



STUDY ON THE IMMUNOMODULATORY ACTIVITY OF KSHIRVIDARI (IPOMOEA DIGITATA LINN) IN ALBINO MICE

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Abstract: The study aimed to assess the Immunomodulatory activity of the tuberous root of Kshirvidari (*Ipomoea digitata* Linn.) Kwath (decoction) and Churna (powder) in experimental animals by following standard procedure. Randomly selected mice were divided into five groups of six animals each. The root of *Ipomoea digitata* Linn.) was administered orally in the form of Kwath and Churna. Parameters like Haemoglobin, RBC count, WBC count, DLC, Platelet count, and measurement of Paw thickness were studied. Kwath (10.4 ml/kg) and Churna (1300mg/kg) of Kshirvidari showed significant improvement in paw volume. From the present study, it can be concluded that the root of Kshirvidari has Immunomodulatory activity.

Keywords: Parameters, Platelet count, volume

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INTRODUCTION

Kshirvidari (*Ipomoea digitata* Linn.), of the Convolvulaceae family, is distributed throughout India's warm and moist regions¹. It was mentioned in various Ganas (groups of drugs) of Ayurvedic classical Samhitas viz. Madhurskandha, Vidarigandaadi Gana, Balya Gana, Varnya Gana, Bruhaniya Gana etc. As per Ayurvedic literature, Kshirvidari mainly works as Vrishya, Mutral, Swarya, Varnya, Rasayana, Jeevaniya, Stanyajanan, Bruhna, etc.^{2,3} Conventionally, the root of this plant has been used in various formulations like Sivagutika, Chawanprash, Mahamash oil, etc. Also, the roots have shown Hepatoprotective activity, Hypoglycaemic activity, etc.⁴ Hence, this study was carried out to assess the Immunomodulatory activity of tuberous root of Kshirvidari (*Ipomoea digitata* auct.) in the form of Kwath (decoction) and Churna (powder) in experimental animals by Delayed-Type hypersensitivity (DTH) Method.

MATERIALS AND METHODS

Animals

Swiss albino mice were taken of either sex weighing approximately 20 to 25 grams and selected from the animal house of the National Toxicology Centre (NTC), Pune. Animals were maintained at room temperature, with 12 hours of the day and dark cycles. Standard laboratory diets were given an unlimited water supply of drinking water. The animal experiment was carried out at National Toxicology Centre (NTC), Pune; after the approval of the Institutional Animal Ethical Committee. Sheep RBCs were used as an antigen for the challenge to get an immune response.

Drug preparation

The tuberous root of the plant Kshirvidari (*Ipomoea digitata* Linn.) was self-collected from Marunji, Pune. Authentication of the plant was done at the Botanical Survey of India, Pune (voucher number-BSI/WRC/100-2/Tech./2017/24). The tuberous root of the plant was washed thoroughly in tap water, cut

into small pieces, and dried under shade. The weight of fresh tuberous root was 2kg approximately. After complete drying for 15-20 days weight of the dry tuber root was 750 grams. Two dosage forms of *Ipomoea digitata* Linn (Churna and Kwath) were prepared by the following methods given in Sharangdhar Samhita⁵. Dried tuber roots were powdered on the grinder. The mesh size of the powder was 100. Fresh Kwath was prepared every day during the experiment.

Pharmacognostic studies

Physiochemical analysis, preliminary phytochemical analysis of Churna and Kwath, and T.S. of tuberous root of Kshirvidari Kand (*Ipomoea digitata* Linn.) were done at Bhide Laboratory, Pune.

Dose selection and schedule

Two dose forms i.e. Churna and Kwath were used in the experiment. The study was carried out at two dose levels, for Kwath namely T1 (low dose - 5.2 ml/kg) and T2 (high dose -10.4ml/kg) respectively; while for Churna namely T3 (low dose – 650mg/kg), and T4 (high dose -1300mg/kg). The dose in mice was calculated by using factor 0.0026. The test drugs were administered orally.⁶

Study protocol

Cell-mediated immune response was studied using Delayed-Type Hypersensitivity (DTH) model^{7, 8}. The selected animals were divided into five groups. The first group (Group 1) was kept as Diseased control (DC), whereas the second (Group 2- T1), and third (Group 3- T2) were administered with kwath at a dose of 5.2 ml/kg and 10.4 ml/kg respectively. While the fourth group (Group 4- T3) and the fifth group (Group 5- T4) were treated with churna at doses of 650mg/kg and 1300mg/kg respectively.

Mice were primed with 0.1 ml of SRBC suspension containing 1×10^8 cells intraperitoneally on Day 0. The kwath and churna were administered orally on day 0 and continued till day 7 of challenge, to the test drug-treated groups. On the 7th day, the thickness of the right hind foot pad was measured using a Vernier calliper. The animals were then challenged on day 7 with 0.1 ml of SRBC suspension containing 1×10^8 cells in the right hind paw. A foot pad thickness was measured again after 24 hr, 48 hr, and 72 hr of the challenge. The difference between pre and post-challenge paw thickness expressed in mm was taken as a measure of DTH response. Haematological parameters viz. Hemoglobin %, RBC count, WBC count, DLC, Platelet count, and Haematocrit Count were also measured as these are supportive criteria for the immunomodulatory effect.

Table 1: Study design

Group	Day	Procedure
Group 1 Diseased control (DC)	D-0	Immunization of mice by injecting SRBC's I.P.
	D1-D6	On normal diet
	D7	Challenge with SRBC's in hind paw
	D8-D9	Measurement of paw thickness on 24 th and 48 th hour
Group 2 Treatment group Kwath (T 1)	D-0	Immunization of mice by injecting SRBC's I.P.
	D1-D6	Treated with Kwath 5.2ml/kg dose
	D7	Challenge with SRBC's in hind paw
	D8-D9	Measurement of paw thickness on 24 th and 48 th hour
Group 3 Treatment group Kwath (T2)	D-0	Immunization of mice by injecting SRBC's I.P.
	D1-D6	Treatment with Kwath 10.4ml/kg dose
	D7	Challenge with SRBC's in hind paw
	D8-D9	Measurement of paw thickness on 24 th and 48 th hour
Group 4 Treatment group Churna (T 3)	D-0	Immunization of mice by injecting SRBC's I.P.
	D1-D6	Treatment with Churna 650mg/kg dose
	D7	Challenge with SRBC's in hind paw
	D8-D9	Measurement of paw thickness on 24 th and 48 th hour

Group 5 Treatment group Churna (T 4)	D-0 D1-D6 D7 D8-D9	Immunization of mice by injecting SRBC's I.P. Treatment with Churna 1300mg/kg dose Challenge with SRBC's in hind paw Measurement of paw thickness on 24 th and 48 th hour
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Statistical analysis

Results were presented as Mean \pm SEM. Data was quantitative ANOVA test was selected for statistical analysis.

RESULT

The organoleptic study of the tuber root of Kshirvidari Kand (Ipomoea digitata Linn.) of the drug was done according to Ayurvedic parameters. Physicochemical and phytochemical analysis of the drug reveals standardization of the tuber root of Kshirvidari Kand (Ipomoea digitata Linn.) Churna and Kwath which includes determination of moisture content, total ash, acid insoluble ash, water soluble ash, T.L.C., pH value, Refractive value, and viscosity. Physicochemical analysis was done according to guidelines at Bhide Laboratory, Pune.

Observations during the experimental study show that there is no specific behavioral change. No specific change in the intake and output (urine and stool) of animals. The food intake of animals was stable throughout the experiment. No specific changes were observed.

The cell-mediated immune response of decoction and powder of Kshirvidari was assessed by DTH reaction, i.e. foot pad reaction. A dose-dependent suppression of SRBC-induced DTH response at 24, 48, and 72 hrs was observed in Kwath low dose (5.2 ml/kg), Kwath high dose (10.4 ml/kg), Churna low dose (650 mg/kg), Churna high dose (1300 mg/kg) treated groups.

The result of Delayed type hypersensitivity (DTH) response to SRBC and haematological parameters are given in Table 2.

Table 2: Effect of Kwath and Churn of I. digitata L. on paw thickness and haematological parameters

Group	Paw Volume in mm			Haematological parameters				
	At 24hr	At 48hr	At 72hr	WBC (10 ³ / μ l)	RBC (10 ⁶ / μ l)	HGB (g/L)	HCT (%)	PLT (10 ³ / μ l)
DC	3.58 \pm 0.134	3.33 \pm 0.203	2.99 \pm 0.25	7.1167	8.1333	13.09	44.967	577
T 1	2.6066 \pm 0.214	2.478 \pm 0.216	2.283 \pm 0.148	6.94	7.4117	12.807	43.617	579.5
T 2	3.18 \pm 0.290	2.915 \pm 0.2874	2.523 \pm 0.279	7.43	7.53	12.95	43.317	577.67
T 3	2.846 \pm 0.20	2.653 \pm 0.184	2.305 \pm 0.208	7.3667	8.2267	12.867	46.1	580
T 4	2.996 \pm 0.10	2.878 \pm 0.17	2.686 \pm 0.11	6.7833	7.1	12.8	43.933	582.83

Group DC-disease control, Group T1: Kwath low dose (5.2 ml/kg); Group T2: Kwath high dose (10.4 ml/kg); Group T3: Churna low dose (650 mg/kg), Group T4: Churna high dose (1300 mg/kg)

DISCUSSION

Immunomodulatory, anti-aging, anti-oxidant, and rejuvenating effects are retained under the Rasayan concept of Ayurveda, which has multiple dimensions to promote physical, and mental health and improve

the defense mechanisms of the body, and enhance longevity^{9, 10}. In classical texts of Ayurveda, Kshirvidari is mentioned as Balya, Bruhaniya, Varnya & Rasayana. Ayurvedic As per Ayurvedic Pharmacopeia of India (Vol. 5, Part 1) and other modern literature, *Ipomoea digitata* Linn. is considered as Kshirvidari. So this study of Kshirvidari Kand (*Ipomoea digitata* Linn.) was carried out to evaluate its immunomodulatory activity.

Delayed type hypersensitivity (DTH) is a mechanism associated with T cell-mediated immunity. Delayed type hypersensitivity (DTH) is a part of the process of graft rejection, tumor immunity, and most important immunity to many intracellular infectious microorganisms especially those causing chronic diseases such as tuberculosis and leprosy.

Two dosage forms i.e. Kwath and Churna were used for this experimental study as it is mentioned in various Rasayana Kalpana. Two dose levels were selected for the experiment as mentioned in Samhita and as recommended in its toxicity study.

Results of screening of immunomodulatory activity show that group treated with Kwath (10.4 ml/kg), immunity response is highest and slightly lower in Churna (1300 mg/kg). Groups treated with low doses of Kwath (5.2 ml/kg) and Churna (650 mg/kg), show lower responses than the above two groups.

In hematological parameters, WBC count is a supportive parameter for immunity response. WBC count is also more in the treated group than in the untreated group. It is higher in group treated with Churna (1300 mg/kg). There is no significant difference in RBC count. Platelet count in treated groups was slightly low than in the diseased control group, but it is not statistically significant. From the above observations, it is seen that the group treated with a high dose of Kwath shows higher overall improvement than other treated groups. These observations are supportive of the immunomodulatory activity of Kshirvidari kand and results suggest its action of immunomodulation to control autoimmune diseases. The key phytochemicals identified to date are Beta-sitosterol, scopoletin taraxerol, t-cinnamic acid, and coumarin. Scopoletin¹¹, which owns immunomodulatory, anti-inflammatory as well as antioxidant activity, and also the presence of Flavonoids are reported to possess immunological effects so it can be said that *Ipomoea digitata* Linn. works as immunomodulatory due to the presence of these chemical constituents.



Fig 1: Induction of Kwath and Churn of *I. digitate* L and measuring paw volume

REFERENCES

1. Anonymous, The Wealth of India, Vol. V, 1st ed. (Publications & Information Directorate, New Delhi), 1959, 248p.

2. Acharya YT. Charaka Samhita with Ayurvedadipika. Commentary by Chakrapani, Sutrasthan 04/02, 04/07, 04/08. Varanasi, India; Chaukhamba Surbharati Prakashan; 2011.166p.
3. Acharya YT. Sushrut Samhita with h Nibandhasamgraha commentary by Dalhana. Sutrasthan 38/04-05, 10. Varanasi; Chaukhambha Orientalia; 1997.142p
4. Anonymous. The Ayurvedic Pharmacopoeia of India. Part 1. 1st edition. Vol. 5. New Delhi; Government of India, Ministry of Health and Family Welfare, Department of Indian Systems of Medicine and Homoeopathy; 2004.33p.
5. Pandit Sharangadharacharya Sharangadhara samhita Adhamalla's Dipika and Kasiramas Gudhartha Dipika, fourth edition. Varanasi: Chaukhambha orientalia; 2000.
6. Paget GE, Barnes JM. In: Evaluation of drug activities. Pharmacometrics. Lawrence DR, Bacharach AL, editors. Vol. 1. New York: Academic Press; 1969. p. 161'
7. Lagrange PH, Mackaness GB, Miller TE. The potential of Tcell-mediated immunity by selective suppression of antibody formation with cyclophosphamide. J Exp Med 1974; 139: 1529-1539.
8. Doherty NS, Selective effects of immunosuppressive agents against the delayed hypersensitivity response and humoral response to sheep red blood cells in mice. Agents Actions, 1981; 11: 237-242.
9. Ayurveda Concept Of Rasayan Therapy (<http://www.deinayurveda.net/> word press /2009/12/ayurveda-concept-of-rasayan-therapy)
10. Science And Philosophy Of Indian Medicine(1990), edited by K.N. Gupta, published by Shree Baidyanath Ayurved Bhavan Ltd (Nagpur)
11. Bot, J & Soc, & Rauniyar, Neha & Srivastava, Deepa. (2020). *Ipomoea digitata*: A therapeutic boon from nature to mankind. The Journal of Indian Botanical Society. 100. 185-191.