et al., 2014).

Scientific evidence confirms that *Curcuma caesia* possesses significant anticancer potential driven by its rich phenolic content and specific terpenoids. It induces apoptosis through mitochondrial-mediated pathways and causes cell cycle arrest at the G2/M phase (Mukunthan et al., 2017).

These findings support its transition from traditional remedy to a source for targeted phytotherapy in modern oncology.

Protective Effects of *Curcuma caesia* Roxb against Cyclophosphamide Toxicity

Cyclophosphamide (CP) is an anti-cancer drug, its metabolic breakdown induces liver toxicity, oxidative stress and damage to healthy tissues (Devi & Mazumder, 2016). Research indicates that methanolic extracts of *Curcuma caesia* Roxb. (MECC), commonly known as black turmeric, possess potent antioxidant properties that can mitigate these adverse effects (Devi 2016).

Anti-Mycobacterial Activity and Agricultural Innovation

Tuberculosis (TB) remains a significant global health challenge caused by *Mycobacterium tuberculosis* (Mtb) (Gupta, 2018). In Madhya Pradesh, India, traditional knowledge from tribal healers has identified 35 plant species used to treat TB-related symptoms (Sieniawska et al., 2020). Scientific validation revealed that 11 of these species, including *Alstonia scholaris*, *Glycyrrhiza glabra*, and *Curcuma caesia*, possess significant anti-mycobacterial activity against both standard (H37Rv) and multidrug-resistant (MDR) strains (Gupta, 2018). Cytotoxicity assays confirmed that these extracts are safe for human macrophages at effective inhibitory concentrations (Logesh et al., 2020).

Beyond direct medicinal use, plants like *Curcuma caesia* serve as hosts for beneficial endophytic bacteria such as *Paenibacillus* sp. CCB36 (Panichikkal, 2021). Research into agricultural productivity shows that supplementing these bacteria with chitosan nanoparticles (CNPs) can modulate biofilm formation and enhance biocontrol efficacy against pathogens like *Rhizoctonia solani* (Panichikkal, 2021). While zinc oxide nanoparticles (ZnONPs) tend to reduce biofilm development, CNPs improve the colonization and functional persistence of endophytes in soil environments (Mburu et al., 2021; Dev et al., 2018; Garg et al., 2016). The integration of traditional tribal knowledge with modern ethnopharmacology and nanotechnology offers a dual pathway for progress: identifying novel treatments for drug-resistant TB and developing sustainable "green" solutions for agricultural productivity. These studies provide the first documented scientific evidence for the anti-Mtb potential of specific Indian flora and demonstrate the power of nanoparticles in optimizing microbial biological controls (Dev et al., 2018; Garg et al., 2016).

Conclusion

The plant has distinctive bluish-black rhizome, rich aromatic profile, and the presence of bioactive constituents such as curcumin, camphor, ar-turmerone, germacrone, and various essential oils. *In vitro* and *in vivo* studies demonstrated

significant antioxidant, cytotoxic, antidiabetic, antiepileptic, adaptogenic, and anti-ulcer activities, suggesting promising applications in modern therapeutics. Due to high medicinal value, *Curcuma caesia* remains underutilized and is categorized as a threatened species in several regions due to overharvesting and limited cultivation practices.

References

- Ain Ibrahim NN, Kamal N, Mediani A, Sajak AAB, Lee SY, Shaari K, Rahman HA. 2023. (1)H NMR-based metabolomics approach revealing metabolite variation of black Turmeric (*Curcuma caesia*) extracts and correlation with its antioxidant and α -glucosidase inhibitory activities. *Food Technology and Biotechnology* 61(1):107-117. doi: 10.17113/ftb.61.01.23.7711.
- Al-Amin M, Eltayeb NM, Khairuddean M, Salhimi SM. 2021. Bioactive chemical constituents from *Curcuma caesia* Roxb. rhizomes and inhibitory effect of curcuzederone on the migration of triple-negative breast cancer cell line MDA-MB-231. *Natural Product Research*. 35(18):3166-3170. doi: 10.1080/14786419.2019.1690489.
- Arulmozhi DK, Sridhar N, Veeranjaneyulu A, Arora SK. 2006. Preliminary mechanistic studies on the smooth muscle relaxant effect of hydroalcoholic extract of *Curcuma caesia*. *J Herb Pharmacother*. 6(3-4):117-24. doi: 10.1080/j157v06n03_06.
- Baghel SS, Baghel RS, Sharma K, Sikarwar I. 2013. Pharmacological activities of *Curcuma caesia* Roxb.: A review of traditional uses and phytochemistry. *International Journal of Green Pharmacy*, 7(1):1-5.
- Banerjee A, Nigam SS. 1976. Antifungal activity of the essential oil of *Curcuma caesia* Roxb. *Indian Journal of Medical Research* 64(9):1318-21.
- Bhattacharya P, Patel TN. 2021. A study of deregulated MMR pathways and anticancer potential of curcuma derivatives using computational approach. *Scientific Report* 11(1):10110. doi: 10.1038/s41598-021-89282-5.
- Borah S, Sarkar P, Sharma HK. 2021. Analysing *Curcuma caesia* fractions and essential oil for neuroprotective potential against anxiety, depression, and amnesia. *Biotechnology*. 11(5):240. doi: 10.1007/s13205-021-02793-w.
- Borah S, Sarkar P, Sharma HK. 2022. Zederone Improves the Fecal Microbial Profile in Dementia Induced Rat Model: A First Report. *CNS & Neurological Disorders - Drug Targets*. 21(4):335-342. doi: 10.2174/1871527320666210827114227.
- Chaturvedi M, Rani R, Sharma D, Yadav JP. 2020. Effect of

- temperature and pressure on antimycobacterial activity of *Curcuma caesia* extract by supercritical fluid extraction method. *International Journal of Mycobacteriology* 9(3):296-302. doi: 10.4103/ijmy.ijmy_113_20.
- Chaturvedi M, Rani R, Sharma D, Yadav JP. 2021. Comparison of *Curcuma Caesia* extracts for bioactive metabolite composition, antioxidant and antimicrobial potential. *Natural Product Research*. 35(18):3131-3135. doi: 10.1080/14786419.2019.1687472.
- Chirangini P, Sharma GJ, Sinha SK. 2004. Sulfur free radical reactivity with curcumin as reference for evaluating antioxidant properties of medicinal zingiberales. *Journal of Environmental Pathology, Toxicology and Oncology* 23(3):227-36. doi: 10.1615/jenvpathtoxocol.v23.i3.60.
- Dev SK, Choudhury PK, Srivastava R, Sharma M. 2018. Phytochemical characterization and antioxidant assessment of herbal extracts, *Journal of Drug Delivery and Therapeutics* 8(4):126-33.
- Dev SK, Sharma M, Srivastava R, Choudhury PK. 2017. Phytochemical and Pharmacological aspects of *Sarcostemma acidum* (Roxb.) Voigt. *Journal of Pharmacy Research* 11(11):1429-31.
- Devi HP, Mazumder PB, Devi LP. 2015. Antioxidant and antimutagenic activity of *Curcuma caesia* Roxb. rhizome extracts. *Toxicology Reports* 2:423-428. doi: 10.1016/j.toxrep.2014.12.018.
- Devi HP, Mazumder PB. 2016. Methanolic Extract of *Curcuma caesia* Roxb. Prevents the Toxicity Caused by Cyclophosphamide to Bone Marrow Cells, Liver and Kidney of Mice. *Pharmacognosy Research* 8(1):43-9. doi: 10.4103/0974-8490.171106.
- Gangal A, Duseja M, Sethiya NK, Bisht D, Chaudhary SK, Rana VS. 2023. A Validated High-Performance Thin-Layer Chromatography Technique for Routine Analysis of Curcumin in Four Different Species of *Curcuma* Viz. *C. amada*, *C. caesia*, *C. longa* and *C. zedoaria*. *Journal of Chromatographic Science*, doi: 10.1093/chromsci/bmad063.
- Gangal A, Duseja M, Sethiya NK, Bisht D, Chaudhary SK, Rana VS. 2025. A Validated High-Performance Thin-Layer Chromatography Technique for Routine Analysis of Curcumin in Four Different Species of *Curcuma* Viz. *C. amada*, *C. caesia*, *C. longa* and *C. zedoaria*. *Journal of Chromatographic Science* 63(3):doi: 10.1093/chromsci/bmad063.
- Gangal A, Singh P, Yogita, Manori S, Shukla RK. 2025. Nematic lyotropic liquid crystalline ordering in rhizome powder of *Curcuma* species and water mixtures: rheological properties and antioxidant applications. *Soft Matter*. 21(22):4420-4433. doi: 10.1039/d5sm00191a.
- Garg A, Shukla A, Pandey P, Dev SK. 2016. Inhibitory effect of alcoholic extract of *Tulsi* (*Ocimum sanctum*) on calcium oxalate crystals: An study. *Asian Journal of Pharmacy and Pharmacology* 3(2):77-80.
- Grover M, Shah K, Khullar G, Gupta J, Behl T. 2019. Investigation of the utility of *Curcuma caesia* in the treatment of diabetic neuropathy. *Journal of Pharmacy and Pharmacology* 71(5):725-732. doi: 10.1111/jphp.13075.
- Gupta VK, Kaushik A, Chauhan DS, Ahirwar RK, Sharma S, Bisht D. 2018. Anti-mycobacterial activity of some medicinal plants used traditionally by tribes from Madhya Pradesh, India for treating tuberculosis related symptoms. *Journal of Ethnopharmacology* 227:113-120. doi: 10.1016/j.jep.2018.08.031.
- Hadem KL, Sharan RN, Kma L. 2014. Inhibitory potential of methanolic extracts of *Aristolochia tagala* and *Curcuma caesia* on hepatocellular carcinoma induced by diethylnitrosamine in BALB/c mice. *Carcinogenesis* 13:7. doi: 10.4103/1477-3163.133520.
- Hadem KL, Sharan RN, Kma L. 2015. Phytochemicals of *Aristolochia tagala* and *Curcuma caesia* exert anticancer effect by tumor necrosis factor- α -mediated decrease in nuclear factor kappaB binding activity. *Journal of Basic and Clinical Pharmacy* 7(1):1-11. doi: 10.4103/0976-0105.170585.
- Ibrahim NNA, Wan Mustapha WA, Sofian-Seng NS, Lim SJ, Mohd Razali NS, Teh AH, Rahman HA, Mediani A. 2023. A Comprehensive Review with Future Prospects on the Medicinal Properties and Biological Activities of *Curcuma caesia* Roxb. *Evidence-Based Complementary and Alternative Medicine* 2023:7006565. doi: 10.1155/2023/7006565.
- Israr F, Hassan F, Naqvi BS, Azhar I, Jabeen S, Hasan SM. 2012. Report: Studies on antibacterial activity of some traditional medicinal plants used in folk medicine. *Pakistan Journal of Pharmaceutical Sciences* 25(3):669-74.
- Jain A, Jain P, Soni P, Tiwari A, Tiwari SP. 2023. Design and Characterization of Silver Nanoparticles of Different Species of *Curcuma* in the Treatment of Cancer Using Human Colon Cancer Cell Line (HT-29). *Journal of Gastrointestinal Cancer* 54(1):90-95. doi: 10.1007/s12029-021-00788-7.
- Jaiswal D, Agrawal M, Agrawal SB. 2022. Dose differentiation in elevated UV-B manifests variable response of carbon-nitrogen content with changes in secondary metabolites of *Curcuma caesia* Roxb. *Environmental Science and Pollution Research*

- 29(48):72871-72885. doi: 10.1007/s11356-022-20936-1.
- Jaiswal D, Agrawal SB. 2021. Ultraviolet-B induced changes in physiology, phenylpropanoid pathway, and essential oil composition in two *Curcuma* species (*C. caesia* Roxb. and *C. longa* L.). *Ecotoxicology and Environmental Safety* 208:111739. doi: 10.1016/j.ecoenv.2020.111739.
- Jose E, George A, Mathew A, Neethu TR. 2021. Phytochemicals, antioxidants and antimicrobial components in leaf extracts of *Curcuma caesia* Roxb. *Journal of Phytochemistry & Biochemistry*, 3(1): 114.
- Karmakar I, Dolai N, Suresh Kumar RB, Kar B, Roy SN, Haldar PK. 2013. Antitumor activity and antioxidant property of *Curcuma caesia* against Ehrlich's ascites carcinoma bearing mice. *Pharmaceutical Biology* 51(6):753-9. doi: 10.3109/13880209.2013.764538.
- Lal M, Munda S, Begum T, Gupta T, Paw M, Chanda SK, Lekhak H. 2022. Identification and Registration for High-Yielding Strain through ST and MLT of *Curcuma caesia* Roxb. (Jor Lab KH-2): A High-Value Medicinal Plant. *Genes (Basel)*. 13(10):1807. doi: 10.3390/genes13101807
- Lalitha CR, Raveendran K, Salim N. 1995. Effect of *curcuma caesia* leaves on rice seed germination and seedling establishment. *Ancient Science of Life* 15(2):115-8.
- Lamo JM, Rao SR. 2017. Meiotic behaviour and its implication on species inter-relationship in the genus *Curcuma* (Linnaeus, 1753) (Zingiberaceae). *Comparative Cytogenetics* 11(4):691-702. doi: 10.3897/CompCytogen.v11i4.14726.
- Lawand R, Gandhi SV. 2013. Comparison of *Curcuma caesia* Roxb. with other *Curcuma* species by HPTLC. *Journal of Pharmacognosy and Phytochemistry*, 2(4):126-131.
- Leela NK, Adheeba PK. 2024. *Curcuma caesia* Roxb. — update of phytochemicals and pharmacological properties. *Journal of Spices and Aromatic Crops*, 33(1):1-10. <https://doi.org/10.25081/josac.2024.v33.i1.8477>
- Lenka J, Sahoo BC, Kar B, Sahoo S. 2025. Unlocking terpenoid treasures of rhizome and leaf volatiles of *Curcuma caesia* Roxb. through GC×GC-TOFMS analysis. *Scientific Reports*, 15: 10895.
- Liu Y, Roy SS, Nebie RH, Zhang Y, Nair MG. 2013. Functional food quality of *Curcuma caesia*, *Curcuma zedoaria* and *Curcuma aeruginosa* endemic to Northeastern India. *Plant Foods Hum Nutr.* 68(1):72-7. doi: 10.1007/s11130-013-0333-5.
- Mahanta BP, Sut D, Kemprai P, Paw M, Lal M, Haldar S. 2020. A (1) H-NMR spectroscopic method for the analysis of thermolabile chemical markers from the essential oil of black turmeric (*Curcuma caesia*) rhizome: application in post-harvest analysis. *Phytochemical Analysis* 31(1):28-36. doi: 10.1002/pca.2863.
- Majeed A, Majeed M, Thajuddin N, Arumugam S, Ali F, Beede K, Adams SJ, Gnanamani M. 2019. Bioconversion of curcumin into calebin-A by the endophytic fungus *Ovatospora brasiliensis* EPE-10 MTCC 25236 associated with *Curcuma caesia*. *AMB Express*. 9(1):79. doi: 10.1186/s13568-019-0802-9.
- Mohanty S, Panda MK, Acharya L, Nayak S. 2014. Genetic diversity and gene differentiation among ten species of Zingiberaceae from Eastern India. *Biotechnology*. 4(4):383-390. doi: 10.1007/s13205-013-0166-9.
- Mukunthan KS, Satyan RS, Patel TN. 2017. Pharmacological evaluation of phytochemicals from South Indian Black Turmeric (*Curcuma caesia* Roxb.) to target cancer apoptosis. *Journal of Ethnopharmacology* 209:82-90. doi: 10.1016/j.jep.2017.07.021.
- Pakkirisamy M, Kalakandan SK, Ravichandran K. 2017. Phytochemical screening, GC-MS and FT-IR analysis of methanolic extract of *Curcuma caesia* Roxb (black turmeric). *Pharmacognosy Journal*, 9(6):952-956. <https://doi.org/10.5530/pj.2017.6.149>
- Paliwal P, Pancholi SS, Patel RK. 2011. Pharmacognostic parameters for evaluation of the rhizomes of *Curcuma caesia*. *Journal of Advanced Pharmaceutical Technology & Research* 2(1):56-61. doi: 10.4103/2231-4040.79811.
- Pandey S, Singh D, Yadav JP, Verma A. 2025. Medicinal potential of *Curcuma caesia* Roxb.: Phytochemical composition and TLC profile of hydroalcoholic rhizome extract. *Journal of Drug Delivery and Therapeutics*, 15(6): 41-47. <https://doi.org/10.22270/jddt.v15i6.7181>
- Panichikkal J, Jose A, Sreekumaran S, Ashokan AK, Baby CS, Krishnankutty RE. 2022. Biofilm and Biocontrol Modulation of *Paenibacillus* sp. CCB36 by Supplementation with Zinc Oxide Nanoparticles and Chitosan Nanoparticles. *Applied Biochemistry and Biotechnology* 194(4):1606-1620. doi: 10.1007/s12010-021-03710-w.
- Paw M, Gogoi R, Sarma N, Pandey SK, Borah A, Begum T, Lal M. 2020. Study of anti-oxidant, anti-inflammatory, genotoxicity, and antimicrobial activities and analysis of different constituents found in rhizome essential oil of *Curcuma caesia* Roxb. Collected from North East India. *Curr Pharm Biotechnol.* 21(5):403-413. doi: 10.2174/1389201020666191118121609.
- Reenu J, Azeez S, Bhageerathy C. 2015. In vitro Antioxidant Potential in Sequential Extracts of *Curcuma caesia* Roxb. Rhizomes. *Indian Journal of Pharmaceutical Sciences* 77(1):41-8. doi: 10.4103/0250-474x.151596.

- Saraf S, Chhabra SK, Kaur CD, Saraf S. 2012. Development of photochemoprotective herbs containing cosmetic formulations for improving skin properties. *Journal of Cosmetic Science* 63(2):119-31.
- Shivalingu BR, Vivek HK, Nafeesa Z, Priya BS, Swamy SN. 2015. Comparative analysis of procoagulant and fibrinogenolytic activity of crude protease fractions of turmeric species. *Journal of Ethnopharmacology* 172:261-4. doi: 10.1016/j.jep.2015.06.018.
- Shivalingu BR, Vivek HK, Priya BS, Soujanya KN, Swamy SN. 2016. Purification and characterization of novel fibrin(ogen)olytic protease from *Curcuma aromatica* Salisb.: Role in hemostasis. *Phytomedicine*. 23(13):1691-1698. doi: 10.1016/j.phymed.2016.09.007.
- Srivastava R Choudhury PK, Dev SK, Rathore V. 2022. Formulation and Evaluation of α -Pinene Loaded Self-emulsifying Nanoformulation for In-Vivo Anti-Parkinson's Activity, *Recent Patents on Nanotechnology* 16(2), 139-159.