RESEARCH ARTICLE

Physicochemical Analysis of Simhanada Guggulu Pill

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Abstract

It is being globally recognized that medicinal plants play an important role for providing health benefits to human beings. Maximum Ayurvedic drugs are plant based drugs. The complex composition of plant based drugs has a big challenge for quality control. Physicochemical analysis is the most important part for standardization of the plant base drugs. One most important Ayurvedic drug i.e. *Simhanad Guggulu* pill had been selected from Ayurvedic famous book named *Bhaishajya Ratnavali* for the Physicochemical analysis. It is mainly and commonly used in the treatment of disease *Amavata* (Rheumatoid arthritis). Drug preparing and Physicochemical analysis both had been done in the Institute for Post Graduate Teaching and Research in Ayurveda, Gujarat Ayurved University, Jamnagar. The Physicochemical study revealed that the *Simhanad Guggulu* pill contained sterol, less moisture, more inorganic constituents and less water soluble constituents.

Keywords Simhanad Guggulu, pill, Physicochemical, Amavata, Rheumatoid arthritis



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INTRODUCTION

It is being recognized globally that the medicinal plants play an important role for providing health benefits to human beings. The gradual increased demand for plant based their drugs and eventual commercialization has given a more concentration on their status. Maximum Ayurvedic drugs are plant based drugs. But global acceptances of Indian plant based drugs are still low and perhaps inadequacy of quality control is the most important responsible factor for this. The complex composition of plant based drugs is a big challenge for quality control. These days Physicochemical analysis is the most important way for standardization of the plant based drugs. Many plant based drugs are described in Ayurvedic texts in context of treatment purpose of different diseases. One most important Ayurvedic drugs i.e. Simhanad Guggulu pill had been selected from Ayurvedic book for the Physicochemical study.

Objectives:ToanalysisthePhysicochemicaldataoftheSimhanadGuggulupill.

MATERIALAS AND METHODS

Simhanad Guggulu pill is mainly and commonly used by the Ayurvedic physician's in the treatment of disease Amavata (Rheumatoid arthritis). Amavata disease is more simulated to Rheumatoid arthritis according to its clinical features and pathogenesis ^{1, 2}. Simhanad Guggulu pill is mentioned in *slokas* no. 190 to 195 of 29th chapter of Bhaishajya Ratnavali (Ayurvedic book)³. Simhanad Guggulu pill was prepared and its physicochemical study of was carried out in the Pharmacy of Institute for Post Graduate Teaching and Research in Ayurveda, Gujarat Ayurved University, Jamnagar the Pharmaceutical Laboratory of Institute for Post Graduate Teaching and Research in Ayurveda, Gujarat Ayurved University, Jamnagar. Uniformity of tablet/pill (weight variation) (average weight), Hardness of tablet/ pill (average), of Disintegration time tablet/pill, Determination of Loss on drying at 110 °C, Ash value (% of total ash), Acid insoluble ash value, Water soluble extractive value and Methanol soluble extractive value of Simhanad Guggulu pill had been observed for Physicochemical analysis^{4,5}. Simhanad Guggulu pill is a herbo-mineral Ayurvedic drug and six ingredients are used in it. Out of six ingredients five are herbal ingredients

and one is mineral ingredient^{6,7}. Name of the ingredients (Ayurvedic and Scientific name), used part of the ingredients and

quantity of used part of the ingredients into the one pill are shown in the **Table 1**.

S. No.	Ingredients	Scientific or Botanical	Used part	Quantity
	(Ayurvedic	name		(part)
	name)			
1.	Haritaki	Terminalia chebula Retz.	Dried mature Fruit	1
2.	Amalaki	Emblica officinalis Gaertn.	Dried mature Fruit	1
3.	Bibhitaka	Terminalia bellirica Roxb.	Dried mature Fruit	1
4.	Guggulu	Commiphora wightii	Gum exudates	1
	(Shodhita)	(Arnott)		
		Bhandari		
5.	Gandhak	Sulphar	Mineral	1
	(Shodhita)			
6.	Eranda taila	Ricinus communis Linn.	Seed oil	4

RESULTS AND DISCUSSION

Results of Physicochemical analysis is shown in the **Table 2**.

Table	2	Data	of	Physicochemical	parameters		
(Quantitative test) of Simhanad Guggulu pill							

S.	Parameter	Result
No.		
1.	Uniformity of tablet (weight	502.5 mg
	variation) (average weight)	
2.	Hardness of tablet (average)	1.55 kg./cm ²
3.	Disintegration time of tablet	more than 1
		hour
4.	Determination of Loss on	2.30 % W/W
	drying at 110 °C	
5.	Ash value (% of total ash)	10.00 % W/W
6.	Acid insoluble ash value	2.40 % W/W
7.	Water soluble extractive	28.10 % W/W
	value	
8.	Methanol soluble extractive	22.20 % W/W
	value	

The data of the Table 2 shows that the average weight of Simhanada Guggulu pill was 502.5mg, Hardness of the Simhanada cm^2 , Guggulu pill was 1.55 kg/ Disintegration time of this pill was more than 1 hour, Loss on drying of this pill sample at110 °C was 2.3 % W/W, and Ash value, Acid insoluble ash value, Water soluble extractive value and Methanol soluble extractive value of this pill sample were observed respectively 10% W/W, 2.40 % W/W., 28.1 % W/W., and 22.2 % W/W. On the basis of this information it can be said that the moisture holding capacity was

less in the sample of Simhanada Guggulu pill and hence there may be minimum chance of damage of the pill by moisture, so shelf life or storage capacity is not less in Guggulu Simhanada pill. Inorganic constituents were more in Simhanada Guggulu pill, because it was made by herbomineral ingredients, therefore Ash value was more. Water soluble constituents such as Sugars, Glycosides etc were less in Simhanada Guggulu pill, because Water soluble extractive value of its sample was less. Guggulu contains Sterol substance and it is soluble in alcohol therefpre, Methanol soluble extractive value was more in the sample of Simhanada Guggulu pill.

CONCLUSION

It can be concluded on the basis of this Physicochemical Analysis that the *Simhanada Guggulu* pill contained sterol, less moisture, more inorganic constituents and less water soluble constituents but more research work is necessary on the drug for more information and accuracy so that standardization of drug is feasible.

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