

Journal of Applicable Chemistry

2014, 3 (4): 1423-1431 (International Peer Reviewed Journal)



Ficus racemosa Linn. : A Comprehensive Review

Satish A Bhalerao^{1*}, Deepa R Verma², Nikhil C Teli², Vinodkumar S Didwana² and Saurabh S Thakur²

1. Environmental Sciences Research Laboratory, Department of Botany, Wilson College, Mumbai-400 007, University of Mumbai, INDIA

2. Department of Biological Sciences, VIVA College, Virar (W)-401 303, University of Mumbai, INDIA

Email: drsatishbhalerao@yahoo.com

Accepted on 17th June 2014

ABSTRACT

Ficus racemosa Linn. syn. Ficus glomerata Roxb. (Family - Moraceae) is a large deciduous tree dispersed all over India which is generally known as Gular, Gular fig, Cluster fig or Country fig. It is a sacred tree of Hindus and Buddhists. All parts of this plant (leaves, fruits, bark, latex, and sap of the root) are medicinally significant in a variety of treatments such as diabetes, diarrohoea ulcers, stomachache, piles, skin diseases, dysentry and as carminative etc. Among various pharmacological properties, Ficus Racemosa Linn imparts vital role as anti-oxidant, anti cancer, antidiuretic, anti bacterial, antiinflammatory, memory enhancing and gastro-protective agent etc. In this review, emphasis is lead upon research associated to therapeutic properties, phytochemistry and pharmacological profile of Ficus racemosa Linn.

Keywords: Ficus racemosa Linn., Gular, antidiuretic, carminative, gastro-protective.

INTRODUCTION

Ficus racemosa (Linn) is a moderate sized avenue plant, belongs to family- Moraceae which is usually known as the Cluster Fig Tree, Indian Fig Tree or Goolar (Gular) Fig. This plant is native to Australia, Malaysia, South-East Asia and the Indian Subcontinent [1]. *Ficus racemosa* grows all over India in several forests and hilly areas. It is frequently available around the water streams and is also cultivated. Found along the river banks and inland forests from plains to 1500 m most frequently in India, Sri Lanka, Pakistan, Queensland and South China to New Guinea. The plant can be grown by vegetative as well as sexual propagation (using seeds) [2].

It is unusual in that its figs grow on or close to the tree trunk, termed cauliflory [3]. In India the tree and its fruit are called 'gular' in the north and 'atti' in the south. The fruits are a favourite staple of the common Indian macaque. In Kerala it is consider as one among nalpamara. It serves as a food plant for the caterpillars of the butterfly the Two-brand Crow (*Euploea sylvester*) of northern Australia [4]. The Ovambo people call the fruit of the Cluster Fig 'eenghwiyu' and use it to distill 'Ombike', their traditional liquor [5].

Ficus racemosa Linn (Moraceae) is an evergreen, moderate to large sized spreading, lactiferous, deciduous tree, without much prominent aerial roots. Tree about 20 m tall often with aerial roots, bark whitish-brown, smooth, Leaves grooved minutely hairy, lamina ovate-lanceolate to elliptic-lanceolate, tri-ribbed, 8-10 pairs of lateral pairs from broad to narrowly cuneate, oblique base, margin entire, acuminate at apex, glabrous on both sides, stipules triangular-ovate, brown, sub-persistent, cystoliths present only on lower side. Hypanthodia on long peduncles, borne in large clusters from tubercles on the main trunk and main leafless branches, subpyriform-globose, green, subtended by, broadly triangular-ovate brownish brads, bracts, apical orifice sunken, closed by brown bracts without internal bristles. Male flowers are sessile, ostiolar in 2-3 whorls, united, lobes dentate and stamens. Gall flowers pedicellate, dispersed among female. Female flowers are sessile or subsessile, ovary substipitate, glabrous style, stigma simple. Figs depressed subglobose or pyriform, red when ripe usually streaked. Seeds are lenticular 1 mm. Syconus fruit [6, 7].

Udumbara is considered scared to god Dattaguru. All ficus species possess latex-like material within their vasculatatures that provide defense and self healing from physical assaults [8]. This plant is universally used in traditional system of medicine for the treatment of numerous disorders. It is one of the herbs mentioned in all ancient scriptures of Ayurveda, Siddha, Unani and Homeopathy. Various plant parts such as bark, root, leaf, fruits and latex are used as astringent, vermifuge, carminative and anti-dysentery. It is a good medication for excessive appetite. The extract of fruit is used in leucoderma, menorrhagia and diabetes. It is used locally to relieve inflammation of lymphadenitis, fibrositis, skin wounds and in sprains [9].

Kingdom	Plantae
Division	Magnoliophyta
Class	Magnolipsida
Order	Urticales
Family	Moraceae
Genus	Ficus
Species	racemosa

Table 1: Taxonomic position of *Ficus racemosa* Linn.

 Table 2: Vernacular names of Ficus racemosa Linn.

Sanskrit	Yajnayoga, Sadaphalah, Brahanvrkisha, Shitavalkah, Sutah, Udumbara, Gular, Mashakin, Jantukaphalah, Jantuphalah, Krmiphalah, Vasudrumah, Saumya, Hemadugdhaka, Jantumati, Yagniyah, Audumbara	
Hindi	Pushp-hina, Pani Bhuj, Dumar, Goolar, Umari, Yajnyadumbur, Udumbara, Jantu Phal, Dharma Patra, Goolar	
English	Gular fig, Cluster Fig, Country Fig	
Bengali	Udumbara	
Telugu	Brahmamamidi, Atti, Bodda	
Gujrati	Goolar, Umbaro	
Manipuri	Heibong	
Malayalam	Atthi, Atthi Al, Aththi, Atthi-al, Udumbaram, Jantuphalam	
Marathi	Umbar, Udumbar	
Urdu	Dumar	
Oriya	Dimri	
Konkani	Rhumbud	
Kannada	Atti, Atti Mara	

Irula	Athi
Others	Goolar, Atthi, Atteeka, Athi, Crattock, Country Fig, Dumrii, Cluster Fig, Vellaiatthi, Gular Fig, Indian Fig, Redwood Fig, Rumbodo

 Table 3: Ayurvedic properties of Ficus racemosa Linn.

Rasa	Kashaya, Madhur
Guna	Guru, Ruksha
Virya	Sheet
Vipak	Katu
Doshaghnata	Kapha, Pitta Shama

 Table 4: Phytoconstituents of Ficus racemosa Linn.

Leaf	Sterols, tannins and flavonoids, triterpenoids (Lanosterol) and alkaloids. A new tetracyclic triterpene glauanol acetate which is characterized as 13α , 14β , 17β H, 20α H-lanosta-8, 22-diene-3\beta-acetate and racemosic acid were isolated from the leaves [7].
Stem-Bark	Tannin, wax, saponin gluanol acetate, β -sitosterol, leucocyanidin- $3 - O - \beta - D$ - glucopyrancoside, leucopelargonidin - $3 - O - \beta - D$ - glucopyranoside, leucopelargonidin - $3 - O - \alpha - L$ - rhamnopyranoside, lupeol, ceryl behenate, lupeol acetate, α -amyrin acetate, leucoanthocyanidin, and leucoanthocyanin from trunk bark, lauanol acetate, lupeol, β -sitosterol and stigmasterol were isolated from stem bark[10].
Trunk-Bark	Upenol, β -sistosterol and stigmasterol [10].
Fruit	Glauanol, glauanol acetate, hentriacontane, β sitosterol, glauanolacetate, glucose, tiglic acid, esters of taraxasterol, lupeolacetate, friedelin, higherhydrocarbons and other phytosterol [11].
Root	Cycloartenol, euphorbol and its hexacosanoate, taraxerone, tinyatoxin; bark euphorbol and its hexacosanate, ingenol and its triacetate, taraxerone [12].
Latex	a-amyrin, β-sitosterol, cycloartenol, cycloeuphordenol, 4-deoxyphorbol and its esters, euphol, euphorbinol, isoeuphorbol, palmitic acid, taraxerol, tinyatoxin, tirucallol, trimethyl ellagic acid [13].

Table 5: Ethnobotanical uses of Ficus racemosa Linn.

PLANT PART	USE
Fruits	Used in leprosy, diarrhoea, circulatory and respiratory disorders and menorrhagia [14-16]. Tender fruits are used as astringent, stomachic, refrigerant, in dry cough, loss of voice, diseases of kidney and spleen, astringent to bowel, styptic, tonic, useful in the treatment of leucorrhoea, blood disorder, burning sensation, fatigue, urinary discharges, leprosy, epitasis, carminative and intestinal worms. They are also useful in miscarriage, spermatorrhoea, epididymitis, cancer, myalgia, scabies, haemoptysis, intrinsic haemorrhage and extreme thirst [17, 18].
Roots	Roots are used in dysentery, pectoral complaints, and diabetes, applied in mumps, other inflammatory glandular enlargements and hydrophobia [12, 18, 19].
Bark	It is highly effective in threatened abortion and also recommended to treat Menorrhagia, leucorrhoea, gonorrhoea, urinary diseases, hemorrhage and skin diseases [20]. The bark is highly recommended in urological disorders, diabetes, hiccough, leprosy, dysentery and piles [7, 12, 18, 19].
Leaves	The leaves are excellent wash for wounds and ulcers. They are useful in dysentery and diarrhea. The infusion of bark and leaves is also employed as mouth wash to spongy gums and internally in dysentery, menorrhagia, efficient remedy in glandular swelling, abscess, chronic wounds, cervical adenitis and haemoptysis [17-19].

Latex	It is administered in haemorrhoids, boils, alleviates the edema in adenitis, parotitis, orchitis, traumatic swelling, toothache, vaginal disorders, diarrhoea particular in childrens and also aphrodisiac. Latex is applied externally on chronic infected wounds to alleviate edema, pain and to promote the healing [2]. The latex is reportedly used for treating piles [21].
Root Sap	It is used for treating diabetes [20]. The sap of this plant is a popular remedy for mumps and other inflammatory enlargements [16, 19]. In Sri Lankan indigenous system of medicine, it is used in the treatment of skeletal fracture. The Australian aborigines use this plant in the treatment of mumps, smallpox, heamaturia, menorrhagia and inflammatory conditions. In Siddha the bark, fruits and latex are used to treat constipation, anaemia and dysentery [20, 22].

APPLICATIONS

Pharmacological Profile

Anti cancer activity: *Ficus racemosa* extract at a dose of 200 and 400 mg kg⁻¹ when given orally a considerable decrease in lipid peroxidation, xanthine oxidase, γ -glutamyl transpeptidase and hydrogen peroxide (H₂O₂) generation with decrease in renal glutathione content and antioxidant enzymes generated by Potassium bromate (KBrO₃), a nephrotoxic agent that induces renal carcinoma in rats .There was considerable recovery of renal glutathione content and antioxidant enzymes. These results recommend that *Ficus racemosa* extract is a potent chemopreventive agent and suppresses KBrO₃ mediated nephrotoxicity in rats [23].

Hypolipidemic: Dietary fiber content of fruits when fed to rats in diet induced pronounced hypocholesterolemic effect, as it increased fecal excretion of cholesterol as well as bile acids. Hypolipidemic activities of ethanolic extract of bark were studied at the doses of 100-500 mg/kg b.w to alloxan-induced diabetic rats. Investigation showed that extract had potent anti-diabetic and hypolipidemic effects when compared to that of the standard reference drug, glibenclamide [24].

Anti-ulcer/Gastro-protective: Gastro-protective effect of 50% ethanolic extract of *Ficus Racemosa Linn* known as *F. glomerata* fruit (FGE) was studied in different gastric ulcer models in rats. FGE prevents the oxidative damage of gastric mucosa by blocking lipid peroxidation and by considerable decrease in superoxide dismutase, H+K+ATPase and increase in catalase activity. The H+K+ATPase are the dimeric enzyme responsible for H+ secretion by the gastric parietal cells. H+K+ATPase are selectively blocked by the action of ranitidine, an acid blocker used to treat gastric ulcers [25, 26].

Anti-oxidant activity: Ethanol extract and water extract were subjected to free radical scavenging both by steady state and time resolved methods. The ethanol extract exhibited significantly higher steady state antioxidant activity. It also exhibited concentration dependent DPPH, ABTS, hydroxyl radical and superoxide radical scavenging and inhibition of lipid peroxidation when tested with standard compounds [27-29].

Antidiuretic: The decoction of the bark of *F. racemosa* is claimed as an antidiuretic and its potential is evaluated in rats using three doses (250, 500 or 1000 mg kg⁻¹). It had a rapid onset (within 1 h), peaked at 3 h and lasted throughout the study period (5 h). It also caused a reduction in urinary Na+ level and Na+/K+ ratio, and an increase in urinary osmolarity indicating multiple mechanisms of action [30].

Hepato protective activity: Methanol extract of *Ficus racemosa* stem bark were studied using the model of hepatotoxicity induced by carbon tetrachloride (CCl_4) in rats. Pretreatment with methanol extract

resulted in significant decreases in the activities of AST, ALT and ALP, compared to CCl_4 -treated rats. The results indicate that *F. racemosa* possesses potent hepatoprotective effects against CCl_4 - induced hepatic damage in rats [31].

Antitussive: The methanol extract of stem bark was tested for its antitussive potential against a cough induced model by sulphur dioxide gas in mice. The extract exhibited maximum inhibition of 56.9% at a dose of 200 mg kg⁻¹ (p.o.) 90 min after administration [32].

Hypoglycemic: There is more than one type of hypoglycemic principles, both organic and inorganic, in Ficus racemosa Linn. fruits which produce a significant fall in blood glucose levels in normal and alloxandiabetic rabbits by producing an organotropic effect on the B-cells which results in an increased release of insulin from the pancreatic beta cells. Ficus racemosa Linn. fruits pulp may, in the long run after more detailed studies, prove to be a more valuable anti-diabetic agent as in addition to its insulin releasing and insulin-like activities. Methanol extract of powered fruits at the dose 1, 2, 3, and 4 g kg⁻¹ reduced the blood glucose level in normal and alloxan induced diabetic rabbits. On the other hand the glucose-lowering efficacy of methanol extract of the stem bark was evaluated both in normal and alloxan-induced diabetic rats at the doses of 200 and 400 mg kg⁻¹ p.o. The activity was also comparable to that of the effect produced by a standard antidiabetic agent, glibenclamide (10 mg kg⁻¹) proving its folklore claim as antidiabetic agent. The ethanol extract (250 mg kg⁻¹ day⁻¹, p.o.) lowered blood glucose level within 2 weeks in the alloxan diabetic albino rats confirming its hypoglycemic activity. β-sitosterol isolated from the stem bark was found to possess potent hypoglycemic activity when compared to other isolated compounds. Ethanolic extract of leaves lowered the blood glucose levels by 18.4 and 17.0% at 5 and 24 h, respectively, in sucrose challenged streptozotocin induced diabetic rat model at the dose of 100 mg kg⁻¹ body weight [33, 34].

Anthelmintic: The crude extracts of bark were evaluated for anthelmintic activity using adult earthworms; they exhibited a dose-dependent inhibition of spontaneous motility (paralysis) and evoked responses to pin-prick, which was comparable with that of 3% piperazine citrate. However, there was no final recovery in the case of worms treated with aqueous extract suggesting wormicidal activity [19].

Anti bacterial activity: Different extracts of leaves were tested for antibacterial potential against *Escherichia coli, Bacilus pumitis, Bacillus subtilis and Pseudomonas aureus.* Out of all extracts tested, petroleum ether extract was the most effective extract against the tested microorganism [31].

Anti-diarrhoeal: Ethanol extract of stem bark has shown significant inhibitory activity against castor oil induced diarrhea and PEG2 induced enter pooling in rats and also showed a significant reduction in gastro intestinal motility in charcoal meal test in rats which proves its efficacy as anti-diarrheal agent [35].

Anti-inflammatory: The anti-inflammatory activity of *F. racemosa* extract was evaluated on carrageenin, serotonin, histamine and dextran-induced rat hind paw edema models. The extract (400 mg kg⁻¹) exhibited maximum anti-inflammatory effect of 30.4, 32.2, 33.9 and 32.0% with carrageenin, serotonin, histamine, dextran-induced rat paw oedema, respectively. In a chronic test, the extract (400 mg kg⁻¹) showed 41.5% reduction in granuloma weight, which was comparable to that of phenylbutazone [31]. Bioassay-guided fractionation of the ethanol extract of leaves isolated racemosic acid. It showed potent inhibitory activity against COX-1 and 5-LOX in vitro with IC50 values of 90 and 18 μ M, respectively [36]. International Journal of Recent Advances in Pharmaceutical Research October 2011; 4: 6-15 with IC50 value of 100 ng ml⁻¹ proves that the drug is used in the treatment of inflammatory conditions [37].

Antifungal: The 50% methylene chloride in hexane flash column fraction of the extract of the leaves of *Ficus racemosa* Linn. was found to have antifungal activity. The extract inhibited the growth of several plant pathogens (*Curvularia sp, Colletotrichum gloeosporioides, Alternaria sp, Corynespora cassiicola and Fusarium sp*). Psoralen was identified as the active compound and was shown to be biodegradable,

having the potential to be developed as a fungicide against pathogens causing diseases on crops of economic importance [38].

Anti-filarial: Alcoholic as well as aqueous extracts caused inhibition of spontaneous motility of whole worm and nerve muscle preparation of Setaria cervi characterized by increase in amplitude and tone of contractions. Both extracts caused death of microfilaria in vitro. LC50 and LC90 were 21 and 35 ng/ml respectively for alcoholic, which were 27 and 42 ng ml⁻¹ for aqueous extracts [39].

Larvicidal: The larvicidal activity of crude hexane, ethyl acetate, petroleum ether, acetone and methanol extracts of the leaf and bark were assayed for their toxicity against the early fourth-instar larvae of *Culex quinquefasciatus* (Diptera: Culicidae). The larval mortality was observed after 24-h exposure. All extracts showed moderate larvicidal effects; however, the highest larval mortality was found in acetone extract of bark. The bioassay-guided fractionation of acetone extract led to the separation and identification of a tetracyclic triterpenes derivative. Gluanol acetate was isolated and identified as new mosquito larvicidal compound. Gluanol acetate was quite potent against fourth-instar larvae of *Aedes aegypti* L. (LC (50) 14.55 and LC (90) 64.99 ppm), *Anopheles stephensi* Liston (LC (50) 28.50 and LC (90) 106.50 ppm) and *C. quinquefasciatus* Say (LC (50) 41.42 and LC (90) 192.77 ppm) [40].

Renal anticarcinogenic: *F. racemosa* extract (200 mg kg⁻¹ body weight and 400 mg kg⁻¹ body weight) resulted in a significant decrease in xanthine oxidase, lipid peroxidation, γ - glutamyl transpeptidase and hydrogen peroxide. There was significant recovery of renal glutathione content and antioxidant enzymes, decrease in the enhancement of renal ornithine decarboxylase activity, DNA synthesis, blood urea nitrogen and serum creatinine [41].

Wound healing: Ethanol extract of stem bark showed wound healing in excised and incised wound model in rats [42].

Anticholinesterase: The present study evaluated the anticholinesterase activity of cold and hot aqueous extracts of *Ficus racemosa* stem bark against rat brain acetyl cholinesterase *in vitro*. Both the cold aqueous extract (FRC) and the hot aqueous extract (FRH) exhibited a dose dependent inhibition of rat brain acetyl cholinesterase. FRH showed significantly higher ($P \le 0.001$) cholinesterase inhibitory activity compared to FRC; however, both the extracts did not show 50% inhibition of AChE at the doses tested (200-1000 µg ml⁻¹). The IC50 values of 1813 and 1331 µg ml⁻¹ were deduced for FRC and FRH, respectively (calculated by extrapolation using Boltzmann's dose response analysis). The percentage of anticholinesterase activity was calculated. Both the extracts (FRC and FRH) exhibited a dose dependent inhibition of rat brain acetyl cholinesterase. However, their inhibitory activities were significantly lower ($P \le 0.001$) than that of neostigmine bromide, a standard acetyl cholinesterase inhibitor. Among FRC and FRH, FRH showed significantly higher ($P \le 0.001$) cholinesterase inhibitory activity compared to FRC; however, both the extracts did not show 50% inhibitory activity compared to FRC; however, both the extracts did not show 50% inhibitory activity activity compared to rate brain acetyl cholinesterase. However, their inhibitory activities were significantly lower ($P \le 0.001$) than that of neostigmine bromide, a standard acetyl cholinesterase inhibitory activity compared to FRC; however, both the extracts did not show 50% inhibition of AChE at the doses tested (200-1000 µg ml⁻¹) and hence IC50 values were calculated by extrapolation using Boltzmann's dose response analysis [43].

Anti-pyretic: Methanol extract of stem bark was evaluated on normal body temperature and yeast-induced pyrexia in albino rats, at doses of 100, 200 and 300 mg kg⁻¹ b.w p.o. It showed significant dose dependent reduction in normal body temperature and yeast-provoked elevated temperature which extended up to 5 hr after drug administration. The anti-pyretic effect was comparable to that of paracetamol [42].

Memory Enhancing: Alzheimer's disease (AD) is a progressive neurodegenerative disorder resulting in dementia and enhancement of acetylcholine (Ach) levels in brain using acetyl cholinesterase inhibitors is one of the most important approaches for the treatment of AD. Aqueous extract of Ficus racemosa Linn (Moraceae) bark having antiinflammatory, antioxidant, and Anticholinesterase activity was evaluated for its ability to enhance Ach levels, and to ascertain its antidementia activity in rats. This work was carried

out under the assumption that the *F. racemosa* extract may show combination of actions which could be beneficial in the treatment of AD, such as neuro protection, attributed to antioxidant and antiinflammatory property and may elevate levels of Ach like Ficus hispida extract reported earlier. The plant extract selected for investigation elevated Ach levels and improved memory in rats. The collective pharmacological actions attributed by *F. racemosa* extract may serve as beneficial and supporting agent in the treatment of AD [44].

Analgesic: The ethanol extract of bark and leaves was evaluated for analgesic activity by analgesiometer at 100, 300 and 500 mg/kg and was found to possess dose dependent analgesic activity [45].

CONCLUSIONS

The genus *Ficus* constitutes an important group of trees with immense medicinal value. The medicinal plants are widely used by the traditional medical practitioners for curing various diseases in their day to day practice. In traditional system of medicine, different parts such as root, fruit, leaves, stem, seeds, latex and even whole plant of *Ficus racemosa* (Linn) have been recommended for the treatment of gastric ulcer, diarrhea, wound healing, diabetes, hyper tension etc. It is one of the popular plant in Indigenous system of medicine such as Ayurveda, Siddha, Unani and also homeopathy system of medicine. *Ficus Racemosa Linn*. showed a wide range of pharmacological actions like hypoglycemic, hypolipidemic, anticarcinogenic, anti-diuretic, hepatoprotective, anti-ulcer, anti-inflammatory, anti-fungal etc. Bio active constituents like β -sitosterol, glauanol acetate in *Ficus Racemosa L.*, has been found to be largely responsible for the therapeutic potentials as a boon for ailments of human kind. Hence, the present study shows the therapeutic potential, pharmacological and phytochemical properties of various bioactive compounds present in the *Ficus racemosa* (Linn). However, more Clinical and Pathological studies should be conducted to investigate the active potentials of bioactive compounds present in this plant which can help in proving it to be a promising source in neutraceutical as well as pharmaceutical industry.

ACKNOWLEDGEMENT

The authors would like to express their sincere gratitude to: Dr. V.J. Sirwaiya, Principal, Wilson College; Management, VIVA Trust, Virar (W) Maharashtra. K.P.N. Kutty, Co-Ordinator, VIVA Trust, Virar (W) Maharashtra; Dr. R.D. Bhagat, Principal, VIVA College, Virar (W) Maharashtra; Non-Teaching Staff, Dept. Biological sciences, VIVA College Virar (W) Maharashtra.

REFERENCES

- [1] P.P Joy, J. Thomas, S. Mathew, B.P Skaria, Medicinal Plants. Tropical Horticulture, Naya Prakash, Calcutta, **2001**, 2, 123-125.
- [2] P.M Paarakh, *Ficus racemosa* Linn.-An overview. *Nat Prod Radiance*. 2009, 8, 84-90.
- [3] C. C Berg, Classification and distribution of *Ficus, Experientia*, **1989**, 45, 605-611.
- [4] N.P Manandhar, Fodder trees. *The Rising Nepal*, **1972**, 7, 1-2.
- [5] Anonymous, The Wealth of India, Council of Scientific and Industrial Research, New Delhi, India, **1952**, 35-36.
- [6] P. K Warrier, Indian Medicinal Plants-A Compendium of 500 species, Orient Longman Ltd: Chennai, **1996** (Vol. III), 34-35.
- [7] R. N Chopra, I. C Chopra, K. I Handa, L. D Kapur, Indigenous Drugs of India, U.N. Dhur and Sons Pvt. Ltd, Calcutta, **1958**, 674-675.
- [8] N. Sirisha, M. Sreenivasulu, K. Sangeeta, C. M Chetty, Antioxidant Properties of Ficus Species-A review, *Int J Pharm Tech Res.*, **2010**, 3, 2174-2182.
- [9] C.P. Khare, Encyclopedia of Indian Medicinal Plants Springer publication, 2004, 216-217.

- [10] A. Husain, O. P Virmani, S. P Popli, L. N Misra, M. M Gupta, G. N Srivastava, Z Abraham & A. K Singh, Dictionary of Indian Medicinal Plants, CIMAP, Lucknow, India, **1992**, 546.
- [11] C. Suresh, L. Jawakhar and M. Sabir, Chemical examination of the fruits of *Ficus Glomerata*, J Indian Chem Soc, 1979, 56(12), 1269-1270.
- [12] K. Murti, U. Kumar, M. Panchal, M. Shah, Exploration of preliminary phytochemical studies of roots of *Ficus racemosa*, *Marmara Pharm J*, **2011**, 15, 80-83.
- [13] J. Bheemachari, K. Ashok, N. H Joshi, D. K Suresh, V. R. M Gupta, Antidiarrhoeal evaluation of *Ficus racemosa* Linn. latex, *Acta Pharmaceutica Sciencia*, **2007**,49,133-138.
- [14] B. Joseph, S. J Raj, Phytopharmacological and phytochemical properties of three *Ficus* species an overview, *Int J Pharma Bio Sci*, **2010**, 1, 246-253.
- [15] A. K Nadkarni, Indian Materia Medica, Popular Book Depot:Bombay; **1954**, (3rd edn), 2571-2575.
- [16] S. K Sharma, V. K Gupta, In vitro antioxidant studies of *Ficus racemosa* Linn. root. *Pharmacognosy Magazine*, **2008**, 4, 70-74.
- [17] Y. S Prabhakar, K. D Suresh, A survey of cardioactive drug formulations from Ayurveda II: porridges, oils, clarified butters, electuaries, pastes, ash preparations and calcined powders. *Fitoterapia*, **1990**, 61, 395-416.
- [18] S. Vedavathy, D. N Rao, Herbal folk medicine of Tirumala and Tirupati region of chittoor, District, Anthra Pradesh, *Fitoterapia*, **1995**, 66, 167-171.
- [19] C. H Chandrashekhar, K. P Latha, H. M Vagdevi, V. P Vaidya, Anthelmintic activity of the crude extracts of *Ficus racemosa*, *Int J Green Pharm*, **2008**, 2, 100-103.
- [20] V. V Patil, R. B Pimprikar, N. G Sutar, *et al*, Anti- Hyperglycemic activity of *Ficus racemosa* Linn leaves, *J Pharm Res*, **2009**, 2,54-57.
- [21] P. K Mukherjee, K. Saha, T. Murugesan, S. C Mandal, M. Pal, B. P Saha, Screening of antidiarrhoeal profile of some plant extracts of a specific region of West Bengal. India, J *Ethnopharmacol*, 1998, 60, 85-89.
- [22] J.A Parrotta, Healing Plants of Peninsular India, CABI publishing, USA, 2001, 557-558.
- [23] K. Naghma and S. Sarwat, Modulatory Effect of *Ficus racemosa*:Diminution of potassium Bromate-Induced Renal Oxidative Injury and Cell Proliferation Response, *Basic Clin Pharmacol Toxicol*, 2005, 97(5), 282 – 288.
- [24] V. Agarwal and B. M Chouhan, A study on composition and hypolipidemic effect of dietry fibre from some plant foods, *Plant Foods Hum Nutr*, **1988**, 38(2), 189-197.
- [25] S. M Patel, S. A Vasavada, Studies on *Ficus racemosa-* Part I: antiulcer activity, *Bull Medico Ethnobotany Res*, **1985**, 6, 17-27.
- [26] C. H. V Rao, A. R Verma, K. M Vijay, S. Rastogi, Gastric protective effect of standardized extract of *Ficus glomerata* fruit on experimental gastric ulcers in rats, *J Ethnopharmacol*, 2008, 115, 323-326.
- [27] K. P Channabasavaraj, S. Badami, S. Bhojraj, Hepatoprotective and antioxidant activity of methanol extract of *Ficus glomerata*, *J Nat Med*, **2008**,62,379-383.
- [28] V. P Veerapur, K. R Prabhakar, V. K Parihar, *et al, Ficus racemosa* stem bark extract: a potent antioxidant and a probable natural radioprotector, *Evid Based Complement Altern Med*, **2009**, 6, 317-324.
- [29] I. A Jahan, N. Nahar, M. Mosihuzzaman, *et al*, Hypoglycaemic and antioxidant activities of *Ficus racemosa* Linn. Fruits, *Nat Prod Res*, **2008**, 23,399-408.
- [30] W. D Ratnasooriya, J. R Jayakody, T. Nadarajah, Antidiuretic activity of aqueous bark extract of Sri Lankan *Ficus racemosa* in rats, *Acta Biol Hungary*, **2003**, 54, 357-363.
- [31] S. C Mandal, T. K Maity, J. Das, B. P Saba, M. Pal, Hepatoprotective activity of *Ficus racemosa* leaf extract on liver damage caused by carbon tetrachloride in rats, *Phytother Res*, **1999**, 13, 430-432.

- [32] R. R Bhaskara, T. Murugesan, M. Pal, B. P Saha, S. C Mandal, Antitussive potential of methanol extract of stem bark of *Ficus racemosa* Linn. *Phytother Res*, **2003**, 17, 1117-1118.
- [33] A. Kar, B. K Choudhary and Ng. Bandyopadhyay, comparative evaluation of hypoglycemic activity of some Indian medicinal plants in alloxan diabetic rats, *J Ethno pharmacol*, **2003**, 84 (1), 105-108.
- [34] L. E Swain, K. R Downum, Light-activated toxins of the Moraceae, *Biochem. Sys. Ecol*, 1990, 18, 153-156.
- [35] P. K Mukherjee, K. Saha, T. Murugesan, S. C Mandal, M. Pal and B. P Saha, Screening of antidiarrhoeal profile of some plant extracts of a specific region of west Bengal, India, *J.Ethno pharmacol*, **1998**, 60 (1), 85-89.
- [36] R. W Li, D. N Leach, S. P Myers, G. D Lin, G. J Leach, P. G Waterman, A new anti-inflammatory glucoside from *Ficus racemosa L*, *Planta Med*, **2004**, 70, 421-426.
- [37] R. W Li, S. P Myers, D. N Leach, G. D Lin, G. Leach, A cross-cultural study: anti- inflammatory activity of Australian and Chinese plants, *J Ethnopharmacol*, 2003, 85, 25-32.
- [38] S. A Deraniyagala, R. L. C Wijesundera, O. Weerasena, Journal of the National Science Council of Sri Lanka, *J. Natl. Sci. Counc, Sri Lanka*, **1998**, 26 (1), 19-26.
- [39] V. Mishra, N. U Khan, K. C Singhal, Potential antifilarial activity of fruit extracts of *Ficus racemosa* Linn. against Setaria cervi in vitro, *Indian J. Exp. Biol*, **2005**, 43, 346.
- [40] N. Khan, S. Sultana, Chemomodulatory effect of *Ficus racemosa* extract against chemically induced renal carcinogenesis and oxidative damage response in Wistar rats, *Life Sci*, **2005**, 77, 1194-1210.
- [41] N. Khan, S. Sultana, Modulatory effect of *Ficus racemosa*: diminution of potassium bromateinduced renal oxidative injury and cell proliferation response, *Basic Clin Pharmacol Toxicol*, **2005**, 97, 282-288.
- [42] R. B Rao, K. Anupama, K. R Swaroop, T. Murugesan, M. Pal, S. C Mandal, Evaluation of antipyretic potential of *Ficus racemosa* bark, *Phytomedicine*, **2002**, 9, 731-733.
- [43] A. P Breen, J. A Murphy, Reactions of oxyradicals with DNA, *Free Rad Biol Med*, **1995**, 18, 1033–1077.
- [44] P. K Warrier, Indian medicinal plants-A compendium of 500 species, Orient Longman Ltd.Chennai, **1996**, Vol. III, 38-39.
- [45] P. Malairajan, G. K Geetha, S. Narasimhan, K. V Jessi, Analgesic activity of some Indian Medicinal Plants. *J Ethnopharmacol*, **2006**, 106, 425-428.