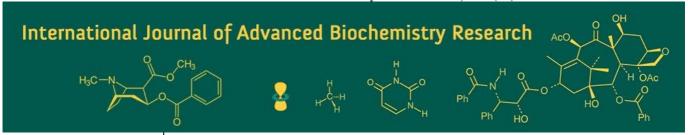
International Journal of Advanced Biochemistry Research 2025; SP-9(10): 2015-2020



ISSN Print: 2617-4693 ISSN Online: 2617-4707 NAAS Rating (2025): 5.29 IJABR 2025; SP-9(10): 2015-2020 www.biochemjournal.com Received: 05-07-2025 Accepted: 07-08-2025

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Phenology and reproductive behaviour of endangered medicinal plant: *Valeriana jatamansi* Jones ex Roxb.

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DOI: https://www.doi.org/10.33545/26174693.2025.v9.i10Sy.6132

Abstract

Understanding a species' phenology and reproductive biology is necessary for the restoration of endangered plant species. *Valeriana jatamansi* is an endangered plant in IUCN Red Data Book due to excessive deforestation and over exploitation. The study delves into the phenological patterns and reproductive strategies of this medicinally significant species across six distinct sources: JK1, JK2, JK3 (from Jammu and Kashmir), and HP1, HP2, HP3 (from Himachal Pradesh). *Valeriana jatamansi* Jones is a perennial herb that ranges in height from 14 to 45 cm belonging to the family Valerianaceae and order Dipsacales. *Valeriana jatamansi* exhibits remontant flowering behavior, re-blooming in spring and autumn. Spring phenophases showed better synchronization and longer duration compared to autumn. Spring flowering initiation occurred between 22 February to 25 with a duration of 53-55 days, while autumn flowering showed greater variation (November 4-8) lasting 42-46 days. The species exhibits a gynomonoecious reproductive system with hermaphrodite flowers. Pollination peaks occurred morning and evening, with *Apis mellifera* and *Musca domestica* as primary pollinators. The data produced by the current study may prove helpful and effective in various breeding methods and the preservation of this medicinal plant species.

Keywords: Valeriana jatamansi, phenology, reproductive biology, endangered medicinal plants

Introduction

Valeriana jatamansi Jones is also known as Indian valerian in English, Mushkbala or Sugandhbala in Hindi and Tagar in Sanskrit (Thakur et al., 2018) [21]. The term "Valerian" was originally used in the ninth century by an Indian physician, and it was taken from the word "Velo," which means "powerful drug" (Bhatt et al., 2012) [3]. Valeriana jatamansi Jones is a perennial herb that ranges in height from 14 to 45 cm. Belonging to the family Valerianaceae and order Dipsacales. The species is mostly found in the Himalayan regions of Afghanistan, Bhutan, Burma, India, Nepal, and Pakistan including cold and temperate regions at elevations of 1500-3000m (Ekhteraei et al., 2010) [5]. V. jatamansi is known for the sedative and tranquilizing effects that are attributed to the presence of the bioactive chemical 'valepotriates' in the roots and rhizome [Wagner et al., 1980; Grusla et al., 1986; Houghton, 1999; Mishra, 2004] [24, 11, 12, 15]. V. jatamansi has antidepressant, anticonvulsive, antispasmodic, antitumor, cytotoxic, analgesic, laxative, carminative, anti-dyspeptic, antimicrobial, and a number of other pharmacological qualities (Dhiman et al., 2020) [4]. While having such valuable medicinal properties and being an endangered species, it is essential to conserve this medicinal herb and for that studying phenology and reproductive behaviour of Valeriana jatamansi Jones is necessary.

Phenology is the investigation of the planning of repeating organic occasions, the causes of their timing concerning biotic and abiotic powers, and the interrelation between periods of the same or different species (Leith, 1974) [13]. The information on timing of phenological occasions and their changeability can assist with getting more steady harvest yields and quality through improved and feasible harvest the board giving dates to ideal water system, fertilizing, and crop security (Ruml and Vulić, 2005) [18].

Study of reproductive biology is a crucial component of plant ecology that plays an important role in the conservation and management of a plant species by scientific way. It is

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Division of Forest Biology and Tree Improvement, Faculty of Forestry, Sher-e-Kashmir University of Agricultural Sciences and Technology of Kashmir, Benhama, Campus, Ganderbal, Jammu and Kashmir, India the pollination that makes the first male and female interactions in plants which is also accompanied by natural biotic and abiotic factors. The successful pollination is represented by the pollinator's potential, daily opening and number of flowers. Therefore, the study of reproductive biology of flowering plants is essential to understand the breeding systems and pollination that helps in regulating the genetic structure of plant populations and determine the barriers of seed and fruit set for the purpose of conservation (Tandon et al., 2003) [22]. Although large efforts have been invested in research, documentation and analysis, of Valeriana jatamansi Jones (Syn. V. wallichii DC.) regarding its pharmacological makeup, chemical composition, and therapeutic applications, there remains a significant gap in understanding the species' phenology and reproductive biology that are key ecological aspects for its sustainable management, conservation and cultivation strategies.

Materials and Methods

The study was conducted at Faculty of Forestry SKUAST-Kashmir, Benhama Ganderbal during the years 2023 – 2024. The study of phenology and reproductive biology was conducted by sourcing rhizomes of *Valeriana jatamansi* of uniform size and weight. These rhizomes were collected from six locations across two regions: Jammu & Kashmir (JK1, JK2, JK3) and Himachal Pradesh (HP1, HP2, HP3). Three sources were selected from each region. The collected rhizomes were then planted in a randomized block design at the onset of spring-2023, and observations were systematically recorded throughout the growing season to capture the plant's growth, reproductive stages, and phenological events.

Phenology

Throughout the growing season, which included both the spring and autumn flowering phases, various observations were made on the phenology and reproductive biology of *Valeriana jatamansi*. The following parameters were recorded: initiation of vegetative bud burst, initiation of flower bud, commencement of flowering, peak period of flowering, duration of peak period of flowering, termination of flowering and duration of flowering. The observations were taken on alternate days.

Morphological observations

Average number of floral spikes per inflorescence, average number of floral clusters per spike and average number of flowers per cluster were recorded by randomly selecting four plants per replication and then evaluated for the same during the peak flowering period.

Reproductive study

The reproductive study of *Valeriana jatamansi* was carried out for sex determination in which flowers from ten randomly selected spikes per source were examined using a magnifying glass to determine whether they were staminate, pistillate, or hermaphrodite. Anthesis was recorded during the peak flowering hours between 6 a.m.–11 a.m. and 4

p.m.-7 p.m. Pollination was observed within the peak hours of anthesis. Pollinating agents, both biotic (such as insects, birds) and abiotic (such as wind), were observed for their role in transferring pollen. Pollinators were identified through direct observation during the peak flowering and anthesis periods.

Results and Discussion

Seasonal Phenology of *Valeriana jatamansi* Jones ex Roxb across different sources.

The seasonal phenology of Valeriana jatamansi Jones ex Roxb. revealed distinct variations between autumn 2023 and spring 2024 across sources JK1-JK3 and HP1-HP3 as mentioned in Table 1. In spring, the initiation of vegetative bud burst occurred earlier (7-15 February) compared to autumn (20–27 October), an effect of favourable temperatures and longer photoperiods promoting early growth. Flower bud initiation followed shortly after vegetative bud burst, occurring between 13-21 February in spring and 26 October–5 November in autumn, with the JK sources consistently showing earlier initiation, reflecting genotypic adaptation to local environments. Flowering commenced earlier and more synchronously in spring (22-29 February) than in autumn (4-12 November), indicating stable climatic conditions during the spring season. Peak flowering was observed between 10-27 March in spring and 10-29 November in autumn, with JK1 peaking earliest and HP2 latest in both seasons, demonstrating a slightly staggered but overlapping pattern. The duration of the peak flowering period was consistent across both seasons (14–16 days), reflecting robust physiological control. Termination of flowering occurred between 11-18 April in spring and 13-19 December in autumn, showing moderate overlap among sources. Overall flowering duration was longer in spring (53–55 days) than in autumn (42–46 days), with JK1 exhibiting the longest duration in both seasons, suggesting superior reproductive potential and adaptability. These patterns underscore how temperature, photoperiod, and genotypic differences govern phenological development, with spring conditions facilitating earlier and longer reproductive phases. The observations align with previous findings by Fenner 1998 ^[7], Ghelardini *et al.*, 2006 ^[10], Liang and Schwartz 2014 ^[14], Reader 1982 ^[19], Elzinga *et* al., 2007 [6], Forrest and Miller-Rushing 2010 [8], and Fox et al., 1997 [9], which collectively highlight the interactive influence of environmental cues and genetic factors on phenological synchronization and duration in temperate species. Figure (1-4).

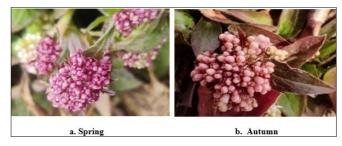


Fig 1: Initiation of vegetative bud burst

Table 1: Seasonal phenology of Valeriana jatamansi Jones ex Roxb. in Autumn 2023 and Spring 2024

| | 1 | T *.* .* . | | | | D 11 0 1 | | |
|--------|---------------|---|---|--|---|--|--|-----------------------|
| Season | Sources | Initiation of vegetative bud burst | Initiation of flower bud | Commencement of flowering | Peak period of flowering | Duration of peak period of flowering | Termination of flowering | Duration of flowering |
| Autumn | JK1 | 20 th Oct to 23 rd Oct | 26 th Oct to 30 th Oct | 4 th Nov to 6 th Nov | 10 th to 24 th Nov | 15 Days | 13 th to 19 th Dec | 46 days |
| Autumn | JK2 | 21st Oct to 25th Oct | 28 th Oct to 2 nd Nov | 6 th Nov to 9 th Nov | 13 th to 27 th Nov | 15 Days | 13th to 18th Dec | 43 days |
| Autumn | JK3 | 20 th Oct to 23 rd Oct | 25 th Oct to 30 th Oct | 5 th Nov to 9 th Nov | 11 th to 25 th Nov | 15 Days | 14 th to 19 th Dec | 45 days |
| Autumn | HP1 | 22 nd Oct to 27 th Oct | 29 th Oct to 4 th Nov | 7 th Nov to 11 th Nov | 14 th to 28 th Nov | 15 Days | 14 th to 18 th Dec | 42 days |
| Autumn | HP2 | 23 rd Oct to 28 th Oct | 31st Oct to 5th Nov | 8 th Nov to 12 th Nov | 15 th to 29 th Nov | 15 Days | 15 th to 19 th Dec | 42 days |
| Autumn | HP3 | 22 nd Oct to 27 th Oct | 30 th Oct to 5 th Nov | 7 th Nov 11 th Nov | 13 th to 28 th Nov | 16 Days | 14th to 19th Dec | 43 days |
| Autumn | Overall range | 20th to 28th Oct | 26 th Oct to 5 th Nov | 4 th to 12 th Nov | 10 th to 29 th Nov | 15 – 16 days | 13th to 19th Dec | 42 to 46 days |
| Spring | JK1 | 7 th Feb to 10 th Feb | 13 th Feb to 17 th Feb | 22 nd Feb to 26 th Feb | 10 th to 25 th Mar | 16 days | 12 th to 17 th April | 55 days |
| Spring | JK2 | 9 th Feb to 12 th Feb | 15 th Feb to 18 th Feb | 22 nd Feb to 26 th Feb | 11 th to 26 th Mar | 16 days | 11 th to 16 th April | 54 days |
| Spring | JK3 | 8 th Feb to 12 th Feb | 15 th Feb to 19 th Feb | 25th Feb to 29th Feb | 13 th to 27 th Mar | 15 days | 13 th to 18 th April | 53 days |
| Spring | HP1 | 11 th Feb to 15 th Feb | 16 th Feb to 20 th Feb | 23 rd Feb to 27 th Feb | 13 th to 27 th Mar | 15 days | 12 th to 16 th April | 53 days |
| Spring | HP2 | 12 th Feb to 15 th Feb | 17 th Feb to 21 st Feb | 24th Feb to 29th Feb | 14 th to 27 th Mar | 14 days | 13 th to 18 th April | 54 days |
| Spring | HP3 | 10 th Feb to 12 th Feb | 14 th Feb to 19 th Feb | 23 rd Feb to 28 th Feb | 12 th to 27 th Mar | 16 days | 12 th to 16 th April | 53 days |
| Spring | Overall range | 7 th to 15 th Feb | 13 th to 21 st Feb | 22 nd to 29 th Feb | 10 th to 27 th Mar | 14 to 16 days | 11 th to 18 th April | 53 to 55 days |



Fig 2: Initiation of flower bud burst



Fig 3: Commencement of flowering



Fig 4: Peak period of flowering

Morphological observations

Morphological observations for Valeriana jatamansi across spring 2024 and autumn 2023 revealed clear seasonal and genotypic influences on floral traits. The average number of floral spikes per inflorescence was higher in spring (9.46-15.55) compared to autumn (7.65-8.82), indicating that favourable spring conditions promote spike development. Similarly, the average number of floral clusters per spike showed a marked seasonal decline, with JK1 achieving the highest values in both seasons (41.00 in spring, 25.45 in autumn) and HP2 and HP3 recording the lowest in autumn (13.76 and 13.73, respectively), reflecting reduced floral productivity under cooler autumn conditions. In terms of flowers per cluster, HP1 (73.52), HP2 (70.10), and JK3 (69.99) led in spring, while autumn values for all sources dropped, with HP3 showing the lowest (46.70), underscoring environmental sensitivity in flower production. JK sources generally performed better in spike and cluster production, while HP sources excelled in flowers per cluster, highlighting distinct genetic strengths. Across all traits, the seasonal reduction in autumn supports the conclusion that lower temperatures, shorter day lengths, and light availability constrain reproductive development, while spring conditions enhance it. These findings corroborate earlier reports by Singh and Gaurav 2010 [20] and Akter et al., 2017 [1], which emphasize the combined influence of genotype and environmental factors on floral trait variability and productivity in medicinal plants.

 Table 2: Morphological variation in Valeriana jatamansi across different sources during Autumn-2023 and Spring-2024.

| Source | Avg. number of floral spikes per inflorescence | Avg. number of floral clusters per spike | Avg. number of flowers per cluster |
|--------|--|--|------------------------------------|
| JK1 | 8.65^{ab} | 25.45 ^a | 65.58 ^a |
| JK2 | 8.82^{a} | 24.57 ^a | 64.19 ^a |
| JK3 | 8.27 ^{ab} | 23.75 ^{ab} | 55.72 ^{ab} |
| HP1 | 8.10 ^{ab} | 19.44 ^b | 62.30 ^a |
| HP2 | 7.65 ^b | 13.76° | 53.77 ^{ab} |
| HP3 | 7.69 ^b | 13.73° | 46.70 ^b |
| JK1 | 10.10 ^b | 41.00° | 61.28 ^a b |
| JK2 | 10.58 ^b | 35.62ª | 55.32 ^b |
| JK3 | 15.55° | 37.01 ^a | 69.99 ^a b |
| HP1 | 9.81 ^b | 24.06 ^b | 73.52 ^a |
| HP2 | 9.46 ^b | 15.75 ^b | 70.10 ^a b |
| HP3 | 10.19 ^b | 23.36 ^b | 60.18 ^{ab} |

^{*}Values followed by the same superscript are not significantly different.

Reproductive Biology

The observations and data collection related to reproductive biology were conducted solely during the spring flowering season. This approach was chosen to ensure consistency and focus on the primary flowering phase when environmental conditions and phenological events are most prominent.

Sex determination

The reproductive biology study of *Valeriana jatamansi* across six sources (JK1–JK3, HP1–HP3) confirmed a gynomonoecious system, with both pistillate

and hermaphrodite flowers (Figure-5-a, b) present but hermaphrodites consistently dominant (ratios 66:34 to 70:30). Hermaphrodite numbers were highest in HP1 (69.82%) and JK1 (69.15%), lowest in HP3 (67.42%), yet maintained across regions, suggesting genetic stability and a mixed mating strategy that supports cross-pollination and reproductive assurance. These stable floral ratios align closely with Mukherjee & Chakraborty 2014 [16] and Thakur *et al.*, 2018 [21], who documented similar patterns in temperate Himalayan populations, highlighting the ecological advantage of this trait in variable environments.

Table 3: Floral morphs in Valeriana jatamansi across six sources in spring 2024.

| Course | Floral Morphs (| Ratio of H:P | |
|--------|-------------------------------|----------------------------|--------------|
| Source | Avg. number of Hermaphrodites | Avg. number of Pistillates | Katio of H:P |
| JK1 | 45.35 | 20.23 | 69.15: 30.85 |
| JK2 | 42.59 | 21.60 | 66.35: 33.65 |
| JK3 | 38.32 | 17.40 | 68.77: 31.23 |
| HP1 | 43.50 | 18.81 | 69.82: 30.18 |
| HP2 | 36.34 | 17.43 | 67.58: 32.42 |
| HP3 | 31.49 | 1 5.21 | 67.42: 35.58 |

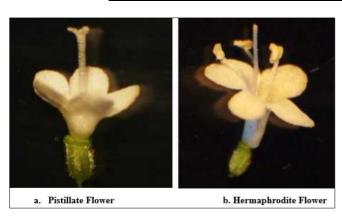


Fig 5: Sex Determination of Valeriana jatamansi

Anthesis and Pollination:

The anthesis period of *Valeriana jatamansi* was observed to occur consistently across six sources during two distinct temporal windows: morning (7:00–12:00 AM) and evening (5:30–6:30 PM). This bimodal anthesis pattern was closely associated with pollinator activity, with bees and flies serving as primary diurnal pollinators in the morning and crepuscular pollinators contributing to evening pollination. Similar observations were reported by Nautiyal *et al.*,

(2009) [17] in *Aconitum heterophyllum*, a critically endangered medicinal plant of the Himalayas. This timing likely enhances the plant's chances of successful pollination by aligning flower availability with the activity periods of different pollinator groups.

Pollinating agent

Observations across all six sources identified two primary pollinators: bees (Apis mellifera) (Figure-6-a) and flies (Musca domestica) (Figure-6-b), with Apis mellifera being the dominant pollinator. The findings align with Verma et al., (2019) [23], who documented similar pollinator diversity in Valeriana wallichii. However, the limited diversity of pollinators in Valeriana jatamansi suggests a more specialized relationship. This specialization may enhance pollination efficiency but also poses risks under environmental changes affecting pollinator populations, as noted by Bhatnagar and Kumari (2024) [2]. The consistency of pollinator types across all sources, suggests that Valeriana jatamansi has evolved specific floral traits that effectively attract these particular pollinators, possibly through visual or olfactory cues, as similarly noted by Verma et al., (2019) [23] in their study of Valeriana wallichii.

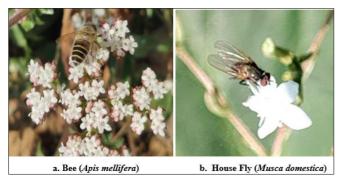


Fig 6: Pollinating Agents of Valeriana jatamansi Jones

Conclusion

In conclusion, *Valeriana jatamansi* has provided a comprehensive understanding of the specie's phenology and reproductive biology. Phenological stages such as vegetative bud burst, flower bud initiation, and peak flowering periods, revealed that genetic and environmental factors significantly influence these traits, with JK sources exhibiting earlier and more consistent flowering patterns. Additionally, the reproductive biology of *Valeriana jatamansi* was characterized by its gynomonoecious nature and crosspollination mechanisms, contributing to genetic stability across varied environmental conditions.

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