



**ORIGINAL RESEARCH ARTICLE** 

# Pharmaceutico-Analytical Study of A PolyherbalAyurvedicFormulationBhallatakaadiTailaIndicated in DadruKushta

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# ABSTRACT

Skin conditions pose significant threat for the person's well-being, mental health, ability to function and social participation. Global Burden of Disease project, has shown that skin diseases are the 4<sup>th</sup> leading cause of non-fatal disease burden world-wide<sup>1</sup>. Superficial fungal infections contribute huge part among the commonly seen dermatological disorders. Worldwide prevalence of Dermatophytosis is about 20000-25000 individuals per 100000, whereas incidence ranges from a low of 10000 per 100000 persons to a high of 15000 per 100000 persons. It is predominance in about 20-25% of the world population. In India, according to recent study prevalence ranges from 36.6-78.4%. One of the reasons for concern is its unresponsiveness to treatment in majority of the cases and the early recurrence. The new drugs have to be discovered and tried in this disease. *Bhallatakaadi Taila*isone of the unutilized new formulations, indicated in the *Dadru Kushta* for *Bahya Parimarjana Chikitsa*, as *Abhyanjana*. It has 14 different ingredients among which *Bhallataka* and *Gunja* are *Upavisha Dravya*. Other ingredients include, *Aksha, Kushta, Triphala, Trikatu, Panchalavana*. The method of preparation adopted was general *Sneha Kalpana Vidhi* as per *Sharangadhara Samhita*. This study is about pharmaceutico-analytical standardization of *Bhallatakaadi Taila*.

Key WordsBhallataka, Bhallatakaadi Taila, Dadru Kushta, Gunja, Pharmaceutico-Analytical study

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# INTRODUCTION

Ayurveda, being the deposit of many unexcavated therapeutics, has various formulations which can be used for treatment of *Dadru*that has similar presentation with dermatophytosis. Superficial fungal infections contribute huge part among the commonly seen dermatological disorders. Worldwide prevalence of dermatophytosis is about 20000-25000 individuals per 100000, whereas incidence ranges





Step 1; Jala Nimajjana



Step 2; Gomutra Sthapana





# Day 1Day 2Day 3Day 4Day 5Day 6Day 7

Step 3; Goksheera Sthapana



Step 4; Ishtika Choorna /Gharshana



Figure 1 Shodhana of Bhallataka





	DOTANICAL NAME		
DRUGS REQUIRED	BOTANICAL NAME	PART USED	QUANTITY
Bhallataka	Semicarpus anacardium	Seeds	5.1428 kg
Maricha	Piper nigrum	Seeds	1.7144 kg
Shunti	Zingiber officinale	Rhizome	1.7144 kg
Pippali	Piper longum	Seeds	1.7144 kg
Aksha (Bhibhitaki)	Terminalia bellarica	Fruit	6.8572 kg
Kushta	Saussurealappa	Whole plant	5.1428 kg
Gunja	Abrus precautarius	Seeds	5.1428 kg
Panchalavana (Saindhava,	-	-	Each 1.02828 kg
Sauvarchala, Saamudhra, Vida,			-
Romaka)			
Haritaki	Terminalia chebula	Fruit	1.7144 kg
Amalaki	Phyllanthus emblica	Fruit	1.7144 kg
Amoorchita Tila Taila	-	_	16 liters

from 10000 to 15000 per 100000 persons. It is predominant in about 20-25% of the world population<sup>1</sup>. In India, according to recent survey prevalence ranges from 36.6-78.4%<sup>2</sup>. Over the last few decades, the frequency of the cases has increased alarmingly constituting at least 5-10% of cases showcasing in dermatology OPD.Sneha Kalpana holds an important place, amidst the various dosage forms of external medications. Advantages of Sneha *Kalpana* areamplified sustainable absorption, longer bioavailability of medicaments and extraction of fat soluble as well as a watersoluble active principle at a time in a single formulation. Bhallatakaadi Taila is the polyherbal formulation selected from Harita Samhita<sup>3</sup>, indicated in Dadru Kushta for Abyanjana. It has 14 ingredients viz., Bhallataka, Gunja, Aksha, Kushta, Triphala, Vyosha, Panchalavana. Among Bhallataka these Gunja and areUpavisha<sup>4</sup>. The Teekshna, Ushna Guna of the Taila acts as counter irritant against the dermatophytes. The present study aims to focus

on the pharmaceutico-analytical study of *Bhallatakaadi Taila* with various parameters.

# AIMS AND OBJECTIVES

Pharmaceutical and Analytical profiling of *Bhallatakaadi Taila*.

# MATERIALS AND METHODS

The *Taila* was prepared in the S.D.M Ayurveda Pharmacy, Kuthpady, Udupi. The drug *Bhallataka* was procured from pharmacy and *Shodhana*was done by method mentioned in API. After *Shodhana*,*Taila* was prepared in S.D.M. Pharmacy.

#### A. Pharmaceutical study;

#### A.1. ShodhanaofBhallataka;

The nuts were introduced to the process of Jala Nimmajjana<sup>4</sup> and those which soak in water were selected for the preparation of the oil. Thenuts were subjected to GomutraSthapana for 7 consecutive days, each day the Mutra was replaced. After 7<sup>th</sup> day they were washed and subjected to GoksheeraSthapana for 7

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consecutive days, replacing the *Ksheera* for 7 days. On 14<sup>th</sup> day nuts were washed and dried. Followed by *IshtikaChoorna Gharshana*<sup>5</sup>for one day to remove the remaining vesicants. Further, nuts were washed and dried for *Taila* preparation. Various processing steps of *Bhallataka Shodhana* is depicted in Figure No.-01.

Oil was prepared by the reference of *Taila Paaka Vidhias* mentioned in *SharangdharaSamhitaMadhyama Khanda*<sup>6</sup>. The ingredients and quantity of the raw drugs used in the *Taila* is mentioned in Table No.-01.

*Kalka*was prepared byusing drugs such as *Shodhita Bhallataka, Gunja, Triphala, Vyosha, Panchalavana, Kushta.* One part of*Kalka*, 4 parts of *Tila Taila* and 16 parts of *Kashaya*wereheated on mild flame until the *Taila Siddhi Lakshana* was obtained. *Sneha Siddhi Pariksha* was done to check if any water content was left in the *Taila*.As per the classics the *Khara Paka* was done, as it was for external application<sup>7</sup>. The process of preparation of *Bhallatakaadi Taila* is shown in Figure No.-02. The Rasa-panchaka of the drugs are mentioned in Table No.-02 and

В.	Analyti	cal Study of	Bhal	latakaadi taila.
Orga	anoleptic	characters	and	Physio-chemical
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made during the preparation of Taila.

parameters- Refractive index, Specific gravity, Viscosity, Acid value, Saponification value, Iodine value, saponification value, Unsaponifiable matter, Peroxide value and HPTLC.

Table No.-03 explains regarding the observations



Figure 2 Preparation of Bhallatakaadi Taila

SI No	Drug	Rasa	Guna	Vipaka	Virya	Karma	Doshagnata
1.	Bhallataka <sup>8</sup>	Katu Tikta Kashaya	Teekshna Laghu Snigdha	Madhura	Ushna	Kushtagna, Arshoghna, Deepani, Bhedani, Mutra- Sangrahana	Kaphavata Shamaka
2.	Gunja <sup>8</sup>	Tikta, Kashaya	Laghu, Ruksha, Tikshna	Katu	Ushna	Keshya	Kaphavata Shamaka
3.	Kushta <sup>8</sup>	Katu, Tikta, Madhura	Laghu Rooksha Teekshna	Katu	Ushna	Sukrala, Kushtaghna Lekhaneeya,	Vata – Kapha Shamaka

 Table 2 Rasa-panchaka of the drugs

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						Vrshya	
4	Shunti <sup>8</sup>	Katu	Guru, Rooksha, Teekshna	Madhura	Ushna	Deepana, Bhedana	Vata- Kaphahara
5	Maricha <sup>8</sup>	Katu, Tikta	Teekshna, Rooksha	Madhura	Naatyushna	Deepana, Avrshya, Pramathi	Kaphavatahara
6	Pippali <sup>8</sup>	Katu	Snigdha, Laghu	Madhura	Anushna	Deepana, Rasayana, Pachana	Vata- Shleshmahari
7.	Pathya <sup>8</sup>	Kashaya	Ruksha Laghu	Madhura	Usna	Anulomani, Deepana, Pachana, Rasayana	Tridoshahara
8.	Amalaki <sup>8</sup>	Amla	Ruksha Laghu	Madhura	Sita	Vrishya, Rasayana, Jwarahara, Raktapittagna	Tridosha Shamaka
9.	Vibhitaki <sup>8</sup>	Kashaya	Ruksha Laghu	Madhura	Usna	Bhedana, Kasahara, Krimighna, Netrya	Tridoshahara
10.	Saindhava Lavana <sup>9</sup>	Lavana, Madhura ,	Laghu, Snigdha, Sukshma	Madhura	Sheeta	Deepana, Pachana, Ruchya, Vrishya, Nethrya	Tridoshahrt
11.	Sauvarchala Lavana <sup>9</sup>	Lavana	Snigdha Laghu Vishada	Madhura	Ushna	Rochana, Hridya,Bhedana, Deepana, Pachana	Vatanuth, Pittala
12.	Vida <sup>9</sup>	Lavana	Laghu Ruksha Vyavayi Teekshna	Madhura	Ushna	Deepana Ruchya	Kapha Vata -Anulomana
13.	Romaka <sup>9</sup>	Madhura	Laghu Teekshna Sukshma	Katu	Ushna	Bhedi Abhish -yandhi	Vataghna Pittala
14.	Saamudhra Lavana <sup>9</sup>	Madhura Katu Lavana	Guru Snigdha	Madhura Katu	Ushna	Deepana Bhedi Saksharam Vidahi	Shleshmala Vatanut

Table 3Observations made during preparation of *Bhallatakaadi Taila*;

Sl. No.	Observation Parameters	Observations
1.	Total quantity of <i>Kalka</i> taken	4 kg
2.	Total quantity of <i>Kwatha</i> taken	64 litres
3.	Quantity of oil	16 litres
4.	Temperature of oil during addition of Kalka	90 <sup>0</sup> C
5.	Temperature of oil during adding of Kwatha	85 <sup>0</sup> C
6.	Average temperature Day 1	101 <sup>0</sup> C
7.	Average temperature Day 2	$100 {}^{0}\mathrm{C}$





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8.	Average temperature Day 3	100 °C
9.	Average temperature Day 4	98 <sup>0</sup> C
10.	Duration of Heating Day 1	4 hours
11.	Duration of Heating Day 2	3 hours 30 minutes
12.	Duration of Heating Day 3	4 hours
13.	Duration of Heating Day 4	4 hours 30 minutes
14.	Total duration of heating	15 hours
15.	Obtained quantity	15.2 litres
16.	Weight loss	0.8 litre
17.	% Weight loss	5%
18.	Temperature during filtration	37 °C

#### Organoleptic Characteristics of Bhallatakaadi

#### Taila-

Colour - Blackish brown

Touch-Smooth

Taste –not applicable

Appearance – Translucent

Clarity - not clear

Smell - Pungent smell

#### **B.1. Methodology**

The following parameters were calculated using the standard procedures.

B.1a. Refractive index<sup>10</sup>

B.1b.Specific gravity<sup>10</sup>

B.1c. Viscosity<sup>10</sup>

B.1d. Acid value<sup>10</sup>

B.1e. Saponification value<sup>10</sup>

B.1f. Iodine value<sup>10</sup>

B.1g. Determination of Unsaponifiable matter<sup>10</sup>

B.1h. Peroxide value<sup>10</sup>

B.1i. Sample preparation for HPTLC<sup>10</sup>

Sample obtained in the procedure for the determination of unsaponifiable matter was dissolved in 10 ml of chloroform this was followed for the sample of *Bhallatakaadi Taila*, and chloroform soluble portion was used for HPTLC.

HPTLC:11,12,13

Aluminium plates were applied with the samples of 3, 6, 9µl of the chloroform fraction of *Bhallatakaadi Taila*to a band width of 8mm using Linomat 5 TLC applicator. Double trough chamber CAMAG in a solvent system Toluene – Ethyl acetate (9:1) was used to develop the plates which were visualized under short UV, long UV and after derivatisation in vanillin-sulphuric acid spraying reagent. And scanned under UV 254nm, 366nm and 620nm (Post derivatisation).  $R_{f}$ , colour of the spots and densitometric scan were recorded.

 Table 4
 Results of standardization parameters for Bhallatakaadi Taila

Parameter	<b>Results</b> $n = 3$	
	Bhallatakaadi taila	
Refractive index	1.33217	
Specific gravity	0.4091	
Viscosity	39.38	
Acid value	137.5	
Saponification value	140.93	
Iodine value	97.29	
Unsaponifiable matter	4.40	
(%w/w)		
Peroxide value	1.2	

## **OBSERVATIONS AND RESULTS**

The results of analytical study of the *Bhallatakaadi Taila*are mentioned in the Table No.-04.The R<sub>f</sub> values of the HPTLC scan are explained in the Table No.-05.The HPTLC photo documentation results are depicted in Figure No.-March 10<sup>th</sup> 2022Volume 16, Issue 2 **Page 148** 







03. And Figure No.-04 shows the densitometric

scan of the study drug.

Short UV	Long UV	Post derivatisation
0.08 (Green)	-	-
-	-	0.11 (Purple)
0.30 (Green)	-	0.30 (Purple)
0.34 (Green)	0.34 (F. blue)	-
0.42 (Green)		0.42 (Purple)
-	0.48 (F. blue)	-
0.54 (Green)	-	0.54 (Purple)
0.59 (Green)	0.59 (F. blue)	0.59 (Purple)
-	0.63 (F. green)	-
0.74 (Green)	-	-
-	0.80 (F. blue)	-
0.84 (Green)	-	0.84 (Purple)
0.88 (Green)	-	-
-	0.96 (F. blue)	-
*F- fluor	rescent	

#### **B.3. Remarks**

The given sample of *Bhallatakaadi Taila*has been standardized as per standard testing protocol. The results of standardization parameters and HPTLC Photo-documentation,  $R_f$ values and Densitometric scan are given in respective tables and figures.



Solvent system - Toluene: Ethyl acetate (9.0:1.0) Track 1 – *Bhallatakaadi taila* – 3μl Track 2 – *Bhallatakaadi taila* – 6μl Track 3 – *Bhallatakaadi taila* – 9μl **Figure 3**: HPTLC photo documentation of chloroform

fraction of Bhallatakaadi Taila



At 620nm

Figure 4 Densitometric scan of Chloroform fraction of *Bhallatakaadi Taila* 

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## DISCUSSION

Semecarpus anacardium Linn.is an organic irritant toxic drug which may induce hypersensitivity in certain individuals, а Shodhanais compulsory processing step conducted to control its default irritant property.Dadru is a Kshudra/Maha Kushta manifested predominantly by vitiated Pitta-Kapha Dosha. The ingredients in Bhallatakaadi Taila such as viz., Bhallataka, Kushta, Gunja, Trikatu, Triphala, Panchalavana alleviates the Dosha and helps in mitigating the Lakshana. Clinical features of Dadru Kushta simulating with Dermatophytosis is caused due to the growth of fungal species on superficial layer of skin. The irritant drugs like Bhallataka and Gunja, act as counter irritants in the superficial mycoses in the dermatophytosis.

Refractive index indicates the possible chances of rancidity in duecourse of time in the oil. Higher is the refractive index, higher is the chances of spoilage due to oxidation. Specific gravity of the oil indicates the weight per ml of oil. This helps in the dosefixation of the medication, as 5 ml of *Bhallatakaadi Taila* contains 2.0455 g of the oil. Viscosity is the resistance to flow in a liquid. As the viscosity of the oil is increased the duration of oil retained in the body for absorption is longer. The oil is stored in adipose tissue prior to get released into the blood stream.Acid value is the relative measure of rancidity as free fatty acids are normally formed during decomposition of triglyceride esters into glycerol and fatty acids. Higher is the acid value higher is the rancidity. Saponification value indicates the components of the fatty substances which are capable of forming soaps when treated with alkali. Unsaponifiable matter includes non-volatile components such as alkanes, sterols etc, also this fraction may involve environmental contaminants and residues such as plasticizers, pesticides, mineral oil hydrocarbons and aromatics. Iodine value determines the amount of unsaturation in oil, mainly due to double bonds which are reactive towards halogens. Higher is the iodine value, more is the unsaturation in the oil.

HPTLC plates when observed under short UV showed 6 bands with different intensity of green. Under long UV it showed 6 bands (all fluorescent blue). Post derivatization with spraying reagent VSA reagent it showed 6 bands (all purple). Densitometric scan at 254 nm showed 10 peaks, among which  $R_f$  0.61 (19.27%) was the major peak. At 366 nm  $R_f$  0.55 (55.05%) was the major constituent. Post derivatization when scanned at 620 nm it showed 3 peaks with absorption around average 25% area namely at  $R_f$  0.58 (27.77%),  $R_f$ 0.33 (26.48 %),  $R_f$  0.48 (22.46%) were the majorly identified possible constituents of *Bhallatakaadi Taila*.

# CONCLUSION

*Bhallatakaadi Taila*having*Dadrugna* property is been standardised pharmaceutically and analytically,being a novel formulation, which was not been standardised in the recent past, was







found pharmaceutically viable and could be carried further for clinical intervention but the safety aspect should be monitored. Further a patch test can be performed to rule-out any adverse drug reaction in sensitive patient population.





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