



Exploring the Cardioprotective and Multifaceted Health Benefits of Terminalia Arjuna

*Sweeya CH, Laxmi Priyanka K, Hemalatha S, Kavitha P, Muni Sireesha S**

Department of Pharmaceutical Chemistry, Sarojini Naidu Vanita Pharmacy Maha Vidyalaya, Osmania University, Tamaka, Hyderabad, India, 500017

Email: sirsnvpmv@gmail.com

DOI : <https://doi.org/10.55248/gengpi.6.0125.0629>

ABSTRACT:

Traditional treatments have been used by various cultures for centuries to treat health problems. Terminalia arjuna, also known as the arjuna tree, is a medicinal plant from the Combretaceae family. It has long been employed to address a variety of health conditions, such as heart diseases, bleeding, diabetes, cough, tumors, leucorrhea, asthma, inflammation, vitiligo, wounds, and anemia. Although all parts of the plant offer medicinal benefits due to the presence of specific bioactive compounds, it is mainly the bark that is commonly used by traditional healers to treat different ailments. The main active ingredients in Terminalia arjuna include glycosides (especially cardiac glycosides), flavonoids (such as arjunone and arjunolone), and triterpenoids. Terminalia arjuna provides various health benefits, including being cardioprotective, anti-inflammatory, anti-asthmatic, anti-cancer, and antimicrobial. This review aims to gather and present information from pharmacological and clinical research to gain a deeper understanding of the therapeutic potential of Arjuna.

KEY WORDS: Arjuna, Terminalia arjuna, Cardiovascular disorders, medicinal plant, herbal remedies.

INTRODUCTION:

Medicinal herbs have been acknowledged as an essential part of traditional healing practices for treating human ailments since ancient times and remain significant today, as natural treatments are perceived as healthier than their synthetic equivalents. Herbal therapies have become increasingly popular due to their benefits, such as easy access, safety, affordability, rare adverse reactions, and cultural preference. These plants contain organic substances that exert physiological effects on the human organism, with bioactive elements like carbohydrates, terpenes, steroids, alkaloids, tannins, flavonoids, and phenolic compounds.

Arjuna, a therapeutic plant, is scientifically known as Terminalia arjuna. It exhibits a variety of healing properties, including antioxidant, blood pressure-lowering, anti-atherosclerotic, anti-inflammatory, anti-cancer, anti-genotoxic, and digestive-enhancing effects. The objective of this review is to compile and refresh the existing information about Terminalia arjuna, focusing on its botanical, pharmacological, chemical composition, and clinical research aspects.

PHYSICAL CHARACTERISTICS AND PROPERTIES:

Taxonomic Rank	Classification
Kingdom	Plantae
Division	Magnoliophyta
Class	Magnoliopsida
Order	Myrtales
Family	Combretaceae
Genus	Terminalia
Species	T. arjuna

HISTORY:

The *Terminalia arjuna* is a deciduous riparian tree native to India, reaching heights of 20 to 27 meters. It thrives in humid, fertile, and red lateritic soils but can adapt to various soil types and grow in shaded areas. The tree features conical, oblong leaves that are green on the top and brown underneath. Its bark is smooth and grey, and it produces pale yellow flowers that bloom between March and June. The fruit is fibrous and woody, characterized by a five-winged structure, measuring about 2 to 5 cm in size, and matures between September and November.

METHOD OF EXTRACTION:

The extraction of *Terminalia arjuna* can be performed through various techniques:

1. SOLVENT FRACTIONATION PROCEDURE:

For the solvent fractionation process, dissolve 2 grams of the aqueous extract in 100 milliliters of water and mix the solution for 30 minutes. Afterward, perform successive extractions using solvents with gradually increasing polarity, such as chloroform, methyl isobutyl ketone (MiBK), ethyl acetate, and n-butanol. Each solvent fraction should be gathered, evaporated, and weighed.

2. AQUEOUS EXTRACTION PROCEDURE:

The aqueous extraction method involves cutting the bark into small pieces, cleaning them thoroughly, drying them in a shaded area, and grinding the dried bark into a fine powder. To prepare the extract, combine 2 grams of this powder with 100 milliliters of distilled water and heat the mixture for 15 to 20 minutes. After boiling, filter the solution first with a tea filter and then with Whatman filter paper. To concentrate the extract, boil it once more, reducing the volume to 5 milliliters, and store it in the refrigerator at 4°C.

3. LIPID NANOPARTICLE EMULSION PROCEDURE:

The creation of a lipid nanoparticle emulsion begins by dissolving the dry bark extract powder in 1,2-pentanediol to form an alcoholic solution. This solution is then blended with an

emulsifier and glycerol monostearate. Heat the mixture to a range of 75 to 85°C, stirring at 650 revolutions per minute for 10 minutes. Afterward, combine the oil phase with distilled water, heat it to the same temperature range, and continue stirring at the same speed. Finally, the mixture is homogenized using a high-pressure homogenizer and allowed to cool to room temperature.

CHEMICAL CONSTITUENTS:

1. Stem Bark:

- **Triterpenoids:** The stem bark contains a variety of triterpenoids, such as arjunolic acid, arjunic acid, arjunin, arjugenin, and ursane triterpenoids. These compounds are known for their anti-inflammatory, anti-cancer, and cardioprotective properties.
- **Glycosides:** It contains several glycosides, including arjunetin, arjunoside I and II, and arjunaphthanolide, which can have various biological activities such as antioxidant and anti-inflammatory effects.
- **Flavonoids:** Key flavonoids like luteolin, quercetin, and kempferol are present, which are powerful antioxidants that help combat oxidative stress and inflammation.
- **Tannins:** The bark also contains tannins such as castalagin, punicalagin, and ellagic acid. Tannins have antioxidant and astringent properties, contributing to wound healing and anti-inflammatory actions.
- **Minerals:** Important trace minerals such as zinc, copper, calcium, and magnesium are also found, contributing to the plant's overall nutritional value and potential health benefits.

2. Roots:

- **Triterpenoids:** Similar to the stem bark, the roots contain arjunolic acid and oleanolic acid, both of which are known for their anti-inflammatory, hepatoprotective, and anti-cancer properties.
- **Glycosides:** The roots contain specific glycosides, including arjunoside I, II, III, and IV, as well as a compound called 2,19-dihydroxy-3-oxo-olean-12-en-28-oic acid 28-O-d-glucopyranoside. These glycosides contribute to the plant's cardiovascular and anti-inflammatory effects.

3. Leaves and Fruits:

- **Glycosides:** Chebuloside II and bellericoside are present in the leaves and fruits, adding to the plant's anti-inflammatory and antioxidative potential.
- **Flavonoids:** The leaves and fruits also contain flavonoids such as luteolin, which have anti-inflammatory, antioxidant, and anticancer effects.

- **Tannins:** Ellagic acid, gallic acid, corilagin, and chebulagic acid are key tannins in the leaves and fruits, offering benefits like antimicrobial, anti-inflammatory, and antioxidant effects.
- **Minerals:** Like the stem bark, the leaves and fruits also provide essential minerals like calcium, magnesium, zinc, and copper, which are vital for overall health.

Arjuna Terminalia plant



Leaves

Arjuna Terminalia bark



fruits

FORMULATIONS:**PHARMACOLOGICAL ACTIVITIES:****1. Anti-Hypertensive Properties**

- **Pulmonary Hypertension Prevention:** Aqueous extract effectively prevented monocrotaline-induced pulmonary hypertension in Wistar rats, likely due to its antioxidant properties and influence on pulmonary arteriolar wall thickening.

2. Anxiolytic (Anti-Anxiety) Effects

- **Behavioral Studies in Mice:** The alcoholic extract showed significant anti-anxiety activity in various behavioral paradigms.

3. Cardiotoxic Activity

- **Prophylactic and Therapeutic Benefits:** The bark extract demonstrated the ability to regulate autonomic function and prevent left ventricular remodeling in rats, promoting overall heart health.

4. Diuretic Activity

- **Cerebral Vascular Leakage Prevention:** Hydroalcoholic extract reduced acute hypobaric hypoxia-induced vascular leakage in mice by modulating the renin-angiotensin-aldosterone system through ANP signaling.

5. Wound Healing

- **Healing Acceleration:** T. arjuna bark's astringent tannins promoted wound healing in rats, accelerating the recovery process.

6. Antimicrobial Activity

- **Effectiveness Against Bacteria:** Methanol, ethanol, acetone, and aqueous extracts showed antimicrobial effects. Acetone leaf extract was particularly effective against *Staphylococcus aureus*, and organic extracts inhibited Gram-negative bacteria, except *Pseudomonas aeruginosa*.

7. Antioxidant Properties

- **Free Radical Scavenging:** Various extracts, especially the ethanolic extract, exhibited high antioxidant activity, with significant effects on DPPH, superoxide radical scavenging, and lipid peroxidation.
- **Antimutagenic Effects:** The alcoholic extract showed protection against DNA damage, reducing polychromatic and normochromatic erythrocytes in a micronucleus test.

8. Liver Enzyme Inhibition

- **Impact on Drug Metabolism:** Both alcoholic and aqueous extracts inhibited key liver enzymes (CYP3A4, CYP2D6, and CYP2C9), impacting drug metabolism. The inhibition was rapid and reversible in human liver microsomes.

9. Kidney Health and Stone Prevention

- **Cytoprotective and Anti-Apoptotic Effects:** The aqueous extract demonstrated anti-apoptotic effects in MDCK cells and prevented the formation of calcium oxalate crystals, which could help in preventing kidney stones.

10. Anti-Dyslipidemic and Lipid-Lowering Effects

- **Studies on Hamsters:** Ethanol and solvent ether fractions of T. arjuna demonstrated lipid-lowering effects, confirming its anti-dyslipidemic properties.

11. DNA Protection and Free Radical Scavenging

- **Protection Against DNA Damage:** Ethanol extracts and ethyl acetate fractions significantly protected against H₂O₂-induced DNA damage and showed inhibition of various free radicals (DPPH, hydroxyl, ABTS, etc.).
- **Metal Chelation:** The ethyl acetate fraction also exhibited strong metal chelation activity, further supporting its antioxidant and protective effects.

12. Anti-Ulcer Effects

- **Free Radical Scavenging and Cytoprotection:** The methanolic extract of T. arjuna bark demonstrated anti-ulcer properties through its free radical scavenging activity and cytoprotective effects.

13. Anti-Hyperglycemic and Diabetic Effects

- **Blood Sugar Regulation:** The methanolic extract of T. arjuna leaves showed significant anti-hyperglycemic effects in diabetic models, largely attributed to its antioxidant properties.

14. Anti-Cancer Properties

- **Protection Against Cancer:** The aqueous extract exhibited increased antioxidant enzyme activity and effective protection against cancerous growth in mice.

CONCLUSION:

The rising fascination with therapeutic plants has resulted in the identification of novel chemical substances and the examination of the medicinal attributes of Terminalia arjuna (arjuna). Numerous laboratory and clinical trials have confirmed its potency as a heart-protective, free radical-scavenging, and plaque-reducing agent. Nevertheless, several critical gaps remain in the investigations, such as the absence of absorption studies, uniform chemical composition of the extracts, and thorough analysis of their extended toxicity. These research areas are essential for gaining a more comprehensive insight into the plant's complete effectiveness and safety for medical application.

The exact function of arjuna in primary and secondary coronary prevention requires more investigation. Furthermore, research should be done to examine its effects on CYP450 enzymes and how it interacts with medications like aspirin, statins, ACE inhibitors, and β -blockers. Increasing knowledge of its therapeutic potential could help doctors deal with issues relating to the treatment of cardiovascular diseases.

ACKNOWLEDGEMENT:

The authors are thankful to the Management, Principal of Sarojini Naidu Vanita Pharmacy Maha Vidhalaya, Osmania University, Hyderabad, India for providing research facilities.

REFERENCE:

1. Kapoor D, Vijayvergiya R, Dhawan V. Terminalia arjuna in coronary artery disease: ethnopharmacology, pre-clinical, clinical & safety evaluation. *J Ethnopharmacol.* 2014;155:1029-1045.
2. Sharma PC, Yelne MB, Dennis TJ. Database on Medicinal Plants Used in Ayurveda. New Delhi: CCRAS (The Central Council for Research in Ayurvedic Sciences); 2005.
3. Chopra RN, Chopra IC, Handa KL, Kapur LD. Terminalia arjuna W&A (Com-bretaceae). In: Chopra RN, Chopra IC, Handa KL, Kapur LD, eds. *Chopra's Indigenous Drugs of India*. 1st ed. Calcutta, India: UNDhur & Sons; 1958:421-424.
4. Nadkarni AK. *Indian Materia Medica*. 1st ed. Mumbai, India: Popular Prakashan; 1976.
5. Ali M. *Text Book of Pharmacognosy*. 1st ed. New Delhi: CBS Publishers; 1994.
6. Row LR, Murty PS, SubbaRao GSR, Sastry CSP, Rao KVJ. Chemical examination of Terminalia arjuna: Part-XII: isolation and structure determination of Arjunic acid, a new trihydroxytriterpene carboxylic acid from Terminalia arjuna bark. *Indian J Chem.* 1970;8:716-721.
7. Honda T, Murae T, Tsuyuki T, Takahashi T, Sawai M. Arjungenin, arjunglucoside I and arjunglucoside II, a new triterpene and new triterpene glucosides from Terminalia arjuna. *Bull Chem Soc Jpn.* 1976;49:3213-3218.
8. Singh DV, Verma RK, Gupta MM, Kumar S. Quantitative determination of oleanane derivatives in Terminalia arjuna by high performance thin layer chromatography. *Phytochem Anal.* 2002;13:207-210.
9. Singh DV, Verma RK, Singh SC, Gupta MM. RP-LC determination of oleanane derivatives in Terminalia arjuna. *J Pharm Biomed Anal.* 2002;28:447-452.
10. Anjaneyulu ASR, Prasad AVR. Structure of terminic acid, a Dihydroxytriterpene carboxylic acid from Terminalia arjuna. *Phytochemistry.* 1983;22:993-998.
11. Singh B, Singh VP, Pandey VB, Rucker G. A new triterpeneglycoside from Terminalia arjuna. *Planta Med.* 1995;61:576-577.
12. Wang W, Ali Z, Shen Y, Li X, Khan IA. Ursane triterpenoids from the bark of Terminalia arjuna. *Fitoterapia.* 2010;81:480-484.
13. Row LR, Murty PS, Subba Rao GSR, Sastry CSP, Rao KVJ. Chemical Examination of Terminalia arjuna: Part-XII: Isolation and structure determination of arjunetin from Terminalia arjuna bark. *Indian J Chem.* 1970;8:772-775.
14. Honda T, Murae T, Tsuyuki T, Takahashi T. The structure of arjungenin: a new sapogenin from Terminalia arjuna. *Chem Pharm Bull.* 1976;24:178-180.
15. Sharma PN, Shoeb A, Kapil RS, Popli SP. Arjunolone: a new flavones from Stem bark of Terminalia arjuna. *Indian J Chem.* 1982;21B:263-264.
16. Tripathi VK, Pandey VB, Udupa KN, Rucker G. Arjunolitin, a triterpene Glycoside from Terminalia arjuna. *Phytochemistry.* 1992;31:349-351.
17. Ali A, Kaur G, Hayat K, Ali M, Ather M. A novel naphthanolglycoside from Terminalia arjuna with antioxidant and nitricoxide inhibitory activities. *Pharmazie.* 2003;58:932-934.
18. Ali A, Kaur G, Hamid H, et al. Terminoside A, a new triterpeneglycoside from The bark of Terminalia arjuna inhibits nitricoxide production in murine macrophages. *J Asian Nat Prod Res.* 2003;5:137-142.
19. Wang W, Ali Z, Li XC, Shen Y, Khan IA. Triterpenoids from two Terminalia species. *Planta Med.* 2010;76:1751-1754.
20. Patnaik T, Dey RK, Gouda P. Isolation of triterpenoidglycoside from bark of Terminalia arjuna using chromatographic technique and investigation of pharmacological behavior upon muscle tissues. *E-J Chem.* 2007;4: 474-479.
21. Alam MS, Kaur G, Ali A, Hamid H, Ali M, Athar M. Two new bioactive oleanane triterpene glycoside from Terminalia arjuna. *Nat Prod Res.* 2008;22: 1279-1288.
22. Ahmad MU, Mullah KB, Norin T, Ulla JK. Terminic acid, a new trihydroxy-triterpene carboxylic acid from bark of Terminalia arjuna. *Indian J Chem.* 1983;22:738-740.

-
23. Pettit GR, Hoard MS, Doubek DL, et al. Antineoplastic agents 338. The Cancer cell growth inhibitory. Constituents of Terminalia arjuna (Combretaceae). *J Ethnopharmacol.* 1996;53:57-63.
 24. Anonymous. Terminalia arjuna. *Altern Med Rev.* 1999;4:436-437.
 25. Saha A, Pawar VM, Jayaraman S. Characterization of polyphenols in Terminalia arjuna bark extract. *Indian J Pharm Sci.* 2012;74:339-347.
 26. Takahashi SH, Tanaka H, Hano Y, Ito K T, Nomura T, Shigenobu K. Hypotensive effects in rats of hydrophilic extract from Terminalia arjuna containing tannin related compounds. *Phytother Res.* 1997;11:424-427.
 27. Lin TC, Chien SC, Chen HF, Hsu FL. Tannins and related compounds from Combretaceae plants. *Chin Pharm J.* 2001;52:1-26.
 28. Kuo PL, Hsu YL, Lin TC, Lin LT, Chang JK, Lin CC. Casuarinin from the bark of Terminalia arjuna induces apoptosis and cell cycle arrest in human breast adenocarcinoma MCF-7 cells. *Planta Med.* 2005;71:237-243.
 29. Dwivedi S, Udupa N. Terminalia arjuna: pharmacognosy, phytochemistry, pharmacology and clinical use: a review. *Fitoterapia.* 1989;60:413-420.
 30. Anjaneyulu ASR, Prasad AVR. Chemical examination of the roots of Terminalia arjuna characterization of two new triterpenoid glycoside. *Indian J Chem.* 1982;21:530-533.
 31. Anjaneyulu ASR, Prasad AVR. Chemical examination of roots of Terminalia arjuna-the structure of arjunoside III and arjunoside IV, two new triterpenoid glycosides. *Phytochemistry.* 1982;21:2057-2060.
 32. Choubey BK, Srivastava SK. Antifungal agents from Terminalia arjuna. *Indian J Chem.* 2001;40B:354-356.
 33. Upadhyay RK, Pandey MB, Jha RN, Singh VP, Pandey VB. Triterpene Glycoside from Terminalia arjuna. *J Asian Nat Prod Res.* 2001;3:207-212.
 34. Rastogi RP, Mehrotra BN. *Compendium of Indian Medicinal Plants.* vol. 3. New.