Development of Standards of Sapistan (*Cordia dichotoma* Forst.f.)

*Abdul Haleem, Abdul Latif, Abdur Rauf, Nazish Siddiqui and Sumbul Rehman

Department of Ilmul Advia, A.K. Tibbiya College, Aligarh Muslim University, Aligarh-202002

Abstract

hysicochemical and Phytochemical standardization is considered a pre-requisite for the assessment of biological activity or determination of biological standards of the plant material. It provides the analytical characteristics which may prove to be useful in fixing the physicochemical standard for the Unani herbal drugs.

Sapistan (*Cordia dichotoma*) belongs to the family Ehretiaceae; its fruits and leaves are used in respiratory diseases such as asthma, cough, diphtheria, pneumonia, sore-throat, coryza. An effort has been made to carry out the physicochemical and phytochemical studies of plant. Physicochemical parameters as Extractive Values: Petroleum Ether (1.9%), Di-ethyl ether (0.85%), Chloroform (0.59%), Acetone (0.12%), Alcoholic (6.13%), Aqueous (10.44%); Solubility: Water (9.44%) & Alcohol (1.16%); Moisture contents (3.45%), Total Ash values (7.188%), pH of 1% (6.76) & 10% solution (6.16) and loss on drying (5.3%). Phytochemical analysis: These revealed the presence of almost all the phyto-constituents in the test drug sample i.e. alkaloid, flavonoids, glycoside, carbohydrate, tannin, protein, amino acids, starch and resins.

Keywords: Sapistan, Physicochemical, Phytochemical, Standardization, *Cordia dichotoma* Forst.f.

Introduction

Sapistan (*Cordia dichotoma* Forst.f.) belongs to the family Ehretiaceae (Anonymous, 2006). Use of the drug 'Sapistan' in Unani system of medicine dates back about a couple of thousands of years. It is mentioned by Theophrastus and was not disregarded through the Arabic, Persian and Urdu authors in their books. It was particularly mentioned by Razi (926 A.D.) Ibne Sina (1037 A.D.) Al- Harwi (10th C.A.D). Sebestan or Sapistan is an abbreviation of sag pistan, derived from sag, adog and pistan dugs. The fruit resembles bitches dugs in shape. Sapistan is used for cure of the respiratory diseases (Khory and Katrak, 1984).

Sapistan is a fruit of tree. The tree is of two types, one type of tree have a large fruits known as 'Rai Gond'. And the other tree having small fruit called 'Kath Gond'. This tree consists of many branches which are pale red in colour. Flowers are found in clusters form. Fruit get ripened in the month of May to July (Ghani, 2011).

Medicinally, the dried fruit is valued on account of its mucilaginous nature and demulcent properties; it is much used in coughs and chest affections, also in

*Author for correspondence



irritation of the urinary passages; in larger quantities it is given in bilious affections as a laxative. Mahometan writers describe two kinds of Sapistan; the greater, the pulp of which is separable from the stone, and the lesser, the pulp of which is adherent.

Botany- brief description:

A middle-sized deciduous tree and usually with a crooked trunk. Bark grey, rough with shallow longitudinal furrows. Leaves thinly coriaceous, variable in shape and size, orbicular, ovate-elliptic. Flowers white, fragrant. Drupes ovoid, apiculate, pink or orange to reddish when ripe, filled with a viscid pulp; stone usually 1. Common in deciduous forests, flowers – March and April; fruits – May to July.

Therapeutic action and uses :

Fruits-astringent, anthelmintic, diuretic, demulcent, expectorant. Used in affection of urinary passages, diseases of the lung and spleen. Bark – used in dyspepsia and fever. Pulp – used in ringworm. Leaves – used in ulcer, headache (Ambasta, 1986; Chopra *et al.*, 1956; Kirtikar and Basu, 1996; Nadkarni, 1989).

No work appears to have been reported regarding standardization of this drug so far. Keeping in mind the medicinal importance of this plant in Indian Systems of Medicine, a physico-chemical and phytochemical study of this drug was carried out in order to fix-up its pharmacopoeial standards.

Material and Method

Collection of plant material: The drug samples of Sapistan (fruits) were collected from Bara Duwari market of Aligarh city and were identified. Voucher specimens were preserved in the herbarium of Medicinal Plants Lab in the Department of Ilmul Advia, F/O Unani Medicine, Aligarh Muslim University, Aligarh (Voucher No. SC-0141/14).

Chemical parameters: First, the organoleptic characters were studied. The dried powder of the fruits of Sapistan was used for chemical analysis. Various physicochemical studies like total ash, acid insoluble ash, water soluble ash, alcohol and water soluble matter, moisture content, successive extractive values using soxhlet extraction method, bulk density and pH studies were carried out as per guidelines of WHO (Anonymous, 1998, 2008). Qualitative analysis of the drug was conducted to identify the organic chemical constituents present in the drug (Overtone, 1963; Harborne, 1973).

The thin layer chromatographic analysis was conducted following Stahl (1969) and Harborne (1973) on precoated silica gel $60F_{254}$ TLC plates. The plates were



visualized in day light, in short UV and Long UV. They were also derivatised using iodine vapors.

Observations

- (a) Organoleptic characters: The powder of the fruits of Sapistan was brown with fruity smell any characteristic odour (Table 1).
- (b) Physico-chemical constants: The analytical values of different physicochemical constants were determined (Table 2).

S.No.	Organoleptic parameters	Observations
1.	Colour	Brwon
2.	Smell	Fruity
3.	Taste	Disagreeable

Table 1: Organoleptic Characters of Cordia dichotoma Forst.f.

Table 2: Physicochemical study of Powder of Sapistan

S.No.	Parameters	Percentage (w/w)*
1	Ash value	
	Total ash	7.188
	Acid insoluble ash	1.22
	Water soluble ash	5.92
2	Soluble Part	
	Ethanol soluble	1.66
	Aqueous soluble	9.44
3	Successive Extractive Values	
	Pet. Ether	1.72
	Di-ethyl ether	0.85
	Chloroform	0.59
	Acetone	0.12
	Alcohol	6.13
	Aqueous	10.44
4	Moisture content	3.45
5	Loss on Drying	5.3
6	pH values	
	1% water solution	6.76
	10% water solution	6.16
7	Bulk density	0.67

*Note: Values are average of three experiments.



- (c) Qualitative analysis of organic chemical constituents of drug: The phytochemicals present in the drug were identified on the basis of different chemical tests given for various plant constituents (Table 3).
- (d) FTAR Analysis: Fluorescence analysis of the successive extract was studied under day light as well as Ultra Violet (short and long wave length) light, results have been summarized in Table-4. FTAR Analysis was also done of the powdered drug after reacting them with various chemical reagents (Table 4).

S.No.	Chemical Constituent	Tests/Reagent	Inference
1	Alkaloids	Dragendorff's reagent	+
		Wagner's reagent	+
		Mayer's reagent	+
2	Carbohydrate	Molisch's Test	+
		Fehling's Test	+
		Benedict Test	+
3	Flavonoids	Mg ribbon and Dil.Hcl	+
4	Glycosides	NaOH Test	+
5	Tannins/Phenols	Ferric Chloride Test	
		Liebermann's Test	+
		Lead Acetate Test	+
6	Proteins	Xanthoproteic Test +	
		Biuret Test	+
7	Starch	Iodine Test	
8	Saponins	Frothing with NaHCO ₃	+
9	Steroid/Terpenes	Salkowski Reaction +	
10	Amino Acids	Ninhydrin Solution –	
11	Resin	Acetic Anhydride test	+

Table 3: Preliminary Screening of major Phyto-chemicals

Indications: '-' Absence and '+' presence of constituent.

Table 4: FTAR	Analysis	of	Sapistan
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S.No.	Extract	Day Light	UV Long	UV Short
1.	Pet. Ether	Brown	Black	Greenish
2.	Di-ethyl ether	Greenish	Bluish	Greenish
3.	Chloroform	Light Green	Black	Light Green
4.	Acetone	Brown	Bluish	Greenish
5.	Alcohol	Brown	Black	Dark Brown
6.	Aqueous	Brown	Black	Black



(e) Thin layer chromatographic profile: Thin layer chromatographic analysis of successive extracts was carried out using different solvent systems and visualizing agents and R_f values were calculated to standardize the drug for its identity and purity (Table 5).

S.	Powdered drug +	Day light	UV short	UV long
No.	Chemical Reagent			
1.	Powdered drug + Conc. HNO ₃	Golden	Light Green	Green
2.	Powdered drug + Conc. Hcl	Brown	Light Green	Dark Green
3.	Powdered drug + Conc. H ₂ SO ₄	Dark Brown	Dark Green	Black
4.	Powdered drug + 2% lodine solution	Brown	Green	Black
5.	Powdered drug + Glacial Acetic acid + HNO ₃	Light Brown	Light Green	Dark Green
6.	Powdered drug + Glacial acetic acid	Pale	Light Green	Black
7.	Powdered drug + NaOH(10%)	Brown	Light Green	Dark Green
8.	Powdered drug + Dil. HNO ₃	Brown	Green	Green
9.	Powdered drug + Dil. H ₂ SO ₄	Brown	Green	Dark Green
10.	Powdered drug +Dil. Hcl	Light Brown	Light Green	Black
11.	Powdered drug + Dragendorff's	Dark Brown	Bright Green	Green
12.	Powdered drug + Wagner's Reagent	Brown	Green	Dark Green
13	Powdered drug + Benedict's reagent	Blackish	Light Green	Black
14	Powdered drug + Fehling reagent	Brown	Light Green	Green
15	Powdered drug + KOH (10%) Methanolic	Brown	Green	Black
16	Powdered drug + CuSO ₄ (5%)	Brown	Light Green	Black
17	Powdered drug + Ninhydrin (2%) in Acetone	Brown	Green	Black
18	Powdered drug + Picric Acid	Yellow	Bright Green	Black
19	Powdered drug + Lead Acetate (5%)	Brown	Light Green	Black

Table 5: Fluorescence Analysis of Sapistan with different chemical reagents



Treatment	Mobile phase:	No of spots	Rf value and colour of spots	
Petroleum Ether Extract				
Day Light	Petroleum ether :	1	0.42 (Brown)	
UV Short	Di-ethyl ether (4:1)	1	0.42 (Bluish)	
Iodine Vapour]	1	0.42 (Yellowish)	
	Aqueo	us Extract		
Day Light	Butanol: Acetic	1	0.75 (Brown)	
UV Short	acid: Water (5:1:4)	3	0.64 (Bluish), 0.78 (White),	
			0.85 (White)	
UV Long		4	0.21 (white), 0.42 (white),	
			0.78 (white), 0.85 (white)	
Iodine Vapour		1	0.75 (Pale Yellow)	
Alcoholic Extract				
Day Light	Chloroform:	1	0.62 (Pale yellow)	
UV Short	Methanol (9:1)	2	0.62 (Purple) , 0.75 (Purple)	
UV Long		2	0.62 (white), 0.75 (White)	
Iodine Vapour		1	0.62 (Pale Yellow)	

Table 6: Thin Layer Chromatography

Results and Discussion

Physico-chemical standardization is of prime importance in quality control of Unani drugs. As the efficacy of many drugs mainly depends upon its physical and chemical properties, therefore, the determination of physico-chemical characters for the authenticity of a drug is necessary before studying any medicinal property. Phyto-chemical constituents present in the drug vary, not only from plant to plant but also among different samples of same species, depending upon various atmospheric factors, storage and drying conditions.

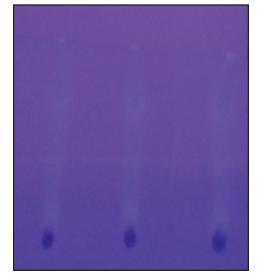
The present study is an attempt to fix-up pharmacopoeial standards of drug 'Sapistan' to ascertain its quality, identity, purity and strength. Powder of the drug has been used for study to bring out several standards like ash, solubility in alcohol and water, successive, extractive values and quality screening of physicochemicals, total alkaloids, total flavonoids, phenol, nitrogen, fatty matter, sterol/ terpenes, protein and carbohydrates.

Based on the various physico-chemical and phytochemical parameters, the drug 'Sapistan' has been standardized to ensure its use in manufacturing quality and genuine herbal preparations.





Sapistan Fruits (Cordia dichotoma Forst.f.)





TLC of Alcoholic Extract of Sapistan



References

Afaq, S.H., Tajuddin, Siddiqui, M.M.H., 1994. Standardization of Herbal Drugs. Publication Division, AMU (Aligarh), pp. 33-34, 41-42, 100, 143-146.

Ambasta, S.P., 1986. The Useful Plants of India. PID, CSIR, New Delhi.

Anonymous, 1997. Standardization of Single Drugs of Unani Medicine, Part III. CCRUM New Delhi, pp. 124-128.

Anonymous, 1998. Quality control methods for medicinal plant materials. World Health Organization, Geneva, pp. 25 -28.



- Anonymous, 2006. National Formulary of Unani Medicine, Part I. CCRUM, Dept. of AYUSH, New Delhi, p. 272.
- Anonymous, 2008. Quality control manual for Ayurveda, Siddha and Unani medicine. Govt. of India, Dept. of AYUSH, New Delhi, pp. 21 29.
- Anonymous, 2008. Unani Medicinal Plants of Tarai Forests in Kumaon Region of Uttarakhand. Central Council for Research in Unani Medicine, p. 60.
- Baitar I., 1999. Al-jamiul-mufridat-ul-advia-wa-ul-aghzia. Urdu Translation. Part.III. CCRUM, New Delhi, pp. 23-24
- Chopra, R.N., Nayar, S.L. and Chopra, I.C., 1956. Glossary of Indian Medicinal Plants, CSIR, New Delhi.
- Dymock, 1891. Pharmacographica Indica, Part II. Published by Institute of Health and Tibbi Research, Hamdard National Foundation, Pakistan, pp. 518-519.
- Ghani Najmul, 2011. Khazainul Advia. Idara Kitabul Shifa, New Delhi, pp. 787-788.
- Harborne, J. B., 1973. Phytochemical methods. Chapman and Hall. London, p. 70.
- Ibn-e-Sina, 2007. Al-Qanoon-fi'l-tib, Vol. II. Urdu translation by Ghulam Hussain Kantoori. Idara Kitabul Shifa, New Delhi, p. 167.
- Jenkins, G.L., Knevel, A.M. and Digangi, F.E., 1967. Quantitative Pharmaceutical Chemistry. 6th edition. The Blackiston Division. McGraw Hill Book Company, U.S.A., pp. 225, 235, 379, 425, 463, 492.
- Khory, R.N. and Katrak, N.N., 1984. Materia Medica of India and their Therapeutics. Neeraj Publishing House, Delhi, p. 421.
- Kiritikar, K.R. and Basu, B.D., 1996. Indian Medicinal Plants, Vol. III. International Book Distributers (Dehradun), pp. 1674-1679.
- Nadkarni, K.M., 1989. The Indian Materia Medica, Vol. II. Bombay Prakashan Pvt. Ltd., Bombay, pp. 379-380.
- Overtone, K.H., 1963. Isolation, Purification and preliminary observation in elucidation of structures by physical and chemical methods. Bentley Interscience Pub., New York, p. 34.

Stahl, 1969. Thin Layer Chromatography: A laboratory handbook, Springer Verlag student edition. Springer Verlag, Berlin, pp. 52 – 86, 127 – 128, 900.



