RESEARCH ARTICLE

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# Analysis of *Madhuraskandha's* Rasayana Drugs through Pharmacognostical, Physicochemical and Phytochemical Parameters

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## Abstract

Acharya Charaka has classified drugs based upon the *Rasa;* as *Rasaskandha* in the context of *Asthapana Bastidravyas*. The drugs included in the *Skandha* are not only based upon their *Rasa* but also according to their *Vipaka* and *Prabhava*. *Madhurarasadravya* attributes actions like *Rasayana*, *Balya*, and *Jivaniya Karma*. Charaka had stated another therapeutic classification containing *dashemani* dravyas. In these groups of drugs, *Rasayana* karma may be attributed to *Jeevaniya*, *Balya*, *Brihmaniya*, *Vayasthapanadashemani* groups. In the present study, a list of ten *Rasayana* drugs was drawn by comparing *Vayasthapana gana* and *Madhuraskandha* drugs i.e. *Atibala*, *Vidari*, *Kantakari*, *Eranda*, *Gokshura*, *Guduchi*, *Shalaparni*, *Jivanti*, *Shatavari* and *Punarnava*. These drugs were evaluated by Pharmacognostical, Physicochemical and Phytochemical studies. The microscopical study of the powder showed the presence of starch grains and calcium oxalate crystals. Physicochemical parameters showed that water soluble extractive value is more than alcohol soluble extractive value. Majority of these drugs showed presence of Carbohydrates, Reducing sugar, Amino acids, Proteins, Tannin, Steroids, Triterpenoids, Anthraquinone, and Saponin. These can be used as standard parameters to generate purity and quality of the herbal drugs.

### Keywords

Madhuraskandha, Pharmacognostical, Phytochemical, Starch, Carbohydrates



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# INTRODUCTION

Acharya Charakahas classified drugs into 50 groups based upon pharmacological activity and rasa i.e., six skandhas were described. Among these *skandhas* maximum numbers of drugs were quoted in Madhuraskandha i.e., eighty five drugs. In these skandha, dravyas were arranged not only based upon their *Rasa* but also according to their *Vipaka* Prabhava. indicates and It that Madhuraskandha contains a drug which may have Madhurarasa or Madhuravipaka or *Madhuraprabhava*<sup>1</sup>.

Madhurarasa drugs mainly have Madhuravipaka and Sheetaveerya. They have Saptadhatuvardhan, Indriyaprasadana, Balavarnakara, Keshya, Kanthya, Balya, Brihmaniya, Preenana, Tarpana, and Jeevaniya activities<sup>2</sup>.

Secondly, based upon pharmacological action, Charaka had described a group of drugs having *Rasayana* karma referred as 'Vayasthapana dashemani'. Most of the drugs of this group Madhuraskandha. included under are Vayasthapana dashemani contains Amruta, Abhaya, Dhatri, Mukta (Rasna), Shweta (Aparajita), Jivanti, Atirasa, Mandukaparni, Sthira (Shalaparni), Punarnava<sup>3</sup>.It is noted

that 6 drugs namely *Guduchi, Shatavari, Punarnava, Shalaparni, Jivanti, Shweta* (*Aparajita*) are common to both the groups (*Madhuraskandha* and *Vayasthapana dashemani*).

Bala, Atibala, Shravani, Mahashravani, Rajadana, Ashwagandha, Vidari, Kshiravidari, Brihati, Kantakari, Eranda, Yashtimadhu, Mridvika, Kashmari, Sariva, Kharjura, Parushaka, Aatmagupta, Gokshura, Kakanasa etc. are the drugs, having Rasayana property, which were not included in Vayasthapana dashemani.

A list is prepared by including five drugs from each group i.e. drugs uncommon and common between *Madhuraskandha* and *Vayasthapana dashemani*. In the present study, the following ten drugs i.e. *Atibala*, *Vidari, Kantakari, Eranda, Gokshura, Guduchi, Shalaparni, Jivanti, Shatavari, Punarnava* are selected to formulate a group having *Rasayana* activity.

The World Health Organization (WHO) has appreciated the importance of medicinal plants for public health care in developing nations and has evolved guidelines to support the member states in their efforts to formulate national policies on traditional medicine and to study their potential usefulness including evaluation, safety, and efficacy. Keeping this in view, attempt has been made to evaluate the pharmacognostical, physicochemical and phytochemical analysis of these ten drugs, to document the qualitative information about the purity and quality.

## MATERIALS AND METHODS

#### **Collection of drugs**

The drugs were collected in their respective season according to the part used. The details are mentioned in Table 1. All the collected drugs were washed, cleaned and dried in shade for several days.

### **Preparation of powder**

The shade dried drugs were pulverized individually with help of grinder and passed through Mesh no.85 to obtain a fine powder<sup>4</sup>.

### **Preparation of extracts**

About 5g of the test drug powders were macerated with methanol (100ml) in a closed flask for 24 hours with frequent shaking for first 6hrs and kept it for 18 hrs. After 24 hours it was filtered and alcoholic extracts were collected in semisolid form. The same procedure was followed to obtain aqueous extracts of the test drugs<sup>7</sup>.

#### **Organoleptic characters**

Organoleptic characters of 10 test drugs such as odor, taste, texture and color were observed.

#### **Powder microscopy**

The characteristic of test powder was found by taking pinch of powder on a glass slide. One drop of Chloral-hydrate solution was put on a slide and covered it with a cover slip, heated over a low flame for a short time<sup>5</sup>. The specimen is treated with Phloroglucinol + Conc. HCl (for Calcium oxalate crystal, lignified cell), Ruthenium red (for mucilage), FeCl<sub>3</sub> to (tannin) and Iodine (for starch grains) <sup>6</sup>. The sample was observed under compound microscope (QUASMO, India) at 10X followed by 40X magnification and photographs were taken by using Kodak Easy Share C140, 8.2 megapixel 3x Optical/5x Digital Zoom HD camera. The same procedure is followed for all the ten test samples.

#### **Physicochemical parameters**

Physico-chemical parameters like loss on drying, ash value, alcohol soluble extractive and water-soluble extractive values and pH were determined as per the API guidelines for the all the test samples<sup>7</sup>.

### Phytochemical parameters

A preliminary phytochemical study of methanolic and aqueous extracts of ten test

drugs was carried out. Presence of various phyto-constituents viz., alkaloids, starch, proteins, amino acids, cardiac glycosides, flavonoids, phenols, saponins, steroids, tannins, phenolic compound and amino acids were evaluated <sup>8, 9</sup>.

#### Sample coding

All the ten samples are coded as follows: S1- Abutilon indicum, S2- Pueraria tuberosa, S3- Solanum xanthocarpum, S4-Ricinus communis, S5- Tribulus terestris, S6- Tinospora cordifolia, S7- Desmodium gangeticum, S8- Leptadenia reticulata, S9-Asparagus racemosus, and S10- Boerhavia diffusa.

## RESULTS

### **Organoleptic characters**

Organoleptic characters of ten test samples i.e., taste, color, odor and texture are mentioned in Table 2. Majority of the drugs are of *Madhura* (Sweet), *Tikta* (Bitter) and *Kashaya* (Astringent) Rasa.

#### **Powder Microscopy**

Microscopic characteristics of all the test samples are described in Table 3 (Plate 2 & 3)

### **Physicochemical parameters**

The physicochemical characters like loss on drying, ash value, water soluble extractive

value and alcohol soluble extractive value had been carried out by using crude drugs and pH was determined by using 5% aqueous solution. The results of physicochemical characters are as mentioned in Table 4.

## **Phytochemical parameters**

Qualitative analysis was carried out by using methanolic and aqueous extracts of all test samples. The test samples were evaluated for carbohydrate, amino acids, proteins, starch, protein, alkaloid, tannin, steroid, flavonoids etc. their results are as quoted in Table 5.

# DISCUSSION

Powder microscopy of all the test drugs were carried out to identify and authenticate the drugs. Among ten tests drugs, samples namely S1,S2, S4, S6, S7, S9, S10 have shown the presence of simple, compound, with or without hilum starch grains, starch grains with concentric lines; all the test drugs except S3 have revealed the presence of calcium oxalate crystals like prismatic, rosette, acicular and rhomboidal crystals.(Plate 2& 3)

No.	Individual	Botanical Source	Part	Time of	Place of
	drug		Used	collection	collection
1.	Atibala	Abutilon indicum Linn. Sweet	Root	May 2014	Periphery of
					Jamnagar
2.	Vidari	Pueraria tuberosa DC.	Tuber	Nov 2014	Junagadh
3.	Kantakari	Solanum xanthocarpum S. &W	Whole	July 2014	Periphery of
			plant		Jamnagar
4.	Eranda	Ricinus communis Linn	Root	May 2014	Periphery of
					Jamnagar
5.	Gokshura	Tribulus terrestris Linn	Fruit	Oct 2014	Periphery of
					Jamnagar
6.	Guduchi	Tinospora cordifolia (Willd.) Miers	Stem	Sept 2014	Periphery of
		ex Hook. f. & Thoms			Jamnagar
7.	Shalaparni	Desmodium gangeticum DC.	Root	July 2014	Junagadh
8.	Jivanti	Leptadenia reticulata W. & A.	Leaves	July 2014	Periphery of
					Jamnagar
9.	Shatavari	Asparagus racemosus Willd.	Tuberous	Jun 2014	Periphery of
			root		Jamnagar
10.	Punarnava	Boerhavia diffusa Linn	Root	July 2014	Periphery of
					Jamnagar

The loss on drying of any sample is directly related to its moisture content. If the moisture content is very high in any drug it may affect its preservation. The maximum loss on drying value was found in S5 (10.18 %w/w) followed by S3 (7.71% w/w) and S8 (5.38 %w/w).The ash value indicates the presence of inorganic and salt materials in the samples. The maximum ash value was

found in S3 (15.29% w/w) followed by S7 (15.15% w/w), S1 (13.12% w/w). Extract is a solid or semisolid preparation made by removing the soluble portion of a compound by using water or alcohol as the solvent and evaporating the solution or the active principle of a drug obtained by distillation or chemical processes<sup>10</sup>.

No	Sample	<b>Botanical Source</b>	Colour	Odour	Taste	Texture
<b>S</b> 1	Atibala	Abutilon indicum Linn.	Light yellow	Not specific	Sweet, Bitter	Fibrous,
		Sweet				coarse
S2	Vidari	Pueraria tuberosa DC.	Yellow	Sweet smell	Sweet	Fine
S3	Kantakari	Solanum xanthocarpum S.	Brownish	Sharp/	Bitter, Pungent	Moderately
		&W	yellow	Pungent		fine
				smell		
S4	Eranda	Ricinus communis Linn	Light brown	Not specific	Bitter, Sweet	Fibrous,
						coarse
S5	Gokshura	Tribulus terrestris Linn	Yellowish	Sharp smell	Sweet, Bitter	Moderately
			brown			fine
S6	Guduchi	Tinospora cordifolia	Yellowish	Bitter smell	Bitter	Moderately
		(Willd.) Miers ex Hook. f.	brown			fine
		&Thoms				
S7	Shalaparni	Desmodium gangeticum	Brownish	Woody	Astringent	Coarse
		DC.	yellow	smell		
S8	Jivanti	Leptadenia reticulata W. &	Dark green	Leafy smell	Bitter, Sweet,	Moderately
		А.			Astringent	fine
S9	Shatavari	Asparagus racemosus	Yellowish	Sweet smell	Bitter, Sweet	Smooth
		Willd.	brown			
S10	Punarnava	Boerhavia diffusa Linn	Brownish	Woody	Astringent	Coarse
			white	smell		

**Table 3** Microscopic characteristics powdered *Rasayana* drugs of *Madhuraskandha* (Plate 2 & 3)

No	<b>Botanical Source</b>	Part Used	Microscopic characters						
<b>S</b> 1	Abutilon indicum Linn. Sweet	Root	Simple and compound starch grains with hilum, Pitted, spiral vessels, Cluster crystal, Prismatic crystal,						
			Rhomboidal crystal, Pitted parenchyma cells, lignified						
			fibres, Yellowish content						
S2	Pueraria tuberosa DC.	Tuber	Simple starch grain with hilum, Compound starch grain						
			with hilum, Annular vessel, Yellowish content, Fibre,						
			Scaleriform vessel, Fragments of fibre, Acicular crystal,						
			Prismatic crystal, Parenchyma cells filled with starch grain						
<b>S</b> 3	Solanum xanthocarpum S.	Whole	Stellate, multi-branched and warty trichomes; Anisocyte						
	&W		stomata; Pitted, annular vessel, spiral vessels; Oil globule;						

		plant	Septate fibre; Epicarp cells of fruit; Black debris; Cork cells
S4	Ricinus communis Linn	Root	Bordered pitted vessels, Rosette, acicular, prismatic crystal,
			Septate fibre, fibre with wide lumen, Simple & compound
			starch grain with hilum, Cortex cell with rosette crystal,
			Brown content, Cork cell with tannin, Annular and spiral
			vessel
S5	Tribulus terrestris Linn	Fruit	Trichome, Pluricellulartrichome, Rossette crystal, Epicarp
			of fruit, Group of fibres, Mesocarp, Spiral vessel, Sclerides,
			Brown content, Epidermal cell, Stratified fibre, Prismatic
			crystal, Parenchyma cell with tannin
<b>S</b> 6	Tinospora cordifolia (Willd.)	Stem	Simple and compound irregular shaped starch grain with or
	Miers ex Hook. f. & Thoms		without hilum, Prismatic crystal
			Bordered pitted vessel , Yellow content; brown content,
			Collenchyma cells, Cork cells, Spiral vessels
<b>S</b> 7	Desmodium gangeticum DC.	Root	Cork cells, Prismatic crystal, Rhomboidal crystal, Brown
			content, Compound starch grain, Pitted vessel, Spiral vessel,
			Border pitted vessel, Epidermal cells with colouring matter,
			Group of simple starch grain without hilum, Oil globule,
			Crystal fibre, Pitted parenchyma
<b>S</b> 8	Leptadenia reticulata W. & A.	Leaf	Fragment of multicellular trichome, Epidermal cells with
			chlorophyll content, Fragments of Spiral vessel, Fragment
			of stomata along with epidermal cell, Fragment of
			multicellular warty trichomes, Paracytic stomata, Prismatic
			crystal, Brown content, Oil globule
<b>S</b> 9	Asparagus racemosus Willd.	Tuberous	Parenchymatous cell; Septate fibre; Acicular, Rod shaped
		root	microcrystals; Pitted parenchyma cells; Starch grain with
			concentric lines; Raphides, Simple starch grain; Filiferous
			hair; Scleriform vessels, Lignified fibres
S10	Boerhavia diffusa Linn	Root	Acicular crystal, Compound starch grain, Oil globule,
			Border pitted vessel, Brown content, Prismatic crystal, Cork
			cell in surface view, Parenchyma cells with starch grain,
			Bunch of Sclerides ,Stone cells

Therefore, water soluble extract indicates the total water soluble contents of the drug and methanol soluble extract indicates the total alcohol soluble contents of the drug. In all the test sample water soluble extractive value is more as compared to alcohol soluble extractive value, except in S3 where alcohol soluble extractive value is more than water soluble extractive value. It can be considered that the above drugs can be used in water based dosage forms like decoction, cold infusions, hot infusions; juice etc. pH of the drug determines acidity or alkalinity of drug<sup>11</sup>. Almost all the test drugs have pH below 7 indicating its acidic nature except S4 and S7 have pH value more than 7 which indicate their alkaline nature. All the test samples except S8 have shown the presence of Carbohydrates followed by reducing amino acids. proteins, sugar, tannin. steroids, triterpenoids, anthraquinone, and saponin.

### CONCLUSION

Madhuraskandha's Rasayana drugs may show the presence of starch grains and calcium oxalate crystals. Physicochemical analysis showed that soluble water extractive values of the all the drugs is more than alcohol soluble extractive values, which indicates more water soluble contents like carbohydrates, starch etc. Phytochemically, these drugs show the presence of carbohydrates, reducing sugar, amino acids, proteins, tannin, steroids, triterpenoids, anthraquinone, and saponin, which may be useful for their Rasayana activity. Pharmacognostical, Physico-chemical characters and phytochemical parameters may be useful to generate standards for quality and purity of Madhuraskadha's Rasayana drugs.

No	Botanical Source	Part Used	Loss on drying	Ash value	Water soluble extractive value	Alcohol soluble extractive value	рН
S1	Abutilon indicum Linn. Sweet	Root	4.9% w/w	13.12% w/w	6.858%w/w	4.226% w/w	6.40
S2	Pueraria tuberosa DC.	Tuber	2.29 %w/w	2.61% w/w	11.52%w/w	2.876%w/w	4.61
S3	Solanum xanthocarpu m S. &W	Whole plant	7.71% w/w	15.29% w/w	5.992%w/w	8.938% w/w	6.93
S4	<i>Ricinus communis</i> Linn	Root	4.13% w/w	7.19% w/w	5.185%w/w	4.876% w/w	7.25
S5	<i>Tribulus terrestris</i> Linn	Fruit	10.18 %w/w	8.68% w/w	7.213%w/w	6.284% w/w	6.47
S6	Tinospora cordifolia	Stem	4.79 %w/w	7.61% w/w	7.020%w/w	6.232% w/w	5.25

Table 4 Physicochemical characteristics of the Rasayana drugs of Madhuraskandha

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	(Willd.) Miers ex Hook. f. &Thoms						
S7	Desmodium gangeticum DC.	Root	1.3 %w/w	15.15% w/w	10.103% w/w	6.626% w/w	7.12
S8	Leptadenia reticulata W. & A.	Leaf	5.38 %w/w	12.46% w/w	29.91%w/w	10.15% w/w	6.22
S9	Asparagus racemosus Willd.	Tubero us root	0.89 %w/w	3.21% w/w	36.52%w/w	22.27% w/w	4.73
S10	<i>Boerhavia</i> diffusa Linn	Root	0.47% w/w	11.95% w/w	10.58%w/w	5.936% w/w	5.13

No	Phyto- constituents	Test performed	<b>S1</b>	S2	<b>S</b> 3	<b>S4</b>	<b>S</b> 5	<b>S6</b>	<b>S7</b>	<b>S8</b>	<b>S9</b>	S10
1.	Carbohydrate	Molish's test	+	+	+	+	+	+	+	-	+	+
2.	Reducing Sugar	Fehling's test	-	+	-	-	-	+	-	-	++	+
3.	Hexose sugar	Selwinoff's test	-	-	+	-	-	-	-	Т	+	-
4.	Amino acids	Ninhydrin test	-	+	-	-	+	+	+	+	+	+
5.	Alkaloid	Dragondroff's	-	-	-	-	-	-	-	-	-	-
		test										
		Hegar's test										
6.	Starch	Iodine test	-	-	-	-	-	-	-	+	-	-
7.	Protein	<b>Biuret's test</b>	-	-	-	+	+	+	-	-	-	-
		Protein										
		containing	-	-	-	-	-	-		-	-	-
		sulphur										
8.	Tannin	Lead acetate test	+	+	-	-	-	-	-	-	-	-
		FeCl <sub>3</sub> test										
					+	+		+	+			
9.	Steroid	Salkowski test	+	-	+	+	-	+	+	-	-	+
10.	Flavonoids	Zinc HCl test			+	+			+			
		Shinoda test										
		VanilineHCl test										
		Lead acetate										
11.	Cardiac glycoside	Legal's test	+	+	+	+	+	+	+			
		Baljet test				+						

 Table 5 Preliminary phytochemical characteristics of the Rasayana drugs of Madhuraskandha

12.	Anthraquinone	Borntreger's test	-	-	-	+	-	+	+	-	-	+
13.	Saponin	Foam Test	•	-	-	-	+	-	-	-	+	-

[T: Trace; +: Present; - : Absent]



Fig 1 Abutilon indicum Linn. Sweet



Fig 3 Solanum xanthocarpum S. &W



Fig 5 Tribulus terestris Linn



Fig 2 Pueraria tuberosa DC



Fig 4 Ricinus communis Linn





Fig 7 Desmodium gangeticum DC



Fig 9 Asparagus racemosus Willd



Fig 8 Leptadenia reticulata W. & A.



Fig 10 Boerhavia diffusa Linn

Plate no.1 Madhuraskandha Rasayana drugs in their natural habitat

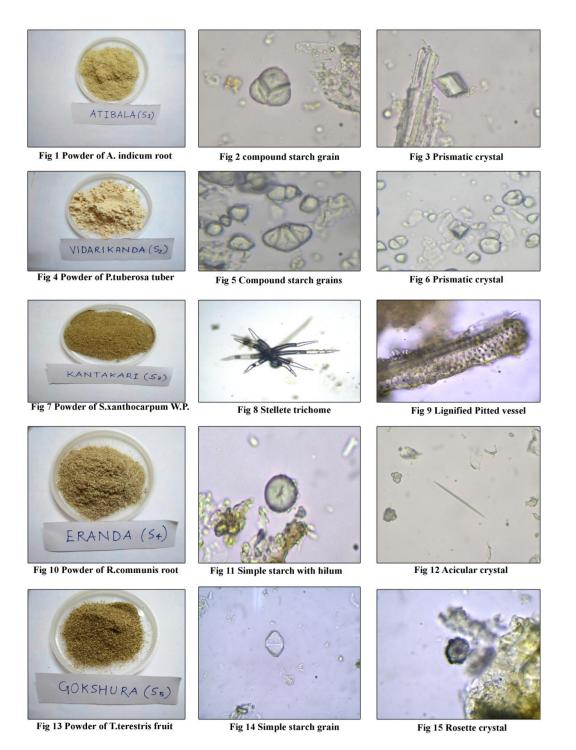


Plate no.2 Powder microscopical characters of Madhuraskandha's Rasayana drugs (S1-S5)

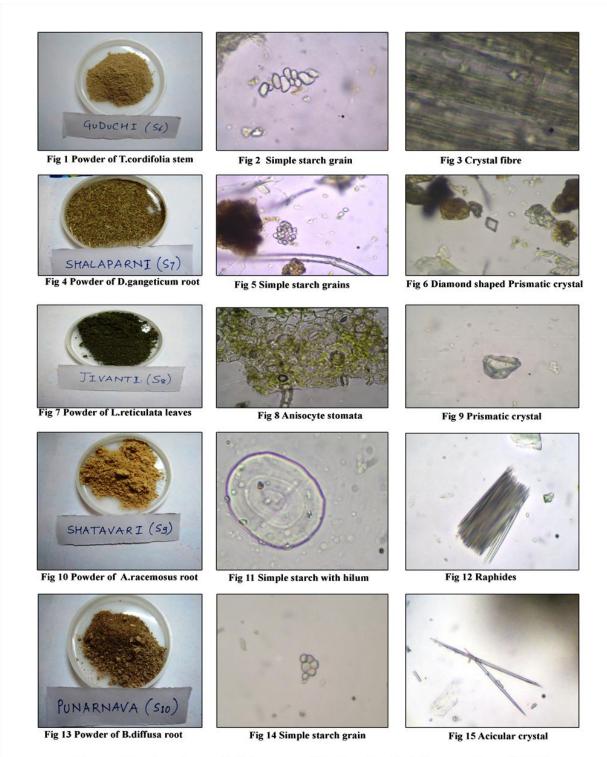


Plate no.3 Powder microscopical characters of Madhuraskandha's Rasayana drugs (S6-S10)

## REFERENCES

Bhuvad and Nishteswar Int J Ayu Pharm Chem 2015 Vol. 3 Issue 1 [e ISSN 2350-0204]

- Yadavaji Trikamaji Acharya, editor, Cakrapanidatta, commentator (2009).Carakasamhita, Vimanasthana 8/138, 1st reprint edition, Varanasi: ChaukhambhaSurbharatiPrakashana;283
- Yadavaji Trikamaji Acharya, editor, Cakrapanidatta, commentator (2009). Carakasamhita, Sutrasthana 26/42,61, 1st reprint edition, Varanasi: Chaukhambha Surbharati Prakashana; 144,146
- Anonymous (2003).Ayurvedic Formulary of India, Part I, Vol I, 2nd edition, New Delhi: Govt. of India publication, Ministry of Health & FW, Dept. of ISM and H;302
- Yadavaji Trikamaji Acharya, editor, Cakrapanidatta, commentator (2009) Carakasamhita, Sutrasthana 4/18, 1st reprint edition, Varanasi: Chaukhambha Surbharati Prakashana;34
- Anonymous (1999) The Ayurvedic Pharmacopoeia of India, Part I, Vol. I, Appendix 2, 1st edition, New Delhi, Govt. of India publication, Ministry of Health& FW, Dept. of ISM and H; 207
- Krishnamurthy K V (1988). Method in the Plant Histochemistry. Madras: Vishwanandan Pvt. Ltd; 1-77
- Anonymous (1999) The Ayurvedic Pharmacopoeia of India, Part-I, Vol. 1-4, New Delhi , Govt. of India, Ministry of

Health & FW, Dept. of ISM and H; 213-14

- Shukla V.J., Bhatt U.B. (2001). Methods of Qualitative Testing of some Ayurvedic Formulations. Jamnagar, Gujarat Ayurved University; 5-10.
- K. R. Khandelwal (2008). Practical Pharmacognosy-techniques and experiments, 19th Edition, Pune, Nirali Prakashan;149-156
- Donald Venes, Editor, Clayton L. (Eds.) Thomas, Clayton L. Thomas, Editor (2005). Taber cyclopaedic medical dictionary, 20th edition, Philadelphia, PA, F. A. Davis Company; 762.
- 11. Donald Venes (Editor), Clayton L.
  (Eds.) Thomas, Clayton L. Thomas, Editor (2005). Taber cyclopaedic medical dictionary, 20th edition, Philadelphia, PA, F. A. Davis Company; 1647.