



EFFECT OF DIFFERENT DRYING METHODS ON THE QUALITY OF STEM BARK OF *TERMINALIA ARJUNA* ROXB.

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ABSTRACT

The effects of Ayurvedic medicine by different drying conditions were studied. Stem bark of *Terminalia arjuna* was chosen and dried by shade, sun and oven. The dried samples were characterized by means of microscopic study, preliminary physicochemical & phytochemical studies and Thin Layer Chromatography (TLC) study. The aim of this study was therefore to develop an understanding of suitable conditions for the processing of stem bark of *T. arjuna*. The objectives of this study were to investigate the effect of drying techniques i.e. shade, sun and oven drying, on the physicochemical, phytochemical and TLC studies of stem bark of *T. arjuna*. The results of physicochemical parameters i.e. loss on drying, total ash, acid insoluble ash, alcohol soluble extractive and water soluble extractive values of shade dried sample showed lower values when compared with sun dried and oven dried samples. Preliminary phytochemical analysis of all samples revealed the presence of Alkaloids, Glycosides, Flavonoids, Steroids, Triterpenoids, Saponins, Tanins and Carbohydrates. Thin layer chromatogram of the methanolic extract of shade dried sample after derivatisation with anisaldehyde sulphuric acid reagent showed nine major spots, where as sun and oven dried samples showed eight spots.

Keywords: Arjuna, *Terminalia arjuna*, stem bark, microscopic study, preliminary physicochemical study, Preliminary phytochemical analysis, TLC study.

INTRODUCTION

The medicinal plants *Terminalia arjuna* (Roxb, Wight Arn) is a large evergreen tree with butterressed trunk. It belongs to *Combretaceae* family, is an important cardiogenic plant described in the Ayurveda¹. The bark is useful as an anti-ischaemic and cardioprotective agent in hypertension and ischaemic heart disease, especially in disturbed cardiac rhythm, angina or myocardial infarction². *Terminalia arjuna* has been shown to be beneficial for coronary artery disease, heart failure, and possibly for high cholesterol levels³. It has also been found to be antibacterial and antimutagenic⁴. It helps in promoting proper gastrointestinal function to regulate gastrointestinal pH, while improving gastrointestinal motility, increasing stool specific gravity and reducing the populations of certain fecal micro-organisms, including yeast [*Candida albicans*]⁵. This might have significant advantages to some individuals by promoting proper dietary protein digestion and absorption and reducing bowel putrefactive processes in the colon⁶. It also improves the cardiac muscle function and subsequent improved pumping activity of the heart⁷. A decoction of its bark with cane sugar and boiled cow's milk is highly recommended for endocarditis, pericarditis and angina⁸. Chemical constituents of different classes such as hydrolysable tannins⁹, triterpenoid acids and their glycosides^{10,11}, flavonoids¹², Phenolics¹³, phytosterol¹⁴, were reported from stem bark portion of *Terminalia arjuna* species. Additionally, Arjunglucoside I-III, arjunic acid, arjunetin, arjunolic acid, and terminic acid also form group of important constituents of the bark¹⁵. A number of previously published papers report the therapeutic properties for *Terminalia arjuna*¹⁶⁻¹⁹. According to Kumar and Jain, people living in the south Surguja district of Madhya Pradesh uses the bark of *T. arjuna* in the treatment of fever and high blood pressure²⁰.

People living in the Malkangiri district of Orissa, chew the fresh bark of *T. arjuna* and the juice is used as antacid²¹. Decoction of the bark is used as ulcer wash, while bark ash is used in the treatment of the snake bite and scorpion sting²².

The kinetics of the drying process can define the final quality properties of the dried material. Artificial drying has been one of the most important processes in preprocessing of agricultural products, aiming to achieve the phytotherapy product needs of the pharmaceutical industry, which does not have infra-structure to use fresh plants in the quantities required for industrial production²³.

The post-harvesting process of medicinal plants has great importance in the production chain, because of its direct influence on the quality and quantity of the active principles in the product sold²⁴. For this reason, adequate dryers are needed, using temperature, velocity and humidity values for drying air that provides a rapid reduction in the moisture content without affecting the quality of the active principles of medicinal plants. The drying process may also contribute to regular supply and facilitate the marketing of plants, because it facilitates the transport and storage²⁴.

Specific processing methods are often required, to reduce drying time, to detoxify the inherent toxic constituents, to reduce side effects or to enhance therapeutic effects. For example, the methods and temperatures used for drying may have a considerable impact on the quality of the resulting medicinal plant materials. Shade drying is the preferred method for drying plant material since it can maintain or minimize loss of color of leaves and flowers; and the lower temperatures can prevent the loss of volatile substances in the plant materials^{25,26}. Taking into account of this wide use bark of *T. arjuna* present study aimed to

evaluate the effect of different drying methods on the quality.

MATERIAL AND METHODS

Plant material

The stem bark of *Terminalia arjuna* were collected from Raipur in March 2009. A voucher specimen was kept at the Department of Dravyaguna, Govt. Ayurvedic College, Raipur, after identification of the plant. The stem bark was subjected to three different drying conditions namely, shade, sun and microwave oven dry, however for all the three strategies, approximately five hundred grams of the fresh plant material was washed, drained and used. For shade dry, the pre washed and drained plant material was placed on a filter paper (90x60 cm) at room temperature (27± 1°C) for 3 days. For sun dry, the fresh material was placed in to the greenhouse for 3 days. For microwave oven drying, the plant material was placed in the middle of the turntable of a commercial microwave oven (SAMSUNG Model CE1031LFB; 900W) for 4 min. Once, the drying process was over, the dry weights were powdered using a laboratory blender and stored for further work.

Macroscopic study

Macroscopy is the study of the form of an object, where the material is known to occur in a particular form. Macroscopic and organoleptic features viz. color, odour, taste, shape, sizes etc. of the stem bark were observed^{27,28}.

Microscopic study

The shade dried stem barks were powdered and passed through a sieve no 85 to obtain the fine powder and then subjected for microscopic examination. About few mg of the powder was warmed with chloral hydrate solution, stained with saffranin, mounted with glycerin and observed under suitable magnification^{27,28}.

Physicochemical parameters study

The physicochemical parameters viz. loss on drying, total ash, acid-insoluble ash, alcohol-soluble extractive and water-soluble extractive were determined for the stem bark of *T. arjuna*²⁸. The results are presented in Table1.

Preliminary Phytochemical Analysis

The methanolic and aqueous extracts were subjected to preliminary phytochemical analysis. The various quantitative chemical tests performed on the extracts were for alkaloids, glycosides, flavonoids, steroids, triterpenes, tannins, carbohydrates, proteins and amino acids²⁹⁻³¹. The results are presented in Table 2.

Table 1: Physicochemical parameters studies of stem bark of Terminalia arjuna

NO.	Physico-Chemical Test	Shade drying	Sun drying	Oven drying
1.	Loss on drying at 105°C	5.12	6.28	7.32
2.	Total ash	12	18	16
3.	Acid-insoluble ash	1.2	1.4	1.6
4.	Alcohol soluble extractive	12.02	14.63	16.05
5.	Water soluble extractive	17.45	21.30	19.22

Table 2: Preliminary phytochemical screening test of stem bark of Terminalia arjuna

S.No	Chemical test	Shade drying samples		Sun drying samples		Oven drying samples	
		Methanolic extract	Aqueous extract	Methanolic extract	Aqueous extract	Methanolic extract	Aqueous extract
1.	Alkaloids	+	-	+	-	+	-
2.	Glycosides	+	+	+	+	+	+
3.	Flavonoids	+	+	+	+	+	+
4.	Steroids	+	+	+	+	+	+
5.	Triterpenoids	+	+	+	+	+	+
6.	Saponin	+	+	+	+	+	+
7.	Tanins	+	+	+	+	+	+
8.	Carbohydrates	+	+	+	+	+	+
9.	Proteins	-	-	-	-	-	-
10.	Amino acids	-	-	-	-	-	-

THIN LAYER CHROMATOGRAM OF Terminalia arjuna

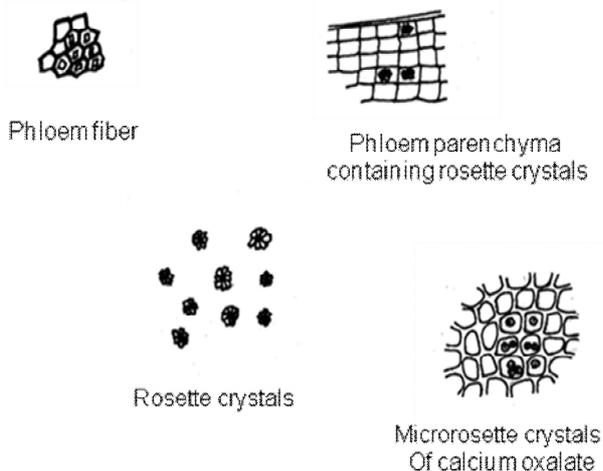
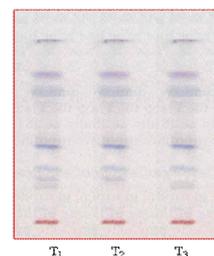


Figure 1: Microscopic study of stem bark of Terminalia arjuna



Tracks : T₁– Shade drying sample
 T₂– Sun drying sample
 T₃– Oven drying sample

Solvent System : Toluene: ethyl acetate : formic acid: methanol (6:3: 0.1:1.0)

Visualization : Anisaldehyde sulphuric acid

Figure 2: Thin Layer Chromatographic study of stem bark of Terminalia arjuna

Thin layer chromatographic study

TLC studies³² of the methanolic extract was carried out on aluminium plates precoated with silica gel 60 F₂₅₄ of 0.2 mm thickness using Toluene: ethyl acetate : formic acid: methanol (6:3: 0.1:1.0) as mobile phase and observed under visible light after derivatisation with anisaldehyde sulphuric acid (5%) followed by heating the plate at 110°C. The colour and R_f values of the resolved spots were noted.

RESULTS AND DISCUSSION

Macroscopic characters

All samples are flat or slightly curved. Outer surface smooth, pale greenish yellow, inner surface finely longitudinally striated and pinkish in colour, bitter taste and characteristic odour.

Microscopic characters (Powder)

Microscopic analysis of all the samples shows the presence of identifying diagnostic characters, that are phloem fiber, phloem parenchyma containing rosette crystals, rosette crystals and micro rosette crystals of calcium oxalate (Figure 1).

Physicochemical parameters study

Physicochemical parameters of *T. arjuna* are tabulated in Table 1. Loss on drying at 105°C is one of the major factors responsible for the deterioration of the drugs and formulations. Low moisture content is always desirable for higher stability of drugs. A high ash value is indicative of contamination, substitution, adulteration, or carelessness in preparing the drug or drug combinations for marketing. Water-soluble and alcohol soluble extractive value plays an important role in evaluation of crude drugs. Less extractive value indicates addition of exhausted material, adulteration or incorrect processing during drying or storage or formulating. The physicochemical parameters of barks of *T. arjuna* were determined as per the standard protocol. The results of loss on drying, total ash, acid insoluble ash, alcohol soluble extractive and water soluble extractive values of shade dried stem showed lower results when compared with sun dried and oven dried stem.

Thin Layer Chromatographic Study

Thin layer chromatogram of the methanolic extract of all samples developed after derivatisation with anisaldehyde sulphuric acid reagent. The shade drying sample showed nine major spots major spots at R_f 0.08(grey), 0.19 (pinkish blue), 0.23 (dark blue), 0.32 (blue), 0.42 (Dark blue), 0.45 (grey), 0.65 (grey), 0.71 (grayish blue) and 0.80 (dark pink), where as sun and oven drying samples showed eight spots. The spot at R_f 0.08 (grey) was absent on sun drying samples and the spot at R_f 0.19 (grey) was absent on oven drying samples (Figure 2).

CONCLUSION

Drying of plant material can be achieved by several processes, including shade, sun and oven drying. Overall results showed that drying methods determine the chemical quality of stem bark of *Terminalia arjuna*. The results of physicochemical parameters i.e. loss on drying, total ash, acid insoluble ash, alcohol soluble extractive and water soluble extractive values of shade dried sample showed lower values when compared with sun dried and

oven dried samples. Preliminary phytochemical analysis of all samples revealed the presence of Alkaloids, Glycosides, Flavonoids, Steroids, Triterpenoids, Saponins, Tanins and Carbohydrates. Thin layer chromatogram of the methanolic extract of shade drying sample after derivatisation with anisaldehyde sulphuric acid reagent showed nine major spots major spots, where as sun and oven drying samples showed eight spots. It concluded that shade drying of plant materials more suitable and is recommended as drying process showed a high potential in improving quantity and quality of medicinal plants.

REFERENCES

1. Tripathi VK, Singh B, Tripathi, RK, Upadhyay and Pandey VB. *Terminalia arjuna* its present status. Oriental J of Chem 1996; 1: 1-16.
2. Bone K. Clinical Applications of Ayurvedic and Chinese Herbs. Warwick, Queensland, Australia. Phytotherapy Press; 1996:131-133.
3. Kapoor LD. Handbook of Ayurvedic Medicinal Plants. Boca Raton, FL. CRC Press; 1990. pp 319-320
4. Singh N, et al. Mechanism of cardiovascular action of *Terminalia arjuna*. Planta Med 1982; 45:102-104.
5. Yadav RN, Rathore K. A new cardenolide from the roots of *Terminalia arjuna*. Fitoterapia 2001; 72: 459-461
6. Dwivedi S, Agarwal MP. Antianginal and cardioprotective effects of *Terminalia arjuna*, an indigenous drug, in coronary artery disease. J Assoc Physicians India 1994; 42: 287-289.
7. Khanna AK, Ramesh C, Kapoor NK. *Terminalia arjuna*: an Ayurvedic cardiotoxic regulates lipid metabolism in hyperlipidaemic rats. Phytotherapy Res 1996; 10: 663-665.
8. Kumar DS, Prabhakar YS. On the ethnomedical significance of the Arjun tree. J Ethnopharmacol 1987; 20:173-190.
9. Kandil FE, Nassar MY. A tannin anti-cancer promotor from *Terminalia arjuna*. Phytochem. 1998; 47: 1567.
10. Tripathi VK, Pandey VB, Udupa KN and Rucker G. Arjunolitin, a triterpene glycoside from *Terminalia arjuna*. Phytochemistry 1992; 31: 349.
11. Ahmad MU, Mullah KB, Nörin and Ulla TJK. Terminic acid, a new trihydroxytriterpene carboxylic acid from bark of *Terminalia arjuna*. Ind. J. Chem. 1983; 22B: 738-740.
12. Sharma PN, Shoeb A, Kapil RS and Popli SP. Arjunolone – a new flavone from stem bark of *Terminalia arjuna*. Ind. J. Chem.1982; 21B: 263-264.
13. Anjaneyulu ASR, Prasad AVR. Chemical Examination of the Roots of *Terminalia arjuna*. Phytochem 1982; 21: 2057-2060.
14. Row LR, Murty PS, Subba Rao GSR, Sastry CSP, Rao KVJ. Chemical examination of *Terminalia* species: part XII--isolation and structure determination of arjunic acid, a new trihydroxytriterpene carboxylic acid from the *Terminalia arjuna* bark. Ind. J. Chem., 1970; 8: 716-721.
15. Khanna AK, Chander C, Kapoor NK. *Terminalia arjuna*: an Ayurvedic cardiotoxic regulates lipid metabolism in hyperlipidemic rats. Phytotherapy Research 1996; 10: 663-665.
16. Dwivedi S, Udupa N. *Terminalia arjuna*: Pharmacognosy, Phytochemistry, Pharmacology and clinical use. A review. Fitoterapia 1989; 60:413-420.
17. Anjaneyulu ASR, Prasad AVR. Chemical examination of roots of *Terminalia arjuna* (Roxb.) Wight & Arnot. Part I. Characterisation of two new triterpenoid glycosides. Ind. J. Chem., 1982; 21B: 530-533.
18. Bharani A, Ganguly A and Bhargava KD. Salutary effect of *Terminalia Arjuna* in patients with severe refractory heart failure. Int. J. Cardiol., 1995;49: 191-199.
19. Shaila HP, Udupa SL, Udupa AL. Hypolipidemic activity of three indigenous drugs in experimentally induced atherosclerosis. Int. J. Cardiol., 1998; 67: 119-124.
20. Kumar V and Jain SK. A contribution to Ethnobotany of Surguja district in Madhya Pradesh, India. Ethnobotany, 1998; 10: 89-96.
21. Prusti AB and Behera KK. Ethnobotanical Exploration of Malkangiri District of Orissa, India. Ethnobotanical Leaflets, 2007; 1:12-15.

22. Yesodharan K and Sujana KA. Ethnomedicinal knowledge among Malamalasar tribe of Parambikulam wildlife sanctuary, Kerala. *Indian J. of Traditional Knowledge*, 2007; 6(3): 481-485.
23. Lorenzi H, Matos FJA. *Plantas medicinais no Brasil: nativas e exóticas*. Nova Odessa: Instituto Plantarum 2002. p. 512p.
24. Silva, F, Casali VWD. *Plantas Mediciniais e aromáticas: Pós-Colheita e Óleos Essenciais*. Viçosa-MG: UFV, DFT, 2000.p.135.
25. Ibanez E, Kubatova A, Senorans FJ, Cavero S, Reglero G, Hawthorne SB. Subcritical water extraction of antioxidant compounds from rosemary plants. *J Agric Food Chem*. 2003;1: 375-382.
26. Bartram T. *Encyclopaedia of Herbal Medicine*. Grace: Dorset. 1995.
27. Johanan DA. *Plant Micro Techniques*, 182, New york, Mcgraw-Hill, 1940.p. 89-93.
28. Anonymous, *Indian Pharmacopoeia*, 1966, 2nd ed. Government of India, New Delhi, 1996.p.23.
29. Trease EG, Evans WC. *Pharmacognosy*. 15th Edition, W.B. saunders, London. 2002. p. 137-149.
30. Kokate CK, Purohit AP, Gokhale SB. *Pharmacognosy*. 16th Edn. Nirali Prakashan, Pune, India;2001.p.105-114
31. Dahiru D, Onibiyi JA, Umaru HA. Phytochemical screening and antiulcerogenic effect of *Moringa oleifera* aqueous extract. *Afr. J. Traditional complementary Alternative med*. 2006; 3:70-75.
32. Stahl Igon. *Thin Layer Chromatography*, Springerverlag Berlin, New York; 1969.p. 843-850.

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