

WORLD JOURNAL OF PHARMACEUTICAL RESEARCH

SJIF Impact Factor 8.074

Volume 7, Issue 19, 1151-1159.

Research Article

ISSN 2277-7105

PHARMACOGNOSTICAL AND PHARMACEUTICAL ANALYSIS OF KHADIRASHTAKA GHANAVATI- AN AYURVEDIC POLYHERBAL FORMULATION

Dr. Anagha Sivanandan*¹, Drashti Shah², Prof. L. P. Dei³, C. R. Harisha⁴ and V. J. Shukla⁵

¹2nd Year MS Scholar, Department of Prasutitantra and Streeroga, IPGT & RA Jamnagar.

²2nd Year MD Scholar, Department of Kayachikitsa, IPGT & RA Jamnagar.

³Dean, HOD Department of Prasutitantra and Streeroga, IPGT & RA Jamnagar.

^{4,5}Head, Pharmacognosy Laboratory, IPGT & RA Jamnagar.

Article Received on 15 October 2018,

Revised on 05 Nov. 2018, Accepted on 26 Nov. 2018

DOI: 10.20959/wjpr201819-13779

*Corresponding Author
Dr. Anagha Sivanandan
2nd Year MS Scholar,
Department of Prasutitantra
and Streeroga, IPGT & RA
Jamnagar.

ABSTRACT

Khadirashtaka is a renowned formulation consisting of eight herbal drugs which had been mentioned in the context of treatment of skin disorders in various classical texts of Ayurveda like Yogaratnakara, Vangasena, Vrindamadhava, Chakradatta etc. Even though the combination had been suggested in the form of decoction in the classical texts, it is converted into the tablet (Ghanavati) form for addressing palatability issues and proper dose fixation. Methods: The finished product is subjected to microscopic evaluation, physic-chemical analysis like hardness, weight variation, loss on drying, ash value, Ph value, water soluble extract, alcohol soluble extract, High

Performance thin layer chromatography(HPTLC) etc. **Results:** Pharmacognostical study showed the presence of certain identifying characters of all of the eight ingredients in the formulation like cork cells of *Khadira*, Crystal fibers of *Nimba*, scleroids of Haritaki, Bibhitaki and Amalaki, pitted vessels of patola, starch cells of guduchi, multicellular trichome of vasa etc. Preliminary physico-chemical analysis showed that hardness 3.5 Kg/cm², ash value 14.88%, loss on drying 12.95%, water soluble extract 35.15 and HPTLC showed two spots in 254nm and one spot in 366nm. **Conclusion:** Present work was carried out to standardize the finished product *Khadirashtaka Ghanavati* in terms of its identity, quality and purity. Pharmacognostical and Physico-chemical observations revealed the specific characters of all active constituents in the preparation.

KEYWORDS: Khadirashtaka, Ghanavati, pharmacognosy, pharmaceutical, HPTLC.

INTRODUCTION

Khadirashtaka refers to a combination of eight herbal drugs namely Khadira (Acacia catechu (Linn.f) Willd), Haritaki (Terminalia chebula Retz), Bibhitaki (Terminalia bellerica Roxb.), Amalaki (Embilica officinalis Gaertn), Nimba (Azadirachta indica A.Juss), Patola (Trichosanthes dioica Roxb.), Guduchi (Tinospora cordifolia (Willd.)Miers) and vasa(Adhatoda vasica Nees). It is explained as an ideal formulation for different kinds of skin disorders in the classical texts of Ayurveda like Yogaratnakara, [1] Vangasena., Chakradatta, Vrinda Madhava, Vaidya Chintamani, Sahasrayoga ,Bharata Bhaishajya Ratnakara etc. In Ayurvedic parlance, the combination is having Tikta (bitter) and Kashaya (astringent) taste, Laghu (light) and Rooksha (rough) properties. The formulation is having the action of pacifying all three Doshas (Vata, Pitta, Kapha) and mainly Kapha-pittahara (pacifying Kapha and Pitta). Kushta (skin disorders), Kandu(itching), Visphota(eruption) are the main indications of *Khadirashtaka*, ^[2] explained in all of the above mentioned classical texts. Apart from these general therapeutic uses, Vangasena explains some specific indications like Visarpa, Pama, Kitibha, Shitapitta, Masoorika. [3] Bharata Bhaishajya Ratnakara mentioned its special therapeutic uses in Romantika and Masoorika. [4] In the classical texts, oral administration of Khadirashtaka in the form of Kwatha (decoction) is mentioned. But, here for better palatability and standardization of dose, it is converted into Ghanavati (tablet) form. The ingredients of Khadirashtaka Ghanavati with their proportions are described in Table 1. Pharmacognosy is also the first step to standardize a drug which is the need of the day. It should be noted that herbal drug standardization is not new in the field of Ayurveda. In the classics it is mentioned in a codified manner, such as Grahya Lakshana, Method of collection etc. It is a timely necessity followed by compulsion to go for quality control of the raw drugs as well as final products using modern parameters. This will not only provide a scientific basis and credibility to Ayurvedic drugs and pharmaceutics but also help in the globalization of Ayurveda.

AIMS AND OBJECTIVES

- 1. To evaluate collected raw drugs and *Khadirashtaka Ghanavati* for authenticity through various pharmacognostical procedures.
- 2. To develop the pharmacognostical and phyto-chemical profile of *Khadirashtaka Ghanavati*.

MATERIALS AND METHODS

Collection, Identification and Authentication of raw drugs

The raw materials were collected from the pharmacy of Gujarat Ayurved University, Jamnagar. All the raw drugs were identified and authenticated in the Pharmacogonosy Department, Institute for Post Graduate Teaching and Research in *Ayurveda*, Gujarat Ayurved University, Jamnagar.

Preparation of Drug

All the ingredient drugs were taken in equal quantity in the *Yavakut* form and *Kwataha* was made as per the classical guidelines. *Kwatha* was heated upto when it is converted into Ghana form and *Ghanavati* is made out of it.

Pharmacognostical study

The Pharmacognostical study comprises of organoleptic study and microscopic study of finished product.

Organoleptic Study

The Organoleptic characters of Ayurvedic drugs are very important and give the general idea regarding the genuinity of the sample. Organoleptic parameters ie. Taste, Colour, odour and touch of *Khadirashtaka Ghanavati* were scientifically studied following standard references.^[5]

Microscopic Study

Khadirashtaka Ghanavati was powdered and dissolved with water and microscopy of the sample was done without stain and after staining with Phloroglucinol + HCl. Microphotographs of Khadirashtaka Ghanavati was also taken under Corl-zeisstrinocular microscope. [6]

Physico-chemical analysis

Khadirashtaka Ghanavati was analyzed using various standard physico-chemical parameters. The common parameters mentioned for compressed tablets in Ayurvedic Pharmacopia of India,^[7] and CCRAS,^[8] guidelines are total ash, pH value and water and alcohol soluble extractives. On this basis these parameters were taken. Presence of more moisture content in a sample can create preser*vati*on problem. Hence loss on drying was also selected as one of the parameters.^[10]

High Performance Thin Layer Chromatography (HPTLC)

HPTLC was performed as per the guideline provided by API. Methanolic extract of drug sample was used for the spotting. HPTLC was performed using Toluene +Ethylacetate + Acetic acid (7:2:1) solvent system and observed under visible light. The colour and Rf values of resolved spots were noted.^[10]

RESULTS AND DISCUSSION

Organoleptic characters of Khadirashtaka Ghanavati

Organoleptic characters contents of *Khadirashtaka Ghanavati* like colour, taste, touch, Odor were recorded and shown in Table- 2.

Microscopic Study

Identifying characters of ingredients of *Khadirashtaka Ghanavati* under the microscope showed cork cells, stone cells, rhomboid crystals and tannin content of *Khadira*, Crystal fibers, prismatic crystals, tannin content and stone cells of *Nimba*, scleroids, epicarp cells and pitted stone cells of *Haritaki*, scleroids and trichomes of *Bibhitaki*, scleroids and silica deposition of *Amalaki*, pitted vessels and simple trichome of patola, starch grains, border pitted vessels, and cork cells of *Guduchi*, multicellular trichome and fragment of cystolith of *Vasa*. All these are showed in Plate 1(a to z).

Physico-chemical analysis

Physico-chemical analysis of *Khadirashtaka Ghanavati* revealed the value as hardness 3.5 Kg/cm², ash value 14.88%, loss on drying 12.95%, water soluble extract 35.15%, alcohol soluble extract 18.34%, pH 5.5 are shown in Table –3.

HPTLC Study

The chromatographic study (HPTLC) was carried out under 254 and 366 nm UV to establish fingerprinting profile. It showed 2 spots at 254 nm and 1 spot at 366nm with $R_{\rm f}$ values were recorded which may be responsible for expression of its pharmacological and clinical actions. Plate 2, Table -4.

Table 1: Ingredients of Khadirashtaka Ghanavati.

Sr.no	Drug	Botanical name	Part used	Ratio
1.	Khadira	Acacia catechu(Linn.f)Willd	Sara/twak	1
2.	Haritaki	Terminalia chebula Retz.	Phala	1
3.	Amalaki	Embilica officinalis Gaertn	Phala	1
4.	Bibhitaka	Terminalia bellerica Roxb.	Phala	1
5.	Nimba	Azadirachta indica A.Juss	Twak	1
6.	Patola	Trichosanthes dioica Roxb.	Patra	1
7	Guduchi	Tinospora cordifolia(Willd.)Miers	Kanda	1
8.	Vasa	Adhatoda vasica Nees	Patra	1

Table 2: Organoleptic parameters of Khadirashtaka Ghanavati

Sl. no:	Character	Observation
1	Colour	Black
2	Odour	Slight fragrant
3	Taste	Astringent
4	Touch	Hard

Table 3: Physico-chemical analysis of Khadirashtaka Ghanavati

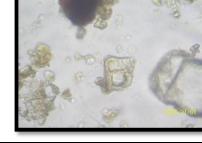
Sl no:	Test	Result
1	Hardness	3.5 kg/cm2
2	Weight Variation	Average Weight-520mg Highest weight-553mg Lowest weight-485mg
3	Ash Value	14.88%
4	Loss on drying	12.95%
5	Water Soluble Extract	35.15%
6	Alcohol Soluble Extract	18.34%
7	рН	5.5
8	Disintegration time	30 minutes

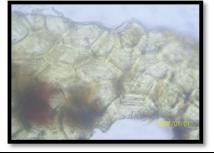
Table 4: HPTLC Study of Khadirashtaka Ghanavati

Wave length	Number of spots	Rf value
254nm	2	0.02,0.09
366nm	1	0.02

Plate no: 1

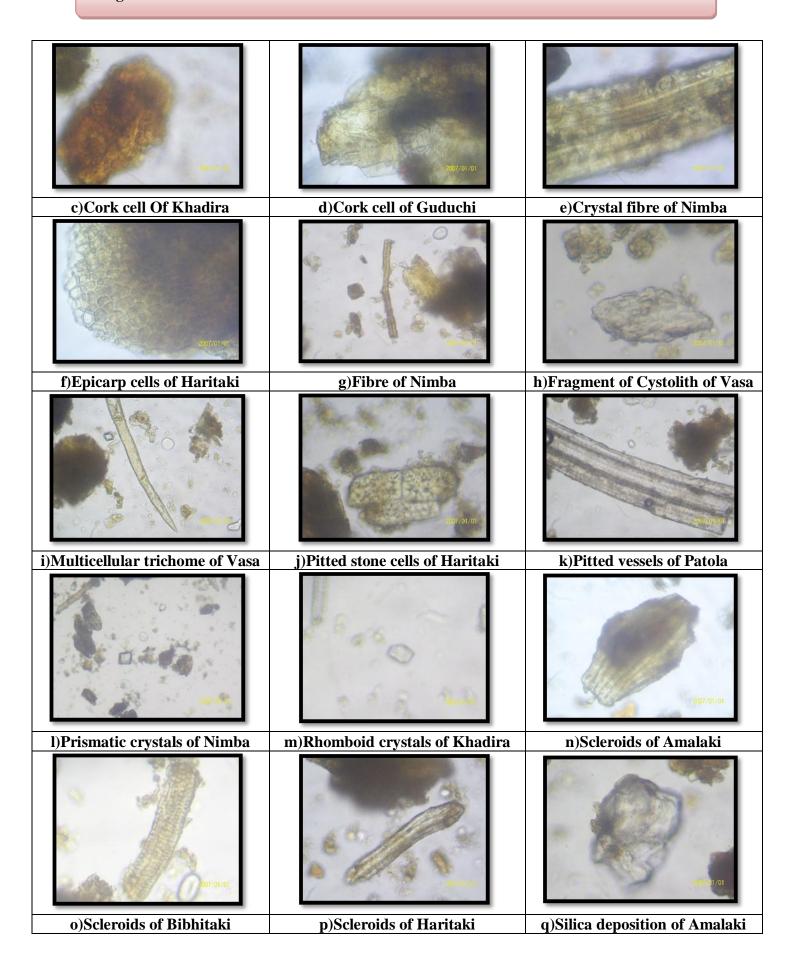






Sample a)Collenchyma of Guduchi

b)Cork cell of Guduchi



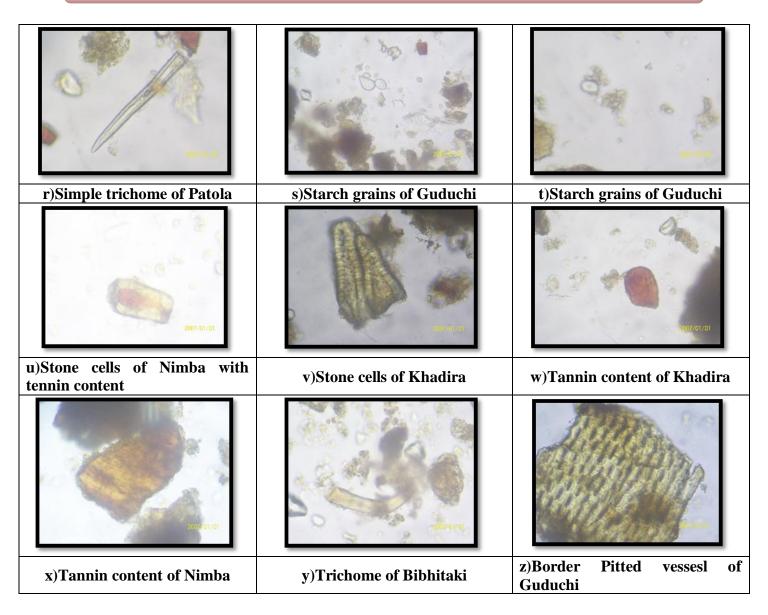
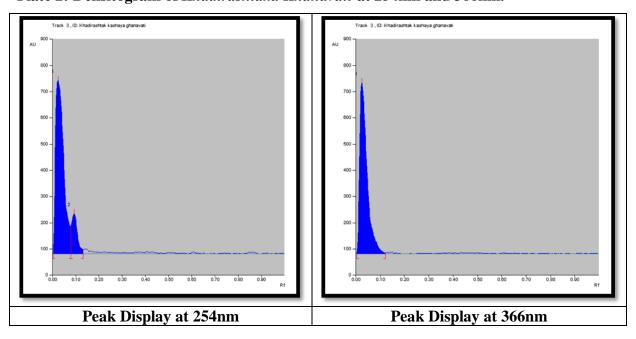


Plate 2: Densitogram of Khadirashtaka Khanavati at 254nm and 366nm.



1157

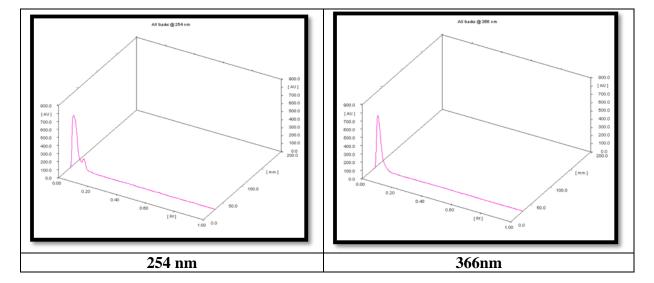


Plate 3: Three dimensional HPTLC (3D) Densitogram

CONCLUSION

The pharmacognostical and physico chemical analysis of *Khadirashtaka ghanavati* confirmed the purity and genuinety of the drug. As there is no published information available on pharmacognostical and physic-chemical profiles of *Khadirashtak Ghanavati*, this preliminary information may be beneficial for future researchers and can be used as a reference standard in the further quality control researchers.

REFERENCES

- 1. Dr. Indradevatripathi and Dr. Dayashankaratripathi, editors, Yogaratnakara, kusthachikitsa/63 2nd edition, Varanasi, Chaukhambha Prakashana, 2007; 648.
- 2. Dr. Indradevatripathi and Dr. Dayashankaratripathi, editors, Yogaratnakara, kusthachikitsa/63 2nd edition, Varanasi, Chaukhambha Prakashana, 2007; 648.
- Vaidya Shankarlalji Jain, Editor, Vangasena, commentary by Kavivar Shaligramji Vaishya, Version 1A, Mumbai, Khemraj Streekrishnadaas Publication, Reprint, 2003; 625.
- 4. Pandit Nivaranchandra Bhattacharya, Editor Bharata Bhaijya Ratnakara by Nagin Das Chandralal Shah, Bhavaprakashika commentary by Pandit Gopinath Gupta, New Delhi, B.Jain publishers, Reprint, 2012; 1: 321.
- 5. Trease and Evans, Pharmacognosy, 15th Ed., W. B. Sunders Company Ltd., 1996; 569: 570.
- 6. Wallis TE, Text book of Pharmacognosy, 5th Ed., New Delhi: CBS Publishers & Distributors, 2002; 123-132: 210-215.

- 7. Anonymous, Protocol for testing of Ayurveda, Siddha & Unani medicines, Pharmacopoeial laboratory for Indian medicines, Ghaziabad, Ministry of AYUSH, Government of India.
- 8. Anonymous, Parameters for qualitative assessment of Ayurveda, Siddha drugs, CCRAS, New Delhi, 2005.
- 9. Anonymous, The Ayurvedic Pharmacopoeia of India, Part II (Formulation), Volume I, First edition, Ministry of AYUSH, Government of India, New Delhi, 2007; 140-147.
- 10. Anonymous Anonymous, Planner Chromatography, Modern Thin layer Chromatography, Switzerland, 1999; 2-16.