# Review on Fern Marsilea Minuta Linn (Marsileaceae)

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Abstract- Marsilea minuta Linn. is a fern belongs to the family Marsileaceae. The plant is distributed throughout India. According to Acharya Charak and Susruth it possess tridosaghan property and grahi in nature. The synonyms of the plant are Sitivara and Svastika. The chemical constituent marsilene, a macrocyclic ketone has been isolated from the plant which possesses sedative and anti-convulsant properties. The plant has been studied for their various pharmacological activities like adaptogenic-antistress, anti-depressant, anti-diabetic, antiaggressive, anti-fertility, anti-tussive, hepatoprotective, analgesic and hypocholesterolemic activity. Ethno botanically the plant is important as it is used in the treatment of diabetes by local people in Javadhu Hills Tamil Nadu, India. Though, systemic information on various aspects of this species is unavailable. In present review, an attempt has been made to present the information regarding plant profile, pharmacological properties and ethno botany.

Keywords- Marsilea minuta Linn., tridosaghna, marsilene, antidepressant, ethnobotanically.

#### I. INTODUCTION

*Marsilea minuta* Linn. is a fern and grows in less moist areas commonly in rice and wheat fields of tropical India. The rhizome is aerial, slender, creeping below the surface of the soil. Roots are borne at the nodes. The young leaves are circinate, four leaflets terminate the petiole, and the leaflets are folded together, till maturity. Leaflets–entire or crenate. Pedicels 2-6, are basal, slightly connate or free. It is cooling, light to digest, appetizer and promotes sleep. <sup>[1]</sup>

## **II. GENUS DESCRIPTION**

*Marsilea* is represented by 53 living and 10 fossil species. The genus *Marsilea* with about 60-65 species is cosmopolitan distribution mainly in warmer parts of the world such as tropical Africa and Australia. About 9 living species have been recorded from India, of them four from Rajasthan. Of these *Marsilea minuta* Linn. is the commonest and *M. brachypus*, *M. quadrifolia*, *M. rajasthanensis* and *M. aegyptiaca* are the other important Indian species. <sup>[2, 3, 4]</sup>

## III. FAMILY FEATURE

Aquatic or marsh plants with slender creeping rhizomes, growing in mud, the leaf with blades (when present) often floating on surface of water and petioles arising from rootstocks, the blades simple or with 2 or 4 pinnae, fanshaped, the veins dichotomous and anostomosing at margin; monoecious, producing megasporangia plants and microsporangia; the sporocarps hard and bean-shaped, borne on the petioles laterally or at their bases, stalked, solitary or numerous. Morphologically, the sporocarps are a modified leaf segment, folded together, containing 2 rows of indusiated sori within. Megasporangia produce megaspores which on germination give rise to egg cells, while the microsporangia produce microspores that give rise to spermproducing antheridia.

#### IV. SCIENTIFIC CLASSIFICATION

The Taxonomical Classification of *M. minuta* Linn. is as follows:

Marsilea minuta L., Mant.pl.Aitera 308.1771[Oct 1771]

Kingdom : Plantae

- Phylum : Polypodiophyta
- Division : Pteridophyta
- Class : Equisetopsida C. Agardh

Subclass : Polypodiidae Cronquist, Takht. & W. Zimm.

- Order : Marsileales
- Family : Marsileaceae Mirb.
- Genus : Marsilea
- Species : *minuta* L. <sup>[5, 6, 7, 8]</sup>

# V. BOTANICAL DESCRIPTION

The plant *Marsilea minuta* L. belonging to the family Marsileaceae is a creeping herb with long, slender rhizomes; leaves alternate, in two rows on the rhizome. Petiole long, slender, with the four obovate–retuse, glabrous leaflets at the tip, arranged in a whorl. Flowers absent. Fructification (sporocarps) produced during summer, dark brown, hard and bean–shaped with two unequal horns. <sup>[9]</sup>

Plant is an aquatic leptosporangiate fern (pteridophyte) which is considered to be highly advanced among pteridophytes for as heterospory and specialization of ganteophytes. Small herbaceous plant with trailing habit shows profuse vegetative growth producing largest nodes in the rhizome. <sup>[10]</sup> (Figure 1).



Fig. 5.1 M. minuta Linn. in rice field

## VI. MEDICINAL USES

The ethanolic extract of *M. minuta* Linn. (Whole plant) produced CNS depressant effects and hypothermia in mice. The aqueous and alcoholic extracts of defatted and fresh leaves of *M. minuta* Linn. was proved effective for anti convulsant and sedative activities.

The microbiological studies on *Marsilea (Sunisannaka)* have been carried out. Marsilea leaves extract showed a mild degree of antifungal activity against *Alternaria alpandi*, *Fusarium nivale, Gleocladium, Phomopsis and Gibberella spp.* Optium antibacterial activity was reported against *Bacillus anthracis, B. pumilus, B. subtilis, Salmonella paratyphi, Vibrio cholera, Xanth. Campestris and Xanth. Malvacearum.*<sup>[10]</sup>

## VII. HISTORICAL REVIEW

1. In Charak Samhita, Sunisannak (Chaupatiya) i.e. Marsilea minuta Linn. is grahi in nature and has tridosaghan property. [11]

2. In *Susruth Samhita*, Susruta has described *Sunisannak* in *Sak Varga*. *Sunisannak* has *tridosanasak* property and helpful in burning sensation and dysentery.<sup>[12]</sup>

3. In *Raj Nighantu, Sunisannak* has been described in *Satahadi Varga*.<sup>[13]</sup>

4. Classical texts describe this plant as a vegetable (*Saka*) and as having four leaflets (*Catuspatri, Caturdalah*), which resemble those of *Cangeri*, i.e. *Oxalis corniculata* (Vaidya Bapalal, 1982: 212). This has been universally accepted as belonging to the genus *Marsilea* (Marsileaceae) and the ancient descriptions compare well this plant. <sup>[14]</sup>

5. However, most of the authors have described it under the name *M. quadrifolia*, the taxonomy of which is in some confusion. <sup>[15]</sup>

6. In a recent paper, *Bharadwaja* (1980) has concluded that the Indian specimens have wrongly been identified as *M. quadrifolia*. The Indian specimens belong to *M. minuta* with which some authors have equated this drug (Singh & Chunekar, 1972:436; Chunekar, 1982:674; Sharma, 1983: 535). <sup>[16]</sup>

7. In *Shankar Nighantu* (1935 AD), *Sunisannak* is harmful for kidney and intestine. The antidote for this is the gum of babul. <sup>[17]</sup>

8. In *Nighantu Adarsh* (1928 AD) the leaves paste of *Sunisannak* is helpful in curing wounds.<sup>[18]</sup>

9. In *Dravyaguna Vijanan* (1997 AD), it is found throughout India, mainly near canals and marshy places. <sup>[19]</sup>

10. *Dalhana* has quoted Brahmadeva while describing the plant as follows:

Sunisanna plant has leaves like Changeri plant, leaves are 4 in number, it is growing in water or watery places. It is known as 'Catuspatri' (plant with 4 leaves).

Dalhana, here, has identified the plant thus:

A plant with 4 leaves. Others say *Sunisanna* is *Siravalika* plant. <sup>[20]</sup>

# VIII. PHARMACOGNOSTICAL REVIEW

1. Distribution: This plant is an emergent aquatic fern commonly found in marshy and shady places by the side of canals and rivers and also in the low flooded rice fields of West Bengal. The Genus Marsilea species are of wide and almost cosmopolitan distribution except for limited occurrence in some areas. <sup>[31, 32]</sup>

2. Synonyms: Sitivara, Svastika, Sunisannaka, Srivaraka, Sucipatra, Parnaka, Kukkuta and Sikhi.<sup>[33]</sup>

- 3. Vernacular names :
  - Sanskrit- Sunnisannaka, Catuspatri
  - Hindi- Choupatiya, Sunasuniya
  - Bengali- Susani Shak
  - Tamil- Arai-kirai
  - Telegu- Mudugo-tamara
  - Malayalam- Chitigina Soppu
  - Kannar- Papalu
  - English- Water clover, pepperwort. <sup>[34]</sup>
- 4. Rasa Pancak (Pharmacodynamics)
  - ✓ Rasa- Kasaya (Astringent), Madhura (Sweet)
  - ✓ Guna-Laghu (Light), Snigdha (Unctuous)
  - ✓ Virya- Sita (Cold)
  - ✓ Vipaka- Katu (Pungent)
  - ✓ *Dosakarma- Tridosaghna* (that which alleviates all three dosas). <sup>[35]</sup>

5. Properties and actions:

- Karma- Arsoghan (alleviates piles), Dipana-grahi (Appetiser- Antilaxative), Raktosodhana (Blood Purifier), Kasahara (Destroys cough), Vrsya (Aphrodisiac), Visaghna (Removes poison), Medhya- nidrajanana- vedanahara (Promotes retentive intelligence- Induces sleep- Alleviates pain), Caksusya (Beneficial for the vision). <sup>[36]</sup>
- Roga- Arsa (Piles), Vatarakta- urustambha (Gout-Stiffness in the thigh muscles), Agnimandya-grahani (Digestive impairment- Malabsorption syndrome), Raktavikara (Disorders of blood), Vataja kasa-svasa (Cough due to vata dosa-Dyspnoea), Sukraksaya (Deficiency of semen), Visa (Poison), Timiraroga (Cataract), Manasaroga-nidranasa (Mental diseases-Insomnia). [37]
- 6. Therapeutic uses

The plant drug is used as nervine tonic in treatment of epilepsy and insomnia. The leaves are used as a remedy in carbuncle in thigh. Leaves roasted in ghee are used in bilious affections. The mature spores with butter milk recover urinary troubles. The plant acts as anti venom drug.

The whole plant of drug *Sunisannaka* is ground and pasted over wounds. Drug is suggested to be wholesome to protect and promote eye-sight. The vegetable of herb (*Sunisannaka Saka*) is fried in ghee (butter) and given in intrinsic haemorrhage (*raktapitta*) as *Sunisannaka belongs* to a group of vegetables wholesome (*pathyasaka*) in *Raktapitta* diseases. In condition of *Urustambha*, the vegetable of drug plant *Sunisannaka* is cooked in water and oil, without salt, and same is prescribed in diet. The drug plant is a major ingredient drug in *Sunisannaka cangeri ghrita* prescribed in management of piles or haemorrhoids (*arsa*). Stalks of leaves eaten as a pot-herb especially in times of scarcity. *Sunisannaka* does not cause heartburn, checks flow through channels. Settles three dosas. <sup>[38, 39, 40]</sup>

## IX. PHYTOCHEMICAL REVIEW

1. Chemical constituents: Marsilene, a macrocyclic Ketone with sedative and anti convulsant properties. <sup>[41]</sup> (Table 1).

2. Phytochemical studies on Marsilea minuta L. [42] The crude extract of *M. minuta* illustrated diverse phyto-profile with reference to solvents of the plant extracts. The phenol is present in all the tested extracts i.e., petroleum ether, chloroform, acetone, benzene and aqueous extracts of M. minuta. The flavonoid is present in petroleum ether, acetone and benzene. The tannin showed its present in all the extracts of M. minuta except ethanol. The coumarin and carbohydrates are present in the chloroform, acetone, benzene and aqueous extracts of M. minuta. The steroid is present in chloroform, petroleum ether, aqueous and ethanol and saponin in petroleum ether, benzene and chloroform extracts of M. minuta. Xanthoproteins is present in petroleum ether, acetone and aqueous extracts of M. minuta. The chloroform and petroleum ether extracts of M. minuta showed the presence of proteins.

# X. PHARMACOLOGICAL REVIEW

1. *M. minuta* Linn. has a wide range of Pharmacological activities. It is traditional used to promote appetite overcome the three dosas – *vata*, *pitta* and *kapha*-excess fat, fever, diabetes, leprosy and other skin diseases. The drug induces sleep and is used in mental and nervous disorders. It is an aphrodisiac; it purifies blood and cures cough,

haematological diseases, dyspepsia, piles and poisons. (Chunekar, 1982:674; Sharma, 1983:535).<sup>[43]</sup>

The various reported Pharmacological activities of the plant highlight the therapeutic potential of *M. minuta* Linn.

2. Adaptogenic Anti-stress Activity <sup>[44]</sup> *M. minuta* adaptogenic anti stress activity as shown by its mitigating effects on several chronic stress induced physiological and behaviour perturbations, comparable to that induced by the well accepted adaptogenic agent, *Panax ginseng*.

3. Antidepressant Activity <sup>[45]</sup> The anti-depressant effect exhibited by *Marsilea minuta* extract may be due to its effect on 5-HT2A density in rat frontal cortex. Anti-depressant activity was studied using forced swimming test (FST), tail suspension test (TST), learned helplessness test (LHT) and 5hydroxytryptophan (5-HTP) induced head twitches response in rodents. Immobility time in FST and TST significantly P (<0.05) reduced by ethanol extract of *Marsilea minuta* treated animals. A decrease in number of escape failures in LHT was also observed in *Marsilea minuta* treated rats. Head twitch response induced by 5HTP was significantly attenuated by *Marsilea minuta* (400 mg/kg, p.o.) and imipramine showing the involvement of serotonergic system.

4. Antidiabetic Activity [46] Marsilea minuta has significant Antidiabetic activity when compared with standard drug Glibenclamide. The study was performed on ethanolic extract of MM leaf in oral glucose tolerance test (OGTT) and alloxan-induced diabetes models in albino rats. Three weeks treatment of diabetic animals with EEMM (250 and 500 mg/kg) showed significant check in rise of blood glucose compared to undertreated diabetic rats along with improved complete lipid profile. The fasting blood glucose, cholesterol, HDL cholesterol and serum triglyceride content were found to be significantly reduced (P<0.05) in EEMM treated rats and the extract also showed the potent elevation in the level of serum HDL cholesterol. On the basis of analysis of data obtained during the study, it may be concluded that EEMM leaf is having significant antihyperglycemic potential and can be further fractionated in order to get a responsible constituent for this very action.

5. Anti-aggressive Activity <sup>[47]</sup> The standardized extract of *M. minuta* was evaluated for its potential effects against defensive and offensive aggression behaviour models of rodents. *M. minuta* extract was orally administered at three dose levels (100, 200 and 400 mg/kg BW) once daily for 14 consecutive days as a suspension in Polyethylene Glycol (PEG), diazepam (2.5 mg/kg, p.o.) was used as a standard

anti-aggressive agent. Control groups animals were given an equal volume of vehicle (10% v/v PEG Suspension). Antiaggressive activity was evaluated using the following validated models of aggression, viz: foot shock-induced aggression, isolation-induced aggression and resident intruder aggression, in rodents. The results show that the extract from *M. minuta* has a promising anti-aggressive activity qualitatively comparable to that of diazepam.

6. Anti-fertility Activity <sup>[48]</sup> The methanol extract of *M*. *minuta* was found to produce significant elevation of the level of total cholesterol and ascorbic acid content of the ovaries of the treated female swiss albino mice. It was showed that a significant reduction produces in the activities of glucose 6 phosphate dehydrogenase enzymes and  $\Delta^5$ -3- $\beta$ hydrooxysteroid dehydrogenase enzymes in mice. The results show that the methanol extract produced anti-fertility activity in mice, which may be due to inhibition of gonadal steroidogenesis.

7. Anti-tussive, expectorant activity <sup>[49]</sup> The anti-tussive activity of M. minuta methanol, ethyl acetate, and petroleum ether extracts was evaluated using ammonia and sulphur dioxide induced mice coughing. The expectorant activity was evaluated by the volume of phenol red in mice's tracheas. Extracts significantly increased mice's cough latent period and inhibited the frequency of cough induced by ammonia and sulphur dioxide, and improved tracheal phenol red output in expectorant evaluation. Methanol extract produced the highest activity in all tested models. Methanol extract at 500 mg/kg showed 59.5% and 55.8% inhibition in the number of coughing induced by ammonium liquor and SO<sub>2</sub>, respectively, while it showed 89.3% increase in phenol red secretion at the same dose, which showed superior activity compared to other extracts. The present study provided evidence for M. minuta to be used as an anti-tussive and expectorant in Indian folk medicine.

8. Biological Activity <sup>[50]</sup> The anti-bacterial activity of ethanolic extract of *M. minuta* by disc diffusion method and the concentration of cadmium and chromium was determined by using the atomic absorption spectroscopy. The metal contents were extracted from the plants and crude extract by using wet digestion process.

9. Hepatoprotective Activity <sup>[51]</sup> The methanolic extract of *M. minuta* has hepatoprotective and anti-hepatotoxic effects in  $CCl_4$  induced hepatotoxicity in rats by its ability to stabilized cell membrane, which may be due to its antioxidant property by in vitro DPPH assay. MMME at 100, 200, 400 mg/kg b.w.p.o showed a significant decrease in serum bilirubin levels suggesting the possibility of the extract's ability to repair the damage of the hepatocytes caused by  $CCl_4$  in prophylactic and curative studies.

10. Hypocholesterolemic Activity <sup>[52]</sup> Feeding of a *Marsilea minuta* leaf extract (Fr.L) reduced serum cholesterol and triglycerides by 31 and 63% respectively, in athero diet fed gerbils. Liver cholesterol and triglycerides were also lowered by 71 and 27% respectively, in comparison to athero fed controls. Moreover, treatment with Fr.I prevented the accumulation of cholesterol and triglycerides in liver and aorta and was able to dissolve atheromatous plaques of thoracic and abdominal aorta. Faecal excretions of cholesterol and triglycerides were significantly increased in Fr.I fed gerbils.

11. Locomotor and Analgesic activity <sup>[53]</sup> *M. minuta* was used in ancient days as valiya varahyrdighrtam to produce sedative action and to control the vatadosha's induced pain. In the study locomotor and analgesic activity of different extracts of aerial parts of *Marsilea minuta* were evaluated in albino mice. The extracts produce significant locomotor and analgesic activity when compared with standard drugsdiazepam and pentazocine respectively.

## XI. MISCELLANEOUS

1. Structure of Marsileagenin A: A new hexahydroxy triterpene from *Marsilea minuta* Linn. <sup>[54]</sup> The crude saponin obtainable from *Marsilea minuta* Linn. on acid hydrolysis yielded a mixture of Sapogenols. The major sapogenol named Marsileagenin A was found to be a hexahydroxy triterpene of oleanene series. From a study of various Spectrometric data together with chemical reactions the structure of this sapogenol has been assigned as olean-12-ene- $2\alpha$ ,  $3\beta$ ,  $16\beta$ ,  $21\beta$ ,  $22\alpha$ , 28-hexol ( $1\alpha$ ).

2. Ultrastructural and biochemical effects of cadmium on the aquatic fern *Marsilea minuta* Linn. <sup>[55]</sup> The uptake of cadmium by cultured vegetative clones of the aquatic fern *Marsilea minuta* Linn. was studied in a static experimental bioassay system as functions of dose and period of exposure. The pathomorphological manifestations of Cd toxicity, as studied by transmission electron microscope, indicated damage to chloroplasts and tonoplasts as well as electron opaque granular deposits. Preliminary evidence based on molecular sieving Chromatography showed the formation of two cadmium binding proteins of 78 and 33kDa in the leaf tissue under cadmium stress.

## XII. ETHNOBOTANICAL REVIEW

Description of Ethno botanical Uses of *M. minuta* Linn. in different localities are enlisted. (Table 2).

#### XIII. CONCLUSION

The plant has long being investigated for its pharmacological activities supporting its vast ethno botanical and alternative medicinal use. The plant has been reported extensively as anti-depressant, anti-stress, anxiolytic, anti-fertility and anti-tussive agent. Though diseases treated indigenously using the plant has not been confirmed in the laboratory so this leaves an opportunity to explore the species both phytochemically and pharmacologically. Ethno pharmacology can bridge between the folklore use and actual pharmacological efficacy of medicinal plant. Therefore it may be used in novel drug discovery in near future.

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## REFERENCES

1. Chunekar KC. Bhavaprakasa of Bhavamisra (Original text along with commentary and translation including Nighantu Portion), Vol-I, 1<sup>st</sup> ed. Chaukhamba Orientalia, Varanasi, 2006: 452.

2. Vashishta PC, Sinha AK, Kumar Anil. Botany for Degree Students Pteridophyta (Vascular Cryptogams), S. Chand & Company Ltd., 1971: 523-24.

3. Rashid A. An Introduction to Pteridophyta. Diversity. Development. Differentiation, Vikas Publishing House Pvt Ltd., 2010: 143.

4. Lawrence George HM. Taxonomy of Vascular Plants. New Delhi: Oxford & IBH Publishing Co. Pvt Ltd., 353.

5. The plant list [http:www.theplantlist.org/tpl/search?q=marsilea+minuta] [cited 2013 Feb 18].

6.Tropicos[http://www.tropicos.org/Name/26604296] [cited 2012 Dec 26].

7. www.iucnredlist.org [cited 2012 Dec 27].

8. Vashishta PC, Sinha AK, Kumar Anil. Botany for Degree Students Pteridophyta (Vascular Cryptogams), S. Chand & Company Ltd., 1971: 550.

9. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co.Pvt.Ltd., 1994: 455.

10. Pandey Gyanendra. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541-45.

11. Charak, Charak Samhita, Sutra Sthana, 27/88, edited by Dr Brahmananda Tripathi, Chaukhamba Surbharati Prakashan, Varanasi, 2005: 511.

12. Susruta, Susruta Samhita, Sutra Sthana, 46/262, 265, edited by Kaviraja Ambikadutta Shastri, Chaukhambha Sanskrit Sansthan, 2005: 206.

13. Pandit Narahari, Raja Nighantu, 50-52, edited by Dr Indradeva Tripathi & Acharya Viswanath Dwivedi, Chowkhamba Krishnadas Academy, Varanasi, 2006: 71-72.

14. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co. Pvt. Ltd., 1994: 455.

15. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co. Pvt. Ltd., 1994: 455.

16. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co. Pvt. Ltd., 1994: 455.

17. Rajvaidya P, Gaud Shankar Dutt, Shankar Nighantu, Chakumbha Oriental, Varanasi, 2002: 25.

18. Bapalal Vaidya, Nighantu Adarsh, Vol-II, 776, Chaukhambha Bharti Academy, Varanasi, 2005: 775.

19. Sharma PV. Dravyaguna vijnana, Vol-II (vegetable drugs), Chaukhambha Bharati Academy; 535.

20. Bapalal Vaidya. Some Controversial Drugs in India Medicine, Chaukhambha Orientalia, Varanasi, 2005: 212-13.

21. Charak, Charak Samhita, Sutra Sthana, 27/88, edited by Dr Brahmananda Tripathi, Chaukhamba Surbharati Prakashan, Varanasi, 2005: 511. 22. Susruta, Susruta Samhita, Sutra Sthana, 46/262, 265, edited by Kaviraja Ambikadutta Shastri, Chaukhambha Sanskrit Sansthan, 2005: 206.

23. Astanga samgraha, Astanga Samgraha, Sutra Sthana, 5/113-114, edited with Saroj' Hindi commentary by Dr Tripathi Dutt Ravi, Chaukhambha Sanskrit Pratishthan, Delhi, 2003: 136.

24. Dhanvantari, Dhanvantari Nighantu, 68/151-152, editedby Sharma PV, Chaukhambha Orientalia, Varanasi, 2005:44.

25. Bhavamisra, Bhavaprakasa Nighantu, 19/29, 31, edited by Dr GS Pandey, Chaukhambha Bharati Academy, Varanasi, 2004.

26. Pandit Narahari, Raja Nighantu, 50-52, edited by Dr Indradeva Tripathi & Acharya Viswanatha Dwivedi, Chowkhamba Krishnadas Academy, Varanasi, 2006: 71-72.

27. Sharma PV. Priyanighantu, Chaukhamba Surbharati Prakashan, Varanasi, 2004: 98.

28. Mishra Umapati. Dravyaguna Sangraha, Chaukhamba Surbharati Prakashan, Varanasi, 1995: 68.

29. Bapalal Vaidya. Some Controversial Drugs in India Medicine, Chaukhambha Orientalia, Varanasi, 2005: 212-213.

30. Bapalal Vaidya. Some Controversial Drugs in India Medicine, Chaukhambha Orientalia, Varanasi, 2005: 212-213.

31. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co. Pvt. Ltd., 1994: 455.

32. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

33. Chunekar KC, Bhavaprakasa of Bhavamisra (Original text along with commentary and translation Including Nighantu Portion), Vol-I, 1<sup>st</sup> ed. Chaukhamba Orientalia, Varanasi, 2006: 452.

34. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

35. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

36. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

37. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

38. Gyanendra Pandey. Dravyaguna Vijnana, (Materia Medica-Vegetable drugs) (English- Sanskrit), Vol-III, Chowkhamba Krishnadas Academy, Varanasi, 2004: 541.

39. Anonymous. The Useful Plants of India, NISCAIR, New Delhi, 2005: 357.

40. Valiathan MS. The Legacy of Susruta. Orient Longman, 2007: 190.

41. Chunekar KC, Bhavaprakasa of Bhavamisra (Original text along with commentary and translation Including Nighantu Portion), Vol-I, 1<sup>st</sup> edition, Chaukhamba Orientalia, Varanasi, 2006: 452.

42. Muraleedharannair JM, Johnson MA, Mony M, Zachariah MP, Solomon J. Phytochemical studies on *Azolla pinnata* R.Br., *Marsilea minuta* L. And *Salvinia molesta* Mitch. Asian Pac J Trop Biomed 2011; 1(1): 526-29.

43. Sivarajan VV, Balachandran Indira. Ayurvedic Drugs and their Plant Sources. New Delhi: Oxford & IBH Publishing Co. Pvt. Ltd., 1994: 455.

44. Tiwari OP, Bhattamisra SK, Singh PN, Kumar Vikas. Adaptogenic Anti stress Activity of Standard Extract of *Marsilea minuta, Pharmacologyonline 2009;* 1: 290-99.

45. Bhattamisra SK, Khanna VK, Agrawal AK, Singh PN, Singh SK. Antidepressant activity of standardised extract of *Marsilea minuta* Linn. J Ethnopharmacol 2008; 117(1): 51-57.

46. Madhu S, Kannabirran V, Frank PRoyal, Reddy MSathish, Gnanasekar N, Pandurangan Kamalakshi. Evaluation of Antidiabetic Activity of *Marsilea minuta* Linn. Against Alloxan Induced Diabetes in Albino Rats. IRJP 2012; 3(8): 223-25.

47. Tiwari Om P, Bhattamisra Subhrata K, Tripathi Pushpendra K, Singh Paras N. Anti-aggressive activity of a standardized extract of *Marsilea minuta* Linn. in rodent models of aggression. *BioSci Trends* 2010; 4(4):190-94.

48. Gupta Malaya, Mazumder VK, Datta Ipsita, Bhattacharya S, Mukherjee S, Manikandan L. Studies of Antifertility Activity of *Marsilea minuta* Linn. Indian J. Pharm. Sci. 2002; 64(2): 176-78.

49. Chakraborty R, De B, Devanna N, Sen S. Antitussive expectorant activity of *Marsilea minuta* L., an Indian vegetable. J Adv Pharm Tech Res 2013; 4: 61-4.

50. Zahid Hussain, Khalid Mohammad Khan, Shahnaz Perveen, Nida Ambreen, Wajid Rahman, Ata Ullah. The Effect of Cadmium and Chromium Concentration, on Biological Activity of *Marsilea minuta*. J. Chem. Soc. Pak 2011; 33(6): 874-76.

51. Praneetha P, Rani V Swaroopa, Kumar B Ravi. Hepatoprotective Activity of Methanolic Extract of Leaves of *Marsilea minuta* Linn. Against CCl Induced Hepatic Damage in Rats. G JP 2011; 5(3): 164 -71.

52. Gupta RS, Kumar Pramod, Sharma Anita, Bharadwaj TN, Dixit VP. Hypocholesterolemic Activity of *Marsilea minuta* in gerbils. Fitoterapia 2000; 71(2): 113-17.

53. Nagavalli D et al., Locomotor and analgesic activity of different extract of aerial parts of *Marsilea minuta* Linn. Hamdard Med. 2009; 52(2): 86-8.

54. Chakravarti D, Debnath NB, Mahato SB, Chakravarti RN. Structure of Marsileagenin A: A new hexahydroxy triterpene from *Marsilea minuta* Linn. Tetrahedron 1975; 31(15): 1781-82.

55. Singh J, Devi S, Chawl G, Gupta M, Viswanathan PN. Ultrastructure and biochemical effects of Cadmium on the aquatic fern *Marsilea minuta* Linn. Ecotox Environ Safe 1991; 21(2): 171-81.

56. Mustafa Kamal, Sultan Mehmood Wazir, Muhammad Hassan, M.Subhan Saad Ullah Khan, Asim Muhammad, Shaheen Taj. Ethnobotanically Important Plants of district Bannu, Pakistan. Pak. J. Pl. Sci. 2009; 15(2):87-93.

57. Thirumalai T, David Beverly C, Sathiyaraj K, Senthikumar B, David E. Ethnobotanical Study of Antidiabetic medicinal Plants used by the local people in Javadhu Hills Tamil Nadu, India. Asian Pac J Trop Biomed. 2012: S910-S913.

58. Upreti Kanchan, Jalal Jeewan S, Tewari Lalit M, Joshi GC, Pangtey YPS, Tewari Geeta. Ethnomedicinal Uses of Pteridophytes of Kumaun Himalaya, Uttarakhand, India. J Ame Sci. 2009; 5(4): 167-70.

59. Mohammed Rahmatullah, AAB Tajbilur Kabir, Md. Mahfuzur Rahman, Md. Shahadat Hossan, Zubaida Khatun, Mst Afsana Khatun, Rownak Jahan. Ethnomedicinal Practices among a Minority Group of Christians Residing in Mirzapur Village of Dinajpur District, Bangladesh. Adv. in. Nat. Appl. Sci. 2010; 4(1): 45-51.

60. Nicole Bultrushes. Medical Ethnobotany Phytochemisry and Bioactivity of the Ferns of Moorea, French Polynesia.

61. Singh Balendra Pratap, Upadhaya Ravi. Ethno-botanical Importance of Pteridophytes used by the tribe of Pachmarhi, Central India. J Med Plants Res. 2012; 6(1):14-18.

62. Munisamy Anbarashan, Narayanaswamy Parthasarthy, Anbarashan Padmavathy. Ethno-floristic Survey in sacred groves Pudukottai district, Tamil Nadu-India. J Med Plants Res. 2011; 5(3): 439-43.

63. Bharti Malay. Ethno Medicinal Importance of Some Common Pteridophytes Used By Tribals of Ranchi and Latehar District Of Jharkhand, India. Socioscan: An International Quaterly Journal of Ethno and Social Sciences 2011; 3(1&2): 5-8.

64. Kumar Kaushal, Abbas SG. Ethnomedicinal Composition depends on Floristic Composition: A case studied in Sal Forests of Jharkhand. International Journal of Pharmacy and Life Sciences 2012; 3(5):1710-19.

65. Rout SD, Panda T and Mishra N. Ethnomedicinal Studies on some Pteridophytes of Similipal Biosphere Reserve, Orissa, India. International Journal of Medicine and Medical Sciences. 2009; 1: 192-97.

66. Pathak A, Singh A, Singh AP. Ethnomedicinal uses of Pteridophytes of Vindhyan Region (M.P). International Journal of Pharmacy & Life Sciences 2011; 2(1): 496-98.

67. Lakshminarasimhan P, Sharma BD. Flora of Nasik district, (1991), Botanical Survey of India. 596.

68. Michael Woods, Alvin R Diamond, Jr. Pteridophytes of Southeast Alabama Dichotomous Keys, Illustrations and Distribution Maps. J Ala Acad Sci. 2008; 79(3-4): 204.

69. Ghosh SR, Ghosh B, Biswas Anjali, Ghosh RK. The Pteridophytic Flora of Eastern India. (2004), Botanical Survey of India, Kolkata, Vol-1: 187-89.

70. Sundas Iltaf, Zaheer-ud-din Khan, Noreen Riaz. A Contribution to the taxonomic study of fern flora of Punjab, Pak. J. Bot. 2012; 44: 315-22.

71. Patil S. Pteridophytes of Chandgad forest of Kolhapur district (Maharashtra), ISRJ Vol-1, (2011): Botany.

72. Shrivastava Neeles, Sinha Sharma. A. Tribes of Surguja and Korea region in Chattisgarh. JPR 2011; 4(9): 3081-82.

73. Mallik Bikram K, Panda Tribhuban, Padhy Rabindra N. Traditional Herbal Practices by the Ethnic People of Kalahandi District of Odisha, India. Asian Pac J Trop Biomed. 2012; S988-S994.

74. Parihar P, Parihar L. Some Pteridophytes of medicinal importance from Rajasthan. NPR. 2006; 5: 297-301.

Parts	Active Constituents
Alcoholic extract of <i>M. minuta</i> Linn.	A saponin (a mixture of sapogenols on hydrolysis).
	Marsileagenin A, the major sapogenol was found to be olean-12-
	ente-2a, 2B, 16B, 21a, 22a, 28-hexol whereas the other two
	sapogenols viz. Marsileagenins B and C were present in small
	quantities. <sup>[10]</sup>
Chloroform extract of leaves of <i>M. minuta</i>	Marsilin <sup>[10]</sup>
Linn.	
Chloroform extract of whole plant	β- sitosterol <sup>[10]</sup>
Roots and Stems of <i>M. minuta</i> Linn.	Marsilin <sup>[10]</sup>
Petroleum ether extract of leaves of M.	An asymmetrical hydroxyketone substance (3-hydroxy-
minuta Linn.	triacontan-11-one and a mixture of secondary alcohol with
	kentriacontane-16-01. <sup>[10]</sup>
<i>M. minuta</i> Linn.	Calcium and Phosphorous <sup>[10]</sup>

Table	1:0	Chemical	Constituents	

S.	Location	Ailment treated/Properties and action
No.		
1.	Bannu District, Pakistan <sup>[56]</sup>	Methanol extract from the leaves is used to prevent accumulation of
		cholesterol and triglyceride in the liver
2.	Javadhu Hills, Tamil Nadu, India <sup>[57]</sup>	Leaf juice is used for diabetes. Local name is Aarakkerai.
3.	Kumaun Himalaya,	Plant used in cough, spastic conditions of leg muscles, in sedation and
	Uttarakhand, India <sup>[58]</sup>	insomnia. A macro cyclic ketone of sedative and convulsant properties has been isolated.
4.	Mirzapur village of Dinajpur	Juice obtained from crushed whole plant is used in gastrointestinal
	district, Bangladesh <sup>[59]</sup>	disorder.
5.	Moarea, French Polynesia <sup>[60]</sup>	Respiratory (Cough) India [Dhiman 1998, Vasudeva 1999]
		Eye diseases India (Dhiman 1998)
		Reduce cholesterol in gerbils (Gupta et al 2000).
6.	Pachmarhi, Central India	The whole part of plant is used to treat cough, spastic conditions of leg
	(Pagara, Bariam Amkhedi,	muscles, in sedation and insomnia. A macrocyclic ketone of sedative
	Neemghan, Singanama,	and convulsant properties has been isolated.
	Tekapar, Chaka and Pisua) <sup>[61]</sup>	
7.	Pudu kottai district, Tamil	The dried and powdered leaves, mixed with hot water, are taken in
	Nadu, India <sup>[62]</sup>	cases of diabetes.
8.	Ranchi & Latehar District of	Plant is used in cough, spastic conditions of leg muscles, in sedation &
	Jharkhand, India <sup>[63]</sup>	insomnia. A macrocyclic ketone of sedative and convulsant properties has been isolated.
9.	Sal forests of Jharkhand, India	The whole plant is used in body ache.
9.	[64]	The whole plant is used in body ache.
10.	Simlipal Biosphere Reserve,	Plants are used in cough, spastic condition of leg & muscle. About 10g
	Orissa, India <sup>[65]</sup>	whole fresh plant paste is mixed with 100g of crud prepared from
		cow's milk. The dosage is given orally once a day in empty stomach
		for one month against epilepsy. Younger leaves are crushed to extract
		the juice and 2 drops of juice are dropped in the nostrils of nose twice a
		day effective in migraine.
11.	Vindhyan Region, (M.P),	Plants are used in cough, spastic condition of leg muscles. Plant is
	India <sup>[66]</sup>	sweet, diuretic and ophthalmic. It is used in Psychopathy, Opthalmia,

		Diarrhoea, Leprosy, Skin diseases, Hemorrhoids and Fever.
12.	Flora of Nasik district, India	Occasional in stagnated waters Karanjul (Surgana range ).
	[67]	Sporocarps: October – April.
		Leaves are cooked and eaten as vegetable.
13.	Southeast Alabama, USA <sup>[68]</sup>	It is known dwarf water clover. It is found in Shoreline and shallow
		water along margin of beaver pond.
14.	Flora of Eastern India <sup>[69]</sup>	a)Marsilea minuta (Leaves entire or slightly crenate)
		It is found in Eastern India and other parts of India. Grows in marshy
		places along ponds, ditches, lake.
		Uses: Leaves are eaten as vegetables.
		b) Marsilea minuta var. Indica (Leaves always crenulate)
		It is found in Rajasthan, Punjab, West Bengal (India). Grows on marshy
		places near ponds, lakes, jhils.
		Uses: Leaves are edible.
15.	Fern Flora of Punjab, Pakistan	This species is just like <i>M. quadrifolia</i> , but smaller in size, its length is
	[70]	about 7.5 cm and margins are crenate.
16.	Flora of Chandgad forest of	It is a semi aquatic fern used by the native people from the hilly region
	Kolhapur distric	of Chandgad.
	(Maharashtra), India <sup>[71]</sup>	
17.	Surguja and Korea region in	Sleeping disorders, tranquilizer.
	Chattisgarh, India <sup>[72]</sup>	
18.	Kalahandi district of Odisha,	Equal amount of spores of the plant, roots of Smilax Zeylanica, root of
	India <sup>[73]</sup>	Lawsonia inermis and white coloured onion bulb is ground and the
		extracted juice is taken orally twice a day upto 7 days to cure jaundice.
		Leaves are cooked and taken to cure insomnia.
19.	Pteridophytes of Rajasthan <sup>[74]</sup>	Garasia and Bheels cook the leaves as vegetable. The decoction of
		leaves along with ginger is used to cure cough and bronchitis in many
		villages of Rajasthan.

Table 2: Ethno botanical uses of *M. minuta* Linn. around the world