**INTRODUCTION**

The genus Valeriana belonging to the family Valerianaceae contains around 250 types of species which is distributed in all temperate regions of the world. Among them in India, 16 species were found and out of them two subspecies of genus and five species in habitat at higher altitude range of Central Himalayas. The most popular name for this genus is "Valerian." Valeriana jatamansi is commonly known as India Valerian, Muskhal, Sugandhbhala (Hindi), and Tagar (Sanskrit) [1]. Indian Valeriana is a small herbaceous species distributed in tropical and temperate Himalaya up to an altitude of 3000 m between 1500 and 1800 m in Khasi and Jaintia hills; this has been used as an ingredient for herbal medicine in Indian system of medicine and substitutes for European Valeriana officinalis in India. Geographically, this species is available in different temperate regions and has diverse genetic and morphological features, which will affect the level of active constituents present [2]. *V. jatamansi* is a hairy dwarf and rhizomatous perennial herb growing up to 0.5 m with thick root stacks, covered with horizontal descending fibers and pubescent stem and radical leaves 1-3 cm in diameter. Flowers are white or tinged in pink color, fruits are crowned by persistent pappus calyx. This species is gynodioecious, the fruiting and flowering time occurs during the period of March–June and mode of proliferation is both sexual through seeds and abiogenetic through rhizome. *V. jatamansi* used in Ayurvedic and Unani system of medicine. The roots and rhizomes are used to treat insomnia, blood and circulatory disorders, asthma, dry cough, jaundice, seminal weakness, cardiac debility, and skin diseases. Its herbal oil is widely used in pharmacological industries and in hair preparations [3,4].

**Phytochemical constituents**

*V. jatamansi* main active constituents are sesquiterpenes, coumarins, iridoids, lignanoids, alkaloids, and flavonoids. From literature, the reported methods performed for different pharmacological activities have been thoroughly discussed and compiled.

**Macросcopical characters of dried rhizome**

Dried rhizome is prolonged cylindrical in shape 2.5–12 cm long and 2–3.5 cm thick covered by groups of thick brownish rootlets bundles framing a network. The leaves are rose colored slightly pink or blue in sense.

- **Color:** Outer layer of rhizome is dark brown
- **Inner layer** is brown or yellow in color.
- **Odor:** Highly agreeable.
- **Taste:** Aromatic, bitter, and slightly camphoraceous.
- **Size:** 2.5–12 cm in length.
- **Shape:** Elongated and cylindrical.

**Vernacular names**

| Language | Vernacular names |
|----------|------------------|
| Hindi    | Balchhari, Mansi, Nihani, Smak, Sumaya, Tagar |
| Kannada  | Jatate, Naati jatamansi, Nandu bathu, Tagara |
| Malayalam| Takaram          |
| Marathi  | Thagar mool      |
| Sanskrit | Jatamansi, Natah, Tagarah |
| Tamil    | Shadamangie, Takaram |
| Telugu   | Tagara           |

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**Taxonomical classification**

| Kingdom: Viridiplantae |
|-----------------------|
| Class: Asterids        |
| Order: Dipsacales      |

**Phytochemical constituents**

*V. jatamansi* main active constituents are sesquiterpenes, coumarins, iridoids, lignanoids, and sesquiterpenes include valeric acid, and other derivatives include valeranone and valerenal. Moreover, lignanoids...
include pinoresinol-4-0-d-glucoside and lignans 8'-hydroxy-pinoresinol. Iridoids include valepotriates (valtrate, didovaltrate, acetylvaltrate, and 3-isovaleryloxyhydroxylvaltrate) [6,7]. Alkaloids include chatinine, nordepoline, norfephbine, thalipherpine, nantenine, phenanthrene, phoebine, dehydropholine, valerine, valerianine, and oxoaporphine. Flavonoids in Valeriana mainly contain acacetin, hesperidin and methylapigenin, luteolin, quercetin, kaempferol, linarin, and luteolin. The other constituents are volatile oil, essential oil, sugar, bitter extractive matter, starch, gum, resin, and ketones [8,9].

Traditional uses
In traditional system of medicine, the roots of plant are used to treat ulcer, convulsion, asthma, dry cough, jaundice, seminal weakness, cardiac debility and skin diseases, leprosy, general debility, and for sleep enhancement. Rhtzones and roots of V. jatamansi recommended to treat insomnia and blood, circulatory, and mental disorders. It stimulates central nervous system, acts as nerve tonic, and used in the treatment of anxiety and tremors. The plant was used as stimulant, hypotensive, and sedative. Its oil is used in perfumery preparation and insect repellent formulation [10].

In vitro and in vivo studies
Antioxidant activity
Methanolic, chloroform, and aqueous extracts of the dried roots of V. jatamansi were extracted and assessed for their polyphenol and flavonoid content. The antioxidant activity of the three solvent extracts of V. jatamansi roots and essential oil (100 µg/ml) and was assessed by 2,2-diphenyl-1-picrylhydrazyl (DPPH) radical scavenging and ferrous chelation power assay [11]. Methanolic extract was found to be best when compared to that of essential oil [12]. In another study, the antioxidant activity and anti-inflammatory activity were studied by hydroalcoholic extracts of V. jatamansi using DPPH free radical scavenging and the percentage (%) inhibition of proteins denaturation, respectively, using diclofenac sodium as the reference standard. A substantial decrease of various mediators of inflammation was seen due to the presence of bioactive compounds such as flavonoids, tannins, and polyphenols [13].

Anti-inflammatory activity
The aqueous and methanolic extracts from the rhizomes of V. jatamansi had shown to have significant anti-inflammatory activity. It decreases the inflammation in carrageenan-induced paw edema method, in mice at dose levels of 100, 150, and 200 mg/kg in the presence of saline as control and reference drug was aspirin. Both the extracts significantly decreased due to inhibition of inflammatory mediators histamine, prostaglandin, and serotonin synthesis and exhibited anti-inflammatory effect at 200 mg/kg dose [14].

Antibacterial activity
The antibacterial activity was explored using hexane, chloroform, and methanolic extracts from rhizomes of V. jatamansi. Extended-spectrum B-lactamase produced by Escherichia coli, Enterobacter aerogenes, Klebsiella pneumoniae, and Hafnia alvei were isolated from urinary tract infection and subjected to double disc diffusion assay using three solvent extracts [15]. The inhibition was significant with hexane extract and can be given along with other antibiotics as a new alternative treatment for urinary tract infection [16]. The methanolic, chloroform, and hexane extracts of V. jatamansi were evaluated for their antimicrobial effect against strains Staphylococcus aureus, Pseudomonas aeruginosa, Micrococcus luteus, E. coli, Salmonella enteritidis, Lactobacillus plantarum, and Staphylococcus epidermidis. Among them, the hydroalcoholic extracts with doses of 0.3–0.7 mg/ml showed better antibacterial activity against S. aureus, P. aeruginosa by cup plate method using Soyabean Casein Digest Agar medium, and chloramphenicol was standard antibiotic used at 1 mg/ml. The antibacterial activity was due to the presence of valerenic acid as an important chemical constituent which was isolated and characterized by thie-layer chromatography (TLC) and high-performance liquid chromatography (HPLC) by comparing the R and R values using reference standard compound can be a better choice as a new antibacterial agent [17].

Cytotoxic activity
The underground parts, roots and rhizomes of all three Valeriana species V. jatamansi (V. wallichii), V. officinalis, and V. edulis Nutt. possess cytotoxic activity which has been proven by different studies. Fresh extracts and stored extract tinctures of three valerian species were explored for active principles baldrinals, valepotriates, and valerenic acid and cytotoxic activity was performed against COLO 320 a human colorectal and GLC (4) a human small cell lung cancer lines, by utilizing the microculture tetrazolium (MTT) assay [18]. A significant highest toxicity was observed by diene-type valepotriates (isovaltrate, acetylvaltrate, and valtrate) with IC50 (1–6 µM), following continuous incubation. The data imply that isovaltral obtained from isovaltrate possess marked effects than its parent compound. The cytotoxic effect got reduced on storage as valepotriates decomposed, which showed weak cytotoxic effect [19].

An in vitro cytotoxic study was conducted on three valepotriate isomers, jataman at valtrates Z, Z, and Z obtained from the air-dried whole plants of V. jatamansi (syn. V. wallichii). It was found that methanolic extracts exhibited moderate cytotoxicity using MTT assay against the metastatic prostate cancer (PC3M), lung adenocarcinoma (A549), hepatoma (Bel7402) cell lines, and colon cancer (HCT-8) [19]. The viability of cell lines was measured by observing absorbance at 570 nm using dimethyl sulfoxide as solvent and paclitaxel as positive control with IC50 values of 2.8–8.3 µM [20].

Earlier studies reported that 10 new valepotriates, jataman valtrates (1–10), and one already reported compound, nordostachinin (11), were extracted and isolated from the entire plants of V. jatamansi. 10 new valepotriates, jataman valtrates, compound’s structures were determined by spectroscopic analysis and were subjected to access cytotoxic action against PC-3M cell lines, and SAR studies were performed and tested for all the above compounds and the in vitro cytotoxic data report proposed some particular functional groups were related to the cytotoxic activity against PC-3M cell lines. An unsaturated bond at C1-C2, the oxirane ring, and at C10 the chlorine atom presence in the valepotriates was highlighted as important structural features [21].

Anxiolytic activity
The anxiolytic properties of compound were explored from radix and rhizomes of Valeriana jatamansi in mice [22]. Male ICR mice were treated with compound V. jatamansi at dose levels of 1.2 g/kg, 2.4 g/kg, and 4.8 g/kg and saline, diazepam 2 mg/kg given orally for 10 days and then examined using techniques elevated plus maze plus (EPM) and light-dark box (LDB) [23,24]. The compounds effect on mice was spontaneous and tested by locomotor activity test. At higher doses, the mice demonstrated more number of entries and additional time was spent on the open arms of EPM [25]. The transitions occurred have also increased, and similarly, more time was spent in light compartment of the LDB. Treatment with flumazenil significantly reduces anxiolytic-like effects, but treatment with V. jatamansi did not affect the spontaneous activity in mice. The study concludes that the compound V. jatamansi can be a better choice for anxiolytic action than producing sedation [26,27].

In another study, the anxiolytic activity was performed using iroididfraction which was obtained from rhizomes and radix of V. jatamansi using D-101 resin. The major constituents were analyzed preliminarily by TLC, UV spectrophotometry, and HPLC, and its anxiostasy effects at 6 mg/kg, 9 mg/kg, and 12 mg/kg were evaluated using elevated plus maze technique, the Vogel's drinking conflict technique, the open field drink technique, and the LDB technique. The mechanism of action was investigated using ELISA technique and it was shown by regulating GABA level [28].

Neuroprotective effect
Neuroprotective activity of methanolic extract of the dried roots of V. jatamansi was explored using three new iridoids, valerianoids
A-C[1-3], and other three already reported compounds [4-6] were subjected against (MPP+) 1-methyl-4-phenylpyridinium-induced neuronal apoptosis in the neuroblastoma of dopaminergic SH-SY5Y utilizing guanosine as positive control. Among these six compounds, four had showed moderate neuroprotective effect [Fig. 1] [8,29].

CONCLUSION

*V. jatamansi* is a perennial herb grown in higher altitude regions of Himalayas is an important ayurvedic plant with multiple remedies. The roots and rhizomes of the plant had substantial number of chemical constituents such as iridoids, lignoids, valerandoids, and valepotriates which are used to treat various ailments. Pharmacologically, it has been evaluated for *in vitro* antimicrobial, antioxidant, and anticancer activity and *in vivo* anti-inflammatory, anxiolytic, and neuroprotective activity [30]. However, continuous research on this plant is needed to know the exact molecular mechanism, and further, elaborative studies can lead to develop safe therapeutic use in modern medicine.

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