



A REVIEW ON PHYTOCHEMISTRY AND PHARMACOLOGICAL IMPORTANCE OF *CORDIA SEBESTENA*

K Ratna Jyothi*, Kasa Sravya, Yaramala Laxmi Prasanna,
Rathlavath Venkatesh, Rama Rao Tadikonda

CMR College of Pharmacy, Kandlakoya, Medchal, Hyderabad, Telangana, India.

*Corresponding author E- mail: kratnajyothi1994@gmail.com

ARTICLE INFO

Key words:

Cordia sebestena,
phytochemistry,
Geiger tree,
monoterpenoids,
larvicidal activity.

Access this article online

Website:

<https://www.jgtps.com>

Quick Response Code:



ABSTRACT

Over the previous decades, plants possessing biologically active metabolites, which have been proven to be powerful natural medicines. *Cordia sebestena* is a topical blooming plant native to the Caribbean, Florida and central America. It is usually referred to as the Geiger tree or Scarlet *Cordia*. It belongs to the family *Boraginaceae* and it is also known as manjack, and gesho. It is commonly known as Geiger tree. This plant is valued for its own medicinal properties and those are used in traditional medicine for various ailments. It is renowned for its vibrant orange-red flowers, which bloom year-round. *Cordia sebestena* is known for its tolerance to the drought conditions and it is popular in arid regions for adding colours and beauty to the gardens and landscapes. The Geiger-tree aesthetic appeal and adaptability make it a valuable addition to urban and coastal gardens. It is cultivated in tropical and subtropical areas for its ornamental value. It has a low risk of being introduced into temperate area as it does not survive frosts. Its striking flowers, edible fruits, and robust wood have led to its use in reforestation, landscaping, traditional medicine, and sustainable development initiatives. With its low maintenance requirements and wide range of uses, *Cordia sebestena* is a valuable resource for environmental conservation, eco-friendly practices, and community development. The present review describes the habitat, phytochemistry and medicinal uses of *Cordia sebestena*. Thus, this overview draws attention to further biological study on *C. Sebestena* to discover its medicinal activities.

INTRODUCTION:

Throughout the human age, plants have been the major source of basic needs for making food, housing, clothing and medicines. The use of plants as a source of medicine in the form of various folklore medicines, decoctions, oils and many other remedies has been described very well in the history. The plants have been continuously providing new remedies and lead to the human race with unique structural diversity. Medicinal plants are very valuable in

Medicine. Many human disorders can be treated using *Cordia sebestena*. Plants have been the main source of essential materials for food, clothing, and medicine from the beginning of time. Medicinal plants are very valuable in medicine. One of the medicinal plants is *Cordia sebestena*. This plant belongs to the family *Boraginaceae*. *C. sebestena* is an ornamental plant, thoroughly cultivated for gorgeous garden [1-3].

Plant profile

Local names: Scarlet Cordia, Orange Geiger Tree, Geiger Tree, Sebesten Plum.

Vernacular names: Telugu - Virigi; Hindi – Bohari; Lal lasora; Bengali - Kamla buhal raktarag; Kannada - Challekendala; Tamil - Accinayuvili.

Table 1: Taxonomy

Kingdom	Plantea
Subkingdom	Viridiplantae
Division	Tracheophyta
Subdivision	Spermatophyte
Class	Magnoliopsida
Order	Boraginales
Family	Boraginaceae
Genus	<i>Cordia</i>
Species	<i>Sebestena</i>

Cultivation: *Cordia Sebestena* is a slow-growing plant and sheds enough leaves and fruit to require some upkeep. This cultivar has a moderate growth rate and has a high tolerance of salt, drought and nutrient poor soils. However, it is sensitive to soil compaction. It is a handy seashore topic that is planted in parking lots and road medians [4-6].

Botanical description: The plants from the genus *Cordia* are broadly scattered in many countries such as Mexico, Brazil, Argentina, Carib-bean Islands, Panama, Salvador, Bolivia, Paraguay, Texas, California, West Indies, Peru, Ecuador, Malaysia, Curacao, Aruba, Colombia, Venezuela, Costa Rica, India, Suriname, Guyana, Myanmar, Thailand, Somalia, Kenya, Pakistan, tropical Africa, Ceylon, Malacca, Java, China and Japan. The systematic pharmacological studies on the genus *Cordia* have given tremendous acceptance to their traditional uses in health problems. The phytochemical studies of the genus *Cordia* revealed isolation of a total of 293 chemical constituents, out of which 173 new compounds were identified and their structure was mentioned in the review. The

newly isolated compounds belong to the class of quinones, triterpenoids, flavonoids, lignans, neolignans, alkaloids, saponins, monoterpenoids, diterpenoids, steroids, phenyl-propanoids, phenolic, polyphenols, porphyrins and coumarins. Most of the pharmacological studies have been performed using crude extract and fractions, while only 37 newly isolated compounds were evaluated for their pharmacological effects. Furthermore, quinones, terpenoids and flavonoids were found to be the major chemical constituents of the genus *Cordia*, and these constituents are known for their biological effectiveness. Thus, these constituents should be explored for their different pharmaco-logical activities. It is a dense, rounded, evergreen native tree grows up to 30 feet tall and 25 feet wide. It can also grow a trunk that is 1 inch thick. The tree grows slowly. The big, stiff, dark green leaves are 4-9 to inches long, hairy and have a rough, sand paper- like sensation to them. Blooming all year round, but particularly in spring and summer, are 2-inch-wide, dark orange blooms that occur in clusters at the terminals of the branches. These flowers are 1-2 inches long and bear egg- shaped fruits that smell nice but aren't very appetizing. It provides new remedies and lead to the human race with unique structural diversity. The genus *Cordia* is characterised by its alternate petiolate leaves with a dentate or entire margin; white, yellow, or orange flowers with a cyme, spike, or head in floral arrangements; a tubular or campanulate calyx with three to five short teeth; an infundibuliform, hypocateriform, or campanulate corolla with four to eight lobes; stamens included or exerted with pubescent or glabrous filaments at the base; normally, the ovary contains four locules, each containing one erect ovule; fruits are ovoid, globoid, or ellipsoid in shape with a bony endocarp and viscid pulp [7-12].

Propagation: Seed - very slow to germinate, the process can be sped up if the seed is

scarified by lightly abrading the seedcoat to allow easier ingress of water.



Figure 1: *Cordia sebestena* L.



Figure 2: Leaf of *Cordia sebestena*

Flower

Colour: orange red, Features: Very colourful, funnel-shaped; appears in cluster at the branches

Flowering: summer is the season with the most flowers



Figure 3: Flower of *Cordia sebestena*

- Shape: oval or egg-shaped
- Measurement: 1 to 2 inches
- Covering: arid
- Colour: turns from green to white when ripen



Figure 4: Fruits of *Cordia sebestena*



Figure 5: Trunk of *Cordia sebestena*

Uses: *Cordia sebestena* fruit is emollient. It is employed in the diagnosis of different types of fever. Emollient leaves are used in the treatment of bronchitis, coughing, fever and influenza. Ripe white fruit of *C. sebestena* is edible and used in the treatment of indigestion, gastro intestinal problems and dyspepsia. Unripe fruit is poisonous. *Cordia sebestena* (Geiger tree) is helpful in following conditions like indigestion, gastro intestinal disorders, kidney pains, renal dysfunction, cough and bronchial ailments. Leaves used as emollient and in the

treatment of bronchitis, cough, fever and influenza [13,14].

Phytochemistry: *Cordia sebestena* whole plant extracts revealed the presence of alkaloids, sterols, tannins, proteins, amino acids, flavonoids, terpenoids, saponins, carbohydrates, total fatty acid (71.1%). Total 19 compounds were identified as major compounds being 9-octadecene (E) (20.29%); 5-Octadecene (E); 9-eicosene (13.99%). Cyclopropane, nonyl (12.42%); 3-eicosene (E) (72.29%); phenol, 2,4- bis(1,1-dimethyl ethyl) (4.71%); 1-nonadecene (3.17%); 7,9-di-tert-butyl-1oxaspiro (4,5) deca-6,9-diene-2,8-dione (2.70%); and 2,6-diisopropyl naphthalene (2.17%) [6,15,16].

Pharmacological importance [8,9,17-22]

Anti-diabetic activity: The levels of biochemical parameters, hematological indices, serum electrolytes level are improved by the ethanolic extract of fruit in streptozotocin (STZ) which is induced by the diabetogenic rodents and consequences are also evaluated. Blood glucose measurements were used to assess anti-diabetic activity in the chronic biological model by using STZ (65 mg/kg/i.p.)- induced diabetes in rodents. Further studies reveal the effect on body weight, aspartate aminotransferases, alanine, total bilirubin, and total protein, transformations in serum electrolytes such as Na⁺, K⁺, Cl⁻ and Ca⁺² along with estimation of hematological indices such as RBC, WBC, hemoglobin, lymphocytes, neutrophils, eosinophils, and monocytes. The results demonstrated the anti- diabetic potential of plant extract in STZ induced diabetes in rodents, and its affiliated complexities inclusive of anemia, diabetic nephropathy, retinopathy, neuropathy, and hepatitis.

Anti-inflammatory activity: The study involved to evaluate the analgesic and anti-inflammatory properties of ethanolic extract of plant leaves in Wistar albino rats. Eddy's hot plate method is used to determine the analgesic activity. The anti- inflammatory

activity was determined by using Carrageenan- induced paw oedema. The dose-dependent extract was exhibited significant anti-inflammatory effects in the carrageenan-induced inflammation test. Further scientific studies and scientific investigation is required to establish its analgesic and anti-inflammatory properties in other experimental models and clinical settings.

Hepatoprotective activity: To evaluate the evaluate the hepatoprotective effect of the ethanolic extract of *Cordia sebestena* fruit extract (CSFE) against simvastatin- induced hepatotoxicity was induce by simvastatin in rodents. Hepatoprotective potential of CSFE was determined at the 200 to 400 mg/kg body weight by estimating the altered levels of biological and chemical parameters like serum glutamic pyruvic transaminase(SGPT), serum glutamic oxaloacetic transaminase (SGPT), cholesterol, bilirubin, albumin, total protein and hematological indices including red blood cells (RBC), white blood cells (WBC), hemoglobin (Hb), platelets and lymphocytes along with impact on body and liver weight of treated rats. These fruit extract at dose 400 mg/kg reversed liver deteriorations induced by simvastatin in rats, therefore manifesting its traditional use as hepatoprotector.

Anti-microbial activity: The anti-mycobacterial activities of these plants were investigated in *Mycobacterium fortuitum*. The agar cup diffusion method was used for the screening at concentration of 10, 20, 100 and 200 mg/ml. Agar dilution method is used to evaluate the minimum inhibitory concentration.

Larvicidal activity: In this study, an attempt has been made to explore one such plant *Cordea sebestena* leaves. Leaves for its larvicidal properties. In the developing countries like India, one of the biggest threats for public is the tiny creature- mosquitoes. These mosquitoes can be controlled through mosquito repellent, which

cause mortality and kill them. Plants serve as a rich source for potential insecticide. The leaves of the plant were collected and extracted using solvents of increasing polarity such as petroleum ether (60- 80), chloroform, ethyl acetate and methanol. The prepared extracts were evaluated for its larvicidal properties at 100, 250, 500, 1000, 2000 µg/ml against third or fourth instar larvae of *Aedes aegypti*. The evolution clearly shows all the extracts exhibit poor larvicidal property. However, the methanolic extract was found to be little effective as compared to the other extracts but not up to a significant level.

Anti-tumor activity: Since antiquated times, plants act as a treasure of effective drugs for cancer therapy. The total content of phytoconstituents such as phenolic, flavonoid, tannins, and nutrient content like carbohydrates, protein are notably observed acetone extract. The natural proficiency of extricates was too assessed by anti-bacterial moment against chosen human pathogens. Confined hesperetin compound altogether uncovered cytotoxicity for HeLa cell line and its anticancer capacity was revalidated by in silico atomic docking study, which shown solid interaction with E6 protein HPV16.

Anti-helminthic activity: Anti-helminthic properties of *Cordia sebestena* leaves extracts, petroleum ether, chloroform, ethyl acetate and methanol were evaluated against Indian earthworms *Pheretima posthuma* at 10mg/ml, 20mg/ml, 30mg/ml, 40mg/ml and 50mg/ml, piperazine citrate is used as standard. This study reveals that *Cordia sebestena* chloroform, and methanolic extract possess anti-helminthic properties. However, the effect of these extracts was found to be less significant as compared to the standard drug piperazine citrate.

Anti-oxidant activity: The chemical composition of the essential oil from the bark of the *Cordia sebestena* obtained by hydrodistillation was determined using gas chromatography –mass spectroscopy and

analysed for its free radical scavenging potential, thereby preventing oxidative stress and damage to cells. This is typically analysed assessed using assays such as DPPH (2,2-diphenyl-1-picrylhydrazyl). The hydrocarbons may well be valuable for chemotaxonomic characterization of *Cordia sebestena*. The total phenolic content was determined using spectrophotometric methods with a Folin-Ciocalteu reagent. The anti-oxidant activity of the extracts was determined by its ability to inhibit DPPH radicals through IC₅₀ values (ppm). The highest total phenolic content (167.61 ± 0.56 mg GAE/g) and best anti-oxidant activity (31.41 ppm) were found in 70% ethanol extract of *C. sebestena* compared to other extracts (ethyl acetate > dichloromethane > n-hexane).

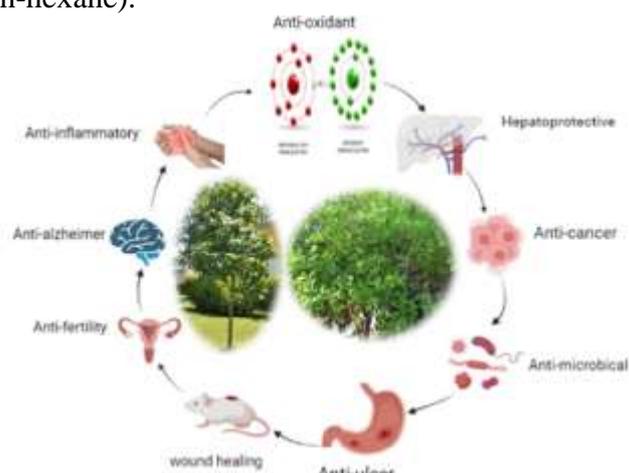


Figure 6: Common pharmacological activities of *Cordia sebestena*

CONCLUSION

The conventional medical system has grown in significance within the pharmaceutical industry from ancient times. A vast majority of people in the majority of developing nations rely on traditional healers who are fervent about healthy herbs to meet their basic medical needs. For historical and cultural reasons, seasoning remedies have maintained their reputation despite the availability of fashionable medications. Due to the increased use of such flavouring medications, concerns and remarks about

their efficacy, safety, and quality have arisen in both developed and developing nations. Scientists are being obliged to objectively evaluate a variety of old claims due to growing curiosity. Because everyone seems to be interested in using traditional medicine in this modern age, it is necessary to verify the conventional claims. Therefore, before using any recommended healthy plants as a medicine, both regular users and medical experts currently get updated, varied information regarding the safety and efficacy of these plants. *Cordia sebestena* has a number of pharmacological properties, including those that are anti-microbial, anti-diabetic, insulin sensitizing, anti-microbial, hepatoprotective, anti-inflammatory, larvicidal, anti-helminthic, anti-oxidant, anti-tumor and that also have some psychiatric and neurological conditions. Thus, the present study also helps to check the falsification of this important medicinal plant and to be significant and encouraging towards the goal for standardization.

Conflicts of interest

None

REFERENCES

1. Thirupathi K et al. A review of medicinal plants of the genus *Cordia*: their chemistry and pharmacological uses. J Nat Remedies 2008; 8:1-10.
2. Manisha J Oza, Yogesh A Kulkarni. Traditional uses, phytochemistry and pharmacology of the medicinal species of the genus *Cordia* (Boraginaceae). Royal Pharmaceutical Society, Journal of Pharmacy and Pharmacology. 2017; 69:755-789.
3. Kompelly A, Kompelly S, Vasudha B, Narender B. *Rosmarinus officinalis* L.: an update review of its phytochemistry and biological activity. Journal of Drug Delivery and Therapeutics. 2019; 9(1):323-330.
4. Nisha M, Chandru N, Pradeep P, Selvarasu P, Surendra Kumar M, Latha ST, Astalakshmi N. A Review on Phytochemical and Pharmacological Activity of *Cordia sebestena*. Int. J. Pharm. Sci. Rev. Res. 2022; 73(1):156-161.
5. Shanmugam Prakash, Selvaraj Kanaga, Nagaraj Elavarasan, Kasivisvanathan Subhashini, Ramamurthy Dhandapani, Magudeewaran Sivandam, Poomani Kumardhas, Chinnaswamy Thirunavukkarasu, Venugopal Sujatha. Isolation of hesperetin- A flavonoid from *Cordia sebestena* flower extract through antioxidant assay guided method and its antibacterial, anticancer effect on cervical cancer via *in-vitro* and *in silico* molecular docking studies. Journal of Molecular Structure. 2020; 1207:127751.
6. Foluso O Agunbiade, Charles B Adeosun, Gbenga G Daramola. Nutritional properties and potential values of *Cordia sebestena* seed and seed oil (GIDA) 2013; 38(3):127-133.
7. CB Adeosun, OS Sojini. The chemical composition of flower lipids of *Cordia sebestena*. Adv Environ Biol. 2012; 6:655-657.
8. CB Adeosun, A Adewuyi. Identification of the chemical constituents of the petroleum ether extract from the flower *Cordia sebestena*. Int J Adv Sci Technol. 2012; 6:3-6.
9. Ahmed Bokhari SW, Sharif H, Umer Gilani SM, Ali ST, Ahmed S, Ahmed Siddiqui MU, Mohtasheemul Hasan M. Pharmacognostic and phytochemical study of the flowers of *Cordia sebestena* L. Pak J Pharm Sci. 2022; 35(1):69-76.

10. Adeosun CB, Olaseinde S, Opeifa AO, Atolani O. Essential oil from the stem bark of *Cordia sebestena* scavengers free radicals. J. of Acute Medicine. 2013; 3:138-141.
11. Adhithya Ragunahan, Lokesh Ravi, Kannabiran Krishnan. Anticancer Cytotoxicity Activity of Pentane-2,4-dione extracted from the leaves of *Cordia sebestena*. Research Journal of Pharmacy and Technology. 2018; 11(6):2191-2196.
12. Kumarseran. M, Palanisamy PN, Kumar PE. Application of eco-friendly natural dye obtained from *Cordia sebestena* on cotton using combination of mordants. J Nat Prod Plant Resour. 2012; 2:32-38.
13. Rao YR, Jena KS, Sahoo D, Rout PK, Ali S. Safety evaluation of ambrette seed oil. J Am Oil Chem Soc. 2005; 82:749-752.
14. Hikmawanti NPE, Hanani E, Sapitri Y, Ningrum W. Total Phenolic Content and Antioxidant Activity of Different Extracts of *Cordia sebestena* L. Leaves. Pharmacogn J. 2020; 12(6):1311-1316.
15. Osho A, Otuechere CA, Adeosun CB, Oluwagbemi T, Atolani O. Phytochemical, sub-acute toxicity, and antibacterial evaluation of *Cordia sebestena* leaf extracts. J Basic Clin Physiol Pharmacol. 2016 Mar;27(2):163-70.
16. Atolani O, Kayode OO, Adeniyi O and Adeosun CB. *In vitro* antioxidant potential of fatty acids obtained by direct transmethylation from fresh *Cordia sebestena* Flowers. Annals of Topical Research. 2014; 36(2):104114.
17. Chaudhary S, Verma HC, Gupta MK, Kumar H, Swain SR, Gupta RK, El-Shorbagi AN. Anti diabetic aptitude of *Cordia sebestena* and its outcome on biological parameters, serum electrolytes, and hematological markers. Pharmacognosy. 2019; 11: 418-423.
18. Meshack S, Gupta S. A Review of Plants with Remarkable Hepatoprotective Activity. Journal of Drug Delivery and Therapeutics. 2022; 12(1):194-202.
19. Chandana R, Basini J, Reddy VJS. Hepatoprotective and antioxidant effect of *Cordia sebestena* in animal model. International Journal of Pharmacology & Toxicology. 2014; (3):181-189.
20. Sarathchandiran I, Gnanavel M. Investigation on hypoglycemic, antioxidant and hypolipidemic activity of ethanolic leaf extract of *Cordia sebestena* in Streptozotocin – induced diabetic rats. International Journal of Research in Pharmaceutical Sciences. 2013; 4(3):336-343.
21. Trivedi MH, Ramana KV, Rao CV. Evaluation of antiinflammatory and analgesic activities of *Cordia sebestena* L. roots. Indo American Journal of Pharm Research. 2015; 5(08):2765-2768.
22. Dai J, Sorribs A, Yoshida WY, Williams PG. Sebestenoids A-D, BACE1 inhibitors from *Cordia sebestena*. J of Phytochemistry. 2013; 71(17-18):2168-2173.