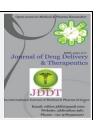


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Research Article

PHYSICO-CHEMICAL STANDARDIZATION OF A UNANI ANALGESIC FORMULATION *HABB-E-SURANJAN*

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ABSTRACT

Habb-e-Suranjaan is an important compound preparation which is used for a centuries in Unani system of medicine as an effective analgesic for all types of Waj-al-Mafasil (Arthritics). The drug is known for its pharmacological actions such as Mushile-Balgham (Phlegmagogue), Mushile-Safra (Cholagogue), Mushile-Sauda (Melanagogue), Mohallile-Waram (Anti-inflammatory), Musakkin-e-Alam (Analgesic), Dafae Niqris (Anti-gout) Muqawwi-e asaab (Nervine tonic) Munawwim (Hypnotic/Soporific), Musaffi-e-Dam (Blood Purifier). Ingredients of Habb-e-Suranjaan play very effective role in management of different types of Waj-al-Mafasil. In this article, standardized value of a specimen of Habb-e Suranjaan assessed on physico-chemical and analytical parameters viz Macroscopic and microscopic features, extractive values, moisture contents, Ash values, loss of weight on drying, pH of 1 % and 10 % solution, TLC and Fluorescence analysis are provided. The moisture content and the ash value were found within the recommended normal range. The value of different hot extracts (Petroleum ether, Methanol, chloroform and Aqueous) is more than the cold extracts of drug.

Keywords- Habb-e-Suranjaan, Waj-al-Mafasil, Analgesic.

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INTRODUCTION

Habb(pill) is an ancient dosage form of Unani system of medicine invented by HakeemSeelon¹. Habb is an Arabic word which means pill but it is used in other meanings also. The constituents of pill may be either one or more than one. It may be prepared either with drugs of the all three origins or two of them or even only one of them. Volume or size of pill may varies, some are very small like size of Sarson (Bressicanigra) seed, Masoor (Lens culinaris)seed, Baajra (Pennissetumglaucum), Matar (Pisumsatvum Linn.) or about to one cm or more. Pills of one cm diameter are called Bunduga (Plural: Banadig). It is generally similar to size of Reetha fruit (Sapindustrifoliatus)^{2, 3}. Habb-e-Suranjaan is an important compound preparation which is used for a long timeinUnani system of medicine as a good analgesic. The compound *Habb-e-Suranjan*is a Pharmacopoeial formulation mentioned in National Formulary of Unani Medicine and Qarabadeene Majeedi. The ingredients and composition of Habb-e-Suranjan are as follows4,5-

Ingredients	Botanical name	Qt.
Suranjan Shireen	Colchicum autumnal	80g
Turbud	Ipomeaturpethum	95g
Aeloa/ Sibr	Aloe barbadensis	35g
Habb al-Neel	Ipomeahederacea	35g
Gugul / Muqil	Commiphoramukul	15g
Mastagi	Pistacialintiscus	15g
Tukhm soya /Shibt	Anethumsowa	35g

Analgesic property of ingredients of are mentioned in Unani literature in abundance and therapeutic use of compound formulation Habb-e Suranjaan are Waja'-al-Mafasil (Arthralgia), Irq al-Nasa (Sciatica) Niqris(Gout), Falij (Paraplegia), Laqwa (Bell's palsy) as per Unani scholars.6,7,8,9,10

MATERIALS AND METHODS

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The drug was purchased from the local market at Khari Baoli, New Delhi, India. The *Haboob*was prepared in Saidla Lab, Dept. of Ilmul Advia, School of Unani Medicine, Jamia Hamdard.

Macroscopic and Microscopic features: were recorded.

Morphological Characters

Detailed study of the morphological character can be helpful in differentiating them from the other habb (pills). The macroscopy of a habb (pills) includes its visual appearance to the naked eye. For each particular morphological group, a particular systemic examination can be carried out. Size, Weight of empty petridish = w1 gm

Weight of petridish + test drug = w2 gm

Weight of petridish + Dried drug = w3 gm

Weight of Test drug = (w2-w1) gm

Weight of moisture = (w2-w3) gm

% age of moisture w2-w3X 100 w2-w1

Determination of pH

pH of 1% solution

Five gram powder of drug was shaken well and suspended in 100ml of distilled water. The resulting solution/ mixture were filtered and pH was measured with a standard glass electrode. Result is summarized in table.

pH of 10% solution

Acid Insoluble Ash Content

HCl (2 N; 25 mL) was added to the crucible containing the total ash, covered with a watch glass, and boiled gently for 5 min. The watch glass was rinsed with 5 mL of hot water and the rinsed contents were added to the crucible. The acid insoluble matter was collected on an ash less filter paper and washed with hot water until the filtrate was neutral. The filter paper containing acid insoluble matter was transferred to the original crucible. The content of crucible incinerated in the furnace at 450 oC. The residue was allowed to cool in a desiccator and weighed. The content of the acid insoluble ash (in mg/g) of air-dried material was calculated as follows:

Percentage of acid insoluble ash = WHCl / WDry×100

WHCl = Wt. of HCl insoluble ash

WDry = Weight of the dried sample

Water soluble ash

Water (25 mL) was added to the crucible containing the total ash, covered with a watch glass and boiled gently for 5 min. The watch glass was rinsed with 5 ml of hot water and added to the crucible. The water insoluble matter was collected on an ash less filter paper and washed with hot water. The filter paper containing the water insoluble matter was transferred to the original crucible and incinerated in the muffle furnace at 450° C.

Percentage of water soluble ash = $W_{H20} / W_{Dry} \times 100$

 $W_{\rm H20}$ = Wt. of water soluble ash = Total ash wt. – wt. of water insoluble matter

Hot Extraction of Habb-e-Suranjaan

50 g of coarse powder of the *Habb-e- Suranjaan* was subjected to extraction with different solvent system like petroleum ether, then chloroform, then ethanol and at last in water through soxhlet apparatus for 6 hr. The liquid extracts

colour, odour and test are important parts of morphology of a particular habb (pills).

Moisture content (Mc)

The drug sample (2g) was taken in a clean petridish of known weight (w1) and the petridish was weighted along with sample drug (w2). The drug containing Petridish was kept in an oven at 100-1050C for 2hrs and then the weight (w3) was noted. Then again, the drug with petridish was kept in an oven at 1050C for next 2 hrs. The process was repeated until constant weight was obtained. The loss of weight (inmg/g) of air dried was calculated as follows:

The experimentation was performed in the same manner as above taking 10gof drug instead of 5g.

Ash value

The powdered material (5g) was accurately weighed and placed in a crucible. The material was spread in an even layer and it was ignited to a constant weight by gradually increasing the heat to $450\,^{\circ}\text{C}$ until it was white indicating the absence of carbon. The residual ash was allowed to cool in a desiccator. The content of total ash(in mg/g) of air-dried material was calculated as follows:

Total Ash value

% Ash (dry basis) = $W_{Ash} / W_{Dry} \times 100$

W_{Ash}= Weight of the ashed sample

W_{Dry} = Weight of the dried sample

Percentage of total ash = W_{Ash} / $W_{Dry} \times 100$ thus obtained were put over water bath and evaporated to dryness. The dried extracts were afterward kept for 5 min in hot oven and their constant successive extractive values were recorded.

Reaction of Habb-e-Suranjan with different reagents

The test drug was treated with various reagents and color in the test tube was noted.

Thin Layer Chromatography (TLC Profile)

Chloroform (CHCl3) extract of *Habb-e- Suranjan* was used for TLC. Chloroform extract was prepared by dissolving 50g of *Habb-e- Suranjan* in 100 ml of Chloroform. Drug solution thereafter filtered and filtrate was concentrated under reduced pressure.

Mobile phase development: Initially, various combination were tried such as chloroform: petroleum ether: ethyl acetate (2:2:1), and other solvents in various ratio. Finally upper layer of chloroform: petroleum ether (8:2) was used as mobile phase for Chloroform extract of *Habb-e-Suranjaan*.

RESULTS

Table-1 Findings of Morphological Characters

S. No.	Parameters	Observation	
1.	Size	Round	
2.	Color	Light olive brown	
3.	Odour	Agreeable	
4.	Taste	Astringent.	
5.	Appearance	Habb(pills)	

Table-2 Findings of Physio-chemical Standardization

S. No.	Parameters	(%) mean
1	Loss on drying	2.6%
1.	Loss on arying	2.6%
2.	Swelling Index	1.1±0.10%
3.	Ash values	
	Total ash (%)	6.4%
	Acid insoluble ash (%)	5.2%
	Water soluble ash (%)	1.6%
4.	рН	
	pH 1% solution	6.54
	pH 10% solution	5.75
5.	Hot extraction	
	Petroleum ether	4.24%
	Chloroform	2.04%
	Methanol	5.6%
1.	Aqueous	28.14%
6.	Cold Extraction	
	Petroleum ether	2.5%
	Chloroform	2.40%
	Methanol	2.90%
	Aqueous	22%
7.	Successive Extract Value	
	Petroleum ether	4.15%
	Chloroform	3.55%
	Methanol	4.98%
	Aqueous	12.05%

 ${\bf Table \hbox{-} 3 \ Fluorescence analysis of} \ \textit{Habb-e-Suranjan}$

S. No.	Reagents	Ob	Observation	
		Day light	UV light	
			(254 nm)	
1.	Powder as such	Light brown	Darkish brown	
2.	Powder treated with dist. water	Light brown	Dark brown	
3.	Powder treated with Ferric chloride	Dark brown	Blackish brown	
4.	Powder treated with Pet. ether	Straw Colour	Yellowish Colour	
5.	Powder treated with sodium hydroxide (NaOH)	Light orange	Brown	
6.	Powder treated with chloroform (CHCl3)	Lemon	Greenish yellow	
7.	Powder treated with methanol	Clear	Clear	
8.	Powder treated with ethyl acetate	Hazy brown	Yellowish	
9.	Powder treated with acetone	Straw Colour	Greenish yellow	
10.	Powder treated with hydrochloric acid (HCl)	Dark brown	Blackish brown	
11.	Powder treated with sulphuric acid	Brick colour	Black colour	
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DISCUSSION AND CONCLUSION

Morphological characterization of a drug helps in identification of drug as well as detection of adulteration. In some cases, quality of crude drug can be checked on the basis of morphology only. In the present study, it has been observed that test drug had moisture content less than 10%. Low moisture content is always desirable for stability of drug. ¹¹ High ash values of 6% suggest presence of inorganic matter. Lower value of the acid insoluble ash suggests the greater physiological availability of drug. Extractive value gives information about availability of soluble phytoconstituents in particular solvent. ¹² Water soluble extractive is more as compared to ethanol, petroleum ether and chloroform extractive value suggesting that aqueous extract

would be more beneficial as compared to methanolic extract for therapeutic purposes.

In the light of above finding it may be concluded that test drug *Habb-e- Suranjaan* is found in good quality on physic-chemical parameters as per gold standard.

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CONFLICT OF INTEREST: None declared.

SOURCE OF SUPPORT: Nil

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