

Formulation and Evaluation of Multipurpose Cream from Berberis Aristata

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ABSTRACT

In the present study, an attempt was made to formulate and evaluate herbal cream of Berberis aristata (B. aristata) for the management of Psoriasis. The study is intended to be carried out because of low cost, lesser or no side effects of herbal formulations and its potent action than the allopathic medication. They therefore provide a viable alternative for psoriasis management. To prepare an herbal cream using an ethanolic extract of Berberis aristata and assess the anti-psoriatic effectiveness of the finished product. We were able to make cream by adding varying concentrations of bees wax and borax, liquid paraffin by learning about various formulation types, such as oil in water. The analysis of many factors, including pH, viscosity, spreadability, and stability, was used to evaluate all formulations. A formulation of ethanolic extract showed antipsoriatic action. No evidence of a separate phase or ease of removal was found in the formation, but it demonstrated good spreadability, consistency, appearance, and Additionally, during irritancy trials, the pH. formulation did not cause redness, oedema, erythema, or irritation. The use of this formulation on the skin is safe. As a result, the study implies that the extract's and the cream compositions are safer and more stable, however they might also have synergistic effects.

KEYWORDS – Anti psoriatic, Berberis aristata, Erythema, Skin Irritancy Test, ethanolic extract.

I. INTRODUCTION

Psoriasis is a very common, noninfectious, inflammatory skin disease characterized by well defined, distinctive erythematous plaques yielding adherent silvery white scales, which may manifest bleeding points when removed. Psoriasis may affect any cutaneous surface, but the commonest sites are the extensor surfaces of the elbow sand knees, scalp (where scales may become extremely dense) and sacral areas. Psoriasis is either benign and localized (hands and feet) or generalized or life threatening, with associated fever, leucocytosis, arthralgias, diffuse cutaneous and mucosal pustules, secondary infection and electrolyte disturbances. Certain European and oriental countries have been exploring use of herbs in practice since the countries.

Creams which are topical formulation are more well-liked by patients since they have higher patient compliance. The herbal cream is basically oil in water type of emulsion. An ointment are semi-solid dosage preparation intended for infected diseases which carries water, waxes, hydrocarbons and volatile oils as a semi-solid. Various plants were being used as medicine much before recorded history, according to a World Health Organization (WHO) report almost 25% of human prescription drugs are made from plants while 80% of people still use conventional medical system. Berberis aristata, (family-Berberidaceae) plant is intended to be used in the herbal formulations because of its potent anti psoriatic action (anti inflammatory) due to the presence of active compound berberine. It shows minimum risk of side effects and having great potential for health management. Herbal products have been applied to human healthcare for long-established time.

In some groups, psoriasis is infrequent or uncommon, probably because of genetic causes. In a survey at a teaching hospital in Nigeria, very few psoriatic patients were identified, and the condition is reportedly uncommon among Eskimos. Indians from North or South America hardly ever exhibit it. The prevalence of psoriasis is equal in males and females, according to the current consensus, despite numerous research in the past showing a varying sex incidence. The age of onset and mean age of affected males and females varied, according to Steinberg and colleagues; the clinical disease manifested in females at a younger age, and the mean age of statistically studied groups of females

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with psoriasis was three to four years lower than that of comparable male.

Herbalism has traditionally been practised outside of the realm of mainstream medicine, but it is growing in popularity as new studies and research demonstrate how successful they are at diagnosing, treating, and preventing disease. Because of the skin's accessibility, size and exposure area medication delivery system through the skin has been considered a promising approach. The primary objective of this study was to formulate and evaluate anti-psoriatic by combining ethanol extract of Berberis aristata with additional components to provide a variety of skin advantages, including antibacterial and antiseptic properties.

***** BERBERIS ARISTATA

1.synonyms: Daruhaldhi 2.Chemical formula: C20H18NO4 3.Molecular formula:336.4g/mol.

*

4.melting point: 204-206 degree celcius

5.Uses: Daruharidra is a definitive remedy for all sorts of hormonal problems. It plays a key role in regulating periods, treating postnatal ailments, and even helps to treat excessive abdominal pain/bleeding. The plant is used in skin diseases, ophthalmic disorders, in menorrhagia, as astringent, antiperiodic jaundice, for treatment of wounds, liver problems, and ear problems.

II. MATERIALS AND METHODS.

✤ Identification, collection and authentication of plant material.

Plants of the Berberis aristata Linn. species were harvested in March 2021 from the Garhwal district of Uttarakhand. The root bark was cleaned and allowed to air dry. The scientist In-charge of the Botanical Survey of India, Allahabad (Central region), India, performed the authentication.



Fig.1 : plant of Berberis aristata

Preparation of Extract from Root-barks

The fresh root barks of Berberis aristata was collected.

The root barks was washed and dried at RT (Room temperature), then converted into a coarse powder by the help of grinder and passes through a sieve number 18 to get the uniform size.

To get rid of fat and other pigments, powdered root bark was defatted using petroleum ether (40–60%). Using a Soxhlet device, the defatted dry root barks were further extracted with ethyl acetate and subsequently with ethanol.

The extracts were dried completely in a vacuum oven until all traces of ethanol were eliminated.

In a refrigerator set between 2-8 $^{\circ}\text{C},$ the extracts were stored



Fig.2 root barks of Berberis aristata

***** FORMULATION OF CREAM

Formulation of Oil in water emulsionbased cream was formulated. The emulsifier (Beeswax) and other oil soluble components (Liquid paraffin) were dissolved in the oil phase (Part A) and heated to 75°C. The preservatives (Methyl paraben) and other wate soluble components (Glycerin, Borax) were dissolved in the aqueous phase (Part B) and heated to 15°C. After heating, the aqueous phase was added in small portions to the oil phase slowly with constant stirring to the wax and oil mixture. Ethanolic extract of the root bark was added to this and thoroughly mixed to create a homogenous material. Continue this process for 2 minutes, stir all the time then remove from the heat and stir until it gets cold then smooth cream is formed.



Sr.No.	Ingrediants	F1H	F2H	F3H
1.	Berberis aristata extract	1ml	2ml	1ml
2.	Beeswax	3.5gm	3gm	4gm
3.	Liquid paraffin	15ml	13ml	13.5ml
4.	Borax	0.4gm	0.3gm	0.5gm
5.	Methyl paraben	1ml	1ml	1ml
6.	Distailled water	q.s	q.s	q.s
7.	Rose oil	3-4 drops	2-3 drops	1-2 drops

The formula for the cream is given in Table no.



Fig 3: Prepared cream

Evaluation of cream

1). Physical evaluation

In this test, the cream was observed for color, odour, texture, state.

2). Irritancy

Mark the area (1 cm2) on the left-hand dorsal surface. Then the cream was applied to that area and the time was noted. Then it is checked for irritancy, erythema, and edema if any for an interval up to 24 h and reported

3). Wash ability

A small amount of cream was applied on the hand and it is then washed with tap water.

4). PH of the Cream

0.5 g cream was taken and dispersed in 50 ml distilled water and then PH was measured by using digital PH meter.

5). Phase separation

Prepared cream was kept in a closed container at a temperature of 25-100 °C away from light. Then phase separation was checked for 24 h for 30 d. Any change in the phase separation was observed/checked

6). Spread ability

The spreadability was expressed in terms of time in seconds taken by two slides to slip off from the cream, placed in between the slides, under certain load. Lesser the time taken for separation of the two slides better the spreadability. Two sets of glass slides of standard dimension were taken. Then one slide of suitable dimension was taken and the cream formulation was placed on that slide. Then other slide was placed on the top of the formulation. Then a weight or certain load was placed on the upper slide so that the cream between the two slides was pressed uniformly to form a thin layer. Then the weight was removed and excess of formulation adhering to the slides was scrapped off. The upper slide was allowed to slip off freely by the force of weight tied to it. The time taken by the upper slide to slip off was noted.

Spread ability= $m \times l/t$

Where,

m= Standard weight which is tied to or placed over the upper slide (30g)

l= length of a glass slide (5 cm)

t= time taken in seconds.



7] Greasiness

Here the cream was applied on the skin surface in the form of smear and checked if the smear was oily or grease-like.

8] Accelerated Stability Testing

The purpose of stability testing is to provide evidence on how the quality of drug substance or drug product varies with time under the influence of variety of environmental factors such as temperature, humidity and light and enables to recommend storage condition and to predict the shelf life. Stability study for cream was performed at accelerated condition Le.40°C 2° C / 75% RH±5% RH. TH formulations were kept both at room and elevated temperature and observed on 0,5th, 10th, 15th and 20th day for the various parameters.

9] Homogeneity

The formulations were tested for the homogeneity by visual appearance and by touch.

III. RESULTS AND DISCUSSION

Evaluation results of all the three formulations are gives below. **1)Physical evaluation**

In this test color, odour, texture and state of the three formulations were checked.

Sr. no.	Parameter	Evolution
1	Colour	Faint white
2	Odour	Pleasant
3	Texture	Smooth
4	State	Semi-solid

Table 2: physical parameter of cream

Sr. no.	Parameters	F1H	F2H	F3H
1	Irritancy	Nil	Nil	Nil
2	Washability	Easily Washable	Easily Washable	Easily Washable
3	pH of cream	6.6	6.2	6.67
4	Phase separation	No phase separation	No phase separation	No phase separation
5	Spreadability	22.18	30.4	15.8
6	Greasiness	Non-greasy	Non-greasy	Non-greasy
7	Homogeneity	Homogeneous	Homogeneous	Homogeneous

 Table 3: Observation and evaluation of parameters of cream

IV. CONCLUSION

By using barbaris aristata the cream showed a multipurpose effect and these herbal ingredients showed significant different activities. Based on results and discussion, the formulations F1H, F2H, F3H were stable at room temperature and can be safely used on the skin.T he result of different tests of cream showed that the formation could be used topically in order to protect skin against damage. Natural remedies are more acceptable in the belief that they are safer with fewer side effects than the synthetic ones.

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ABBREVIATIONS

pH – Hydrogen ion Concentration O/W- oil in water F1H,F2H ,F3H- Formulation trial with different concentration of cream

REFERENCE:-

[1]. Sk Uddandu Saheb, Aduri Prakash Reddy, K Rajitha, B Sravani, B Vanitha. Formulation and evaluation of cream from naturally containing plant extracts. World J Pharm Pharm Sci 2018;7:851-62.

DOI: 10.35629/4494-090222522257 Impact Factor value 7.429 | ISO 9001: 2008 Certified Journal Page 2255



- [2]. Geetha, M. (2014). Anti-psoriatic activity of flavonoids from Cassia tora leaves using the rat ultraviolet B ray photodermatitis model. Revista Brasileira de Farmacognosia, 24, 322-329.
- [3]. Loo, Y.S., Madheswaran, T., Rajendran, R., Bose, R.J. (2020). Encapsulation of berberine into liquid crystalline nanoparticles to enhance its solubility and anticancer activity in MCF7 human breast cancer cells. Journal of Drug Delivery Science and Technology, 57.
- [4]. Zhong, L., Luo, N., Zhong, X., Xu, T., & Hao, P. (2022). The immunoregulatory effects of natural products on psoriasis via its action on Th17 cells versus regulatory T cells balance. International Immunopharmacology, 110, 109032.
- [5]. Chaudhary, A., & Mittal, S. (2022). A study on ethosomes as mode for transdermal delivery of an antipsoriatic drug. Chinese journal of medical genetics, 31(3).
- [6]. Balkrishna, A., Sakat, S., Joshi, K., Singh, R., Verma, S., Nain, P., ... & Varshney, A. (2022). Modulation of psoriatic-like skin inflammation by traditional Indian medicine Divya-Kayakalp-Vati and Oil through attenuation of pro-inflammatory cytokines. Journal of traditional and complementary medicine, 12(4), 335-344.
- [7]. Van Der Fits, L., Mourits, S., Voerman, J. S., Kant, M., Boon, L., Laman, J. D., Lubberts, E. Shim, S.K., Kim, H., Kwak, C.D. (2009). Imiquimod-induced psoriasis-like skin inflammation in mice is mediated via the IL-23/IL-17 axis. The Journal of Immunology, 182(9), 5836-5845.
- [8]. Muruganantham, N., Basavaraj, K. H., Dhanabal, S. P., Praveen, T. K., Shamasundar, N. M., & Rao, K. S. (2011). Screening of Caesalpinia bonduc leaves for antipsoriatic activity. Journal of ethnopharmacology, 133(2), 897-901.
- [9]. Shrivastav, S., Sindhu, R., Kumar, S., & Kumar, P. (2009). Anti-psoriatic and phytochemical evaluation of Thespesia populnea bark extracts. Int J Pharm Pharm Sci, 1(1).
- [10]. Reich K. The concept of psoriasis as a systemic inflammation: Implications for disease management. J Eur Acad Dermatol Venereol. 2012;26(Suppl 2):3–11.

- [11]. Katare OP, Raza K, Singh B, Dogra S. Novel drug delivery systems in topical treatment of psoriasis: Rigors and vigors. Indian J Dermatol Venereol Leprol. 2010;76:612–21. [PubMed] [Google Scholar]
- [12]. Syed TA, Ahmad SA, Holt AH, Ahmad SA, Ahmad SH, Afzal M. Management of psoriasis with Aloe vera extract in a hydrophilic cream: A placebo-controlled, double-blind study. Trop Med Int Health. 1996;1:505–9.
- [13]. Srivastava N, Singh K, Amrit K. Formulation and evaluation of Seabuckthorn leaf extract loaded ethosomal gel. Asian J Pharm Clin Res. 2015;8:309–12. [Google Scholar]
- [14]. Misal J, Dixit G, Gulkari V. Formulation and evaluation of herbal gel. Indian J Nat Prod Res. 2012;3:501–5. [Google Scholar]
- [15]. Ellis J, Parlapally S, Cherukupalli N, Bhumireddy SR, Sripadi P, Anisetti R, et al. Chemical profiling and anti-psoriatic activity of methanolic extract of Andrographis nallamalayana. Natural Product Res. 2015;30:1256–61.
- [16]. Parlapally S, Cherukupalli N, Bhumireddy SR, Sripadi P, Anisetti R, Giri CC, et al. Chemical profiling and anti-psoriatic activity of methanolic extract of Andrographis nallamalayana JL Ellis. Nat Prod Res. 2016;30:1256–61.
- [17]. More B, Sakharwade S, Tembhurne S, Sakarkar D. Evaluation for skin irritancy testing of developed formulations containing extract of Butea monosperma for its topical application. Int J Toxicol Appl Pharmacol. 2013;3:10–3. [Google Scholar]
- [18]. Ghanbarzadeh S, Arami S. Enhanced transdermal delivery of diclofenac sodium via conventional liposomes, ethosomes, and transfersomes. Biomed Res Int. 2013;2013:616810. [PMC free article] [PubMed] [Google Scholar]
- [19]. Lichtenberg D, Opatowski E, Kozlov MM. Phase boundaries in mixtures of membrane-forming amphiphiles and micelle-forming amphiphiles. Biochim Biophys Acta. 2000;1508:1–19. [PubMed] [Google Scholar]
- [20]. Yoshioka T, Sternberg B, Florence AT. Preparation and properties of vesicles

DOI: 10.35629/4494-090222522257 Impact Factor value 7.429 | ISO 9001: 2008 Certified Journal Page 2256



(niosomes) of sorbitan monoesters (Span 20, 40, 60 and 80) and a sorbitan triester (Span 85) Int J Pharm. 1994;105:1–6. [Google Scholar] Nimisha Fatima Z, Kaur C. Formulation and Performance evaluation of Berberis aristata extract loaded ethosomal gel. Asian J Pharm. 2017;11:1–9. [Google Scholar]

- [21]. Elsayed MM, Abdallah OY, Naggar VF, Khalafallah NM. Deformable liposomes and ethosomes: Mechanism of enhanced skin delivery. Int J Pharm. 2006;322:60–6. [PubMed] [Google Scholar]
- [22]. Jain S, Jain P, Umamaheshwari R, Jain N. Transfersomes – A novel vesicular carrier for enhanced transdermal delivery: Development, characterization, and performance evaluation. Drug Dev Ind Pharm. 2003;29:1013–26. [PubMed] [Google Scholar]
- [23]. Kaur CD, Saraf S. Topical vesicular formulations of Curcuma longa extract on recuperating the ultraviolet radiationdamaged skin. J Cosmet Dermatol. 2011;10:260–5. [PubMed] [Google Scholar]
- [24]. Kaur CD, Saraf S. Photoprotective herbal extract loaded nanovesicular creams inhibiting ultraviolet radiations induced photoaging. Int J Drug Deliv. 2011;3:699. [Google Scholar]
- [25]. Duangjit S, Opanasopit P, Rojanarata T, Ngawhirunpat T. Evaluation of meloxicam-loaded cationic transfersomes as transdermal drug delivery carriers. AAPS PharmSciTech. 2013;14:133–40.
- [26]. Khan NR, Wong TW. Microwave-aided skin drug penetration and retention of 5fluorouracil-loaded ethosomes. Expert Opin Drug Deliv. 2016;13:1209–19.
 [PubMed] [Google Scholar]