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GLOBAL JOURNAL OF ENGINEERING SCIENCE AND RESEARCHES PHYTOCHEMICAL INVESTIGATION OF ASPARAGUS RACEMOSUS

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ABSTRACT

The very important medicinal plant *Asparagus racemosus* is widely distributed in the tropical and sub-tropical regions of India. The ancient and traditional systems of medicines i.e. Ayurveda and Unani etc. have reported it's broadly usages treating many diseases and disorders in indigenous system of medicine i.e. antioxidants, aphrodisiac, anti-diarrheal agent, curing and treating the immune system modulator and menopause. It is very much useful threatened miscarriage, decreased libido, infertility, menopause and leucorrhea, balancing pH in the cervical area. The main purpose of this work is to study and identify the medicinally active substances present in the methanolic extract found from the root powder of the plant *Asparagus racemosus*. The preliminary phytochemical screening of the extract gave the glimpses of the presence of the carbohydrates, tannins, alkaloids, sterols and flavonoids.

Keywords: Asparagus racemosus, Liliaceae, Antioxidants, Medicinal plant, Alkaloids.

I. INTRODUCTION

Shatavari (*Asparagus racemosusWilld.*) is a well knownAyurvedic drug (Sharma et al., 2005). *Asparagus racemosus*Willd.is belonging to both Liliaceae and Asparagaceae plant families (Madhavan et al., 2010). This is a woody climbing plant growing to 1-2m in height that grows in low forest areas throughout India (Gomase et al., 2010). In Indian system of medicine *A. racemosus*is an important medicinal plant and its root paste or root juice has been used in various ailments and as health tonic (Krtikar et al., 1975), (Goyal et al., 2003). The root is used to prepare medicine.

It is an important monocot medicinal plant which is distributed in tropical and subtropical forest and in central parts of India, *Asparagus racemosus*Willd., is a perennial shrub, with a tuberous rootstock, stems covered with recurved spines, linear leaves arranged in a tuft, white flowers which is sweet-scented appears in October (Patel et al., 2013). The promising potential of antimicrobial plant derived substances has attracted the attention of pharmaceutical and scientific communities during the last few years. The primary benefit of plant derived medicines is that they are relatively safer than their synthetic counterparts and offer profound therapeutic benefits and more affordable treatment (Rahman et al., 2018). *Asparagus racemosus*is used to treat various diseases such as ulcer, dyspepsia and debility. In Indian medicine it is well known as an antispasmodic, aphrodisiac, demulcent, diuretic, galactogogue, nervine tonic and refrigerant. It is also used in the treatment of diarrhoea, rheumatism, diabetes, brain complaints, jaundice, urinary disorders, blood diseases, cough and bronchitis (Battu and Madahavan et al., 2010), (Raval et al., 2012).

II. MATERIAL AND METHOD

From Nature Nursery Pipliyapala, Choithram Square, Dist. - Indore (M.P.), India the fresh and healthy roots of plant *Asparagus racemosus* were collected. The plant was identified and authenticated by the Dept. of Botany from PMB Gujarati Science College Indore.

Preparation of Extract

First of all the small amount of dried root powder was mixed in a flask with the 70% Methanol and allowed to stand for 6-7 days for the extraction after covering the flask with the aluminium foil. The extraction was allowed to put on the rotary evaporator at 50° C temperature after being filtered with the whatman filter paper no.1. Thus the stock solution 50mg/ml was prepared.





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Preliminary Phytochemical Screening

Using the standard procedures to indentify tennins, sterols, saponins, alkaloids, flavonoids, carbohydrates and amino acids the qualitative phytochemical analysis of the extract of methanol was carried out.

Table 1. Ferric Chloride Test

Experiment	Observation	Inference
In small amount of water some part	Blue black colour appeared.	Presence of tannins.
of the powdered plant sample was		
boiled and filtered, then in about 5		
ml of the filtrate a few drops of		
freshly prepared 0.2% ferric chloride		
were added.		

Table 2. Salkowski's Test

Experiment	Observation	Inference
To the methanolic extract few drops of concentrated sulphuric acid was added and shaken well before allowing to the stand.	_ 	Presence of sterol.

Table 3. Foam Test

Experiment	Observation	Inference
With the little quantity of distilled	Foam produced.	Presence of saponins.
water the small amount of extract		
was mixed and shaken well and was		
allowed to stand for 10 minutes.		

Table 4. Wagner's Reagent Test

Experiment	Observation	Inference
In the extract solution few drops of	Reddish brown colour appeared.	Presence of alkaloids.
dilute hydrochloric acid and few		
drops of Wagner's reagent were		
mixed.		

Table 5.Shinoda's Test

Experiment	Observation	Inference
The extract was dissolved in ethanol	Reddish pink colour appeared.	Presence of flavonoids.
in a test tube with few drops of		
diluted hydrochloric acid and		
magnesium turnings.		

Table 6. Benedict's Test

Experiment	Observation	Inference
With the few drops of Benedict's	Green reddish brown precipitate	Presence of carbohydrate.
reagent a small amount of extract	appeared.	
was boiled in water bath.		

Table 7. Ninhydrin's Test

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Experiment	Observation	Inference







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A small amount of extract was	Purple colour should be appeared.	Found negative result.
heated and was stand for 10 minutes		
in boiling water bath after being		
added few drops of Ninhydrin's		
solution.		

III. RESULT AND DISCUSSION

The presence of various secondary metabolites are very important for the curative properties of medicinal plants. Whatever be the results of the preliminary phytochemical screening, they all will be very helpful in future for qualifying the root powder of *Asparagus racemosus*. The below given table 8 reveals the result of the phytochemical investigation of methanolic extract of the root of *Asparagus racemosus*.

The alcoholic, aqueous and benzene extracts of *Asparagus racemosus* were subjected to different chemical tests for the detection of phyto-constituents such as Sterols, Saponins, Alkaloids, Tannins, Carbohydrats, Flavonoids, Lactones, Amino acids/Proteins, Resins and Starch (Garabadu and Sairam et al., 2009).

The root of *Asparagus racemosus* revealed the presence of the following in the phytochemical investigation of methanolic extract i.e. tannins, sterols, saponins, alkaloids, flavonoids, carbohydrates and amino acids.

Table 8: Phytochemical investigation of metabolic extract of the root of Asparagus racemosus

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Secondary Metabolites	Metabolic extract of the root of Asparagus racemosus
Tannins	+
Sterols	+
Saponins	+
Alkaloids	+
Flavonoids	+
Carbohydrates	+
Amino acid	-

IV. CONCLUSION

For the identification and authentication of a drug the importance of the standardization of crude drugs has increased very much today. In various cases identifying drug techniques fail it's originality and may exploit the uses of drug from it's medicinal traditional system. That's why this investigation was aimed and found to be significant and encouraging as well towards the result and goal.

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REFERENCES

- [1] Sharma, P. V., Dravyaguna, V. (2005), Vol. II (in Hindi), reprint, Chaukambha Bharti Academy, Varanasi.
- [2] Madhavan, V., Tijare, R. D., Mythreyi, R., Gurudeva, M. R., Yoganarasimhan, S. N. (2010). Pharmacognostical studies on the root tubers of Asparagus gonocladosBaker Alternate source for the Ayurvedic drug Satavari, Indian J Nature Resour., 1(1), 57-62.
- [3] Gomase, V. S., Sherkhane, A. S. (2010). Isolation, structure elucidation and biotransformation studies on secondary metabolites from Asparagus racemosus, Inter J Microbio., 2, 7-9.



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- [4] Krtikar, K. R., Basu, B. D. (1975). Indian Materia Medica, India, 3, 2499-2501.
- [5] Goyal, R. K., Singh, J., Lal, H. (2003). Asparagus racemosus- An update. Ind. J Med Sci., 57, 408-414.
- [6] Patel, L. S., Patel, R. S. (2013). Preliminary phytochemical analysis of root extracts of Asparagus racemosusWilld. Life Sciences Leaflets, 5, 72-77.
- [7] Rahman, Magbool, F. Al., Elnima, I., Shayoub, M. E., Hussein, S. E. (2018). Antifungul Potential of QuercusInfectoria Galls against Candida Abican-An Invitro Study, EuropeanJournal of Biomedical and Pharmaceutical Sciences, 5(2), 22-27.
- [8] Battu, G. R., Kumar, B. M. (2010). Anti-inflammatory activity of leaf extract of Asparagus racemosus Willd. Int. J. Chem. Sci., 8(2), 1329-1338.
- [9] Madahavan, V., Tijare, R. D., Mythreyi, R., Gurudeva, Yoganarasimhan (2010). Pharmacolgnostical studies on the root tubers of Asparagus racemosus Baker- Alternative source for the Ayurvedic drug Shatavari, 1(1), 57-62.
- [10]Raval, P. K., Nishteshwar, K., Patel, B.R., Shukla, J. (2012). Asparagus racemosus Wild. A Comparative Phytochemical analysis of fresh dried roots of Shatavari. International Journal of Pharmaceutical & Biological Archives, 3(6), 1458-1461.
- [11] Garabadu, D., Murugananadam, A.V., Joshi, V. K., Krishnamurthy, S. (2009). PharmacolBiochemBehav, 91(3), 283-90.
- [12]Sairam, K., Priyambada, S., Aryya, N., CandGoel, R. K. (2009)., J Ethanopharmacol, 86(1), 1-10.

