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PHARMACOLOGICAL ACTIVITIES OF SELECTED MEDICINAL PLANTS - A REVIEW

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ABSTRACT

The focuses on the medicinal plants were increased all over the world due to its immense potential in controlling several diseases from ancient civilization. The present review documented on the Morphological distribution, Phytochemicals, Traditional medicine and Pharmacological activities in selected medicinal plants. Medicinal plants under the review are *Plumbago zeylanica*, *Curculigo orchioides*, *Tinospora cordifolia*, *Semecarpus anacardium* and *Asparagus racemosus* which contribute in several poly herbal formulations. The pharmacological effects may be mainly due the presence of its principle active components and several secondary metabolites.

Keywords: Medicinal plants, antioxidants, Phytochemistry, Secondary metabolite.

INTRODUCTION

Medicinal plants are from the dawn of civilization to combat diseases and have been considered as a valuable and cheap source of unique phytoconstituents for the human society. They are used extensively in the development of drugs against various diseases [1]. In the ethno medical advance, credence is given to written or oral information on medicinal use of the plant and based on the information for the plants collected and evaluated for the drug discovery process [2]. A retrospective analysis of the NCI program described that the proportion of active medicine was substantially based on taxonomy [3]. In the last few decades, the field of herbal medicine is getting popularized in both developed and developing countries. This is because the herbal medicines are cheap, natural origin with higher safety margins and lesser or no side effects [4].

Selected plants under the review

Plumbago zeylanica

Taxonomical description

Kingdom: Plantae
Family : Plumbaganaceae
Order: Plumbaginales
Genus: *Plumbago*
Species: *zeylanica*

Morphological description

The plant is an evergreen small, perennial shrub which grows upto the height of 3-4 feet. The leaves are

simple, alternate, oblong, spirally arranged, hairy margin, pointed to the tip. The flowers are white in color, 10-25 cm long [5]. The fruits are like a small cocklebur with glue on the soft spines. The root is stout, cylindrical, blackish red in color with a pungent odor [6].

Traditional medicine

Plumbago zeylanica has been reported to be used in variety of folk medicine in Africa and Asia. Traditionally, *P. zeylanica* is used to kill intestinal parasites, anemia due to stagnant blood, external as well as internal trauma and malignant furunculosis scabies [7]. In India it is usually used to treat fever or malaria. It is used as an irritant of the skin in the treatment of leprosy, dyspepsia, piles, diarrhea and skin diseases [8]. It has been specially recommended in the treatment of rheumatism [9].

Phytochemistry

Based on the literature survey several phytochemicals isolated are Naphthoquinones, Binaphthoquinones [10], Coumarins, Steroid glycosides [11], Di-phenyl sulfone [12], Carboxylic acid and Esters [13], Triterpenoids [14], Meroterpenes [15], Sugars [16] etc.,

The root is the chief medicinal property of the plant that contain several alkaloids such as plumbagin, a natural naphthaquinone (5-hydroxy-2-methyl-1,4-aphthoquinone), chitranone, zeylanone, plumbazeylanone, 2- methyl naphthaquin, dihydrosterone, lupeol and teraxesterol.

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Pharmacological activity

The root of this plant and its constituents are accredited with several therapeutic properties including anti atherogenic, cardiogenic, hepatoprotective, neuro protective and central nervous system stimulating properties [17].

Antidiabetic activity

The ethanolic extract of root of *P. zeylanica* offers a strong hypoglycemic effect and antioxidant protection in streptozotocin induced diabetic rats [18].

Antiallergic properties

The antiallergic properties of the 70% ethanol extract from *Plumbago zeylanica* stem (EPZ) were investigated. EPZ inhibits mast cell-dependent immediate allergic reactions, which is probably mediated by histamine from mast cells via elevating intracellular cAMP level and thus weaken the inflammatory action of mediators [19].

Antihyperlipidaemic activity

In hyperlipidaemic rabbits plumbagin, isolated from roots of *P. zeylanica* reduced the serum cholesterol and Low Density Lipoproteins. Plumbagin treated hyperlipidaemic subjects prevented the accumulation of triglycerides and cholesterol in liver and aorta and lowered atheromatous plaques of thoracic and abdominal aorta [20].

CNS activity

In rats the effects of 50% ethanol extract from *P. zeylanica* roots were tested on locomotor behavior and central dopaminergic activity. The extracts significantly raised the spontaneous motility of the animals. The root extracts enhanced the spontaneous ambulatory activity without inducing stereotypic behavior [17].

Antileishmanial activity

The study by Mishra discloses strong *in vitro* antileishmanial activity of 2-methyl-5 (3'-methyl-but-2'-enyloxy)-[1, 4] naphthoquinone isolated from *Plumbago zeylanica* [21].

Anticancer activity

The ethanolic extract of *Plumbago zeylanica* Linn possess significant anticancer activity and also reduces the level of lipid peroxidation. This is mainly due to higher content of terpenoids and flavonoids in Ehrlich ascites carcinoma in animal model.

The anticancer effect of plumbagin was investigated *in vivo* using NB4 tumor xenograft in NOD/SCID mice. The results indicate that plumbagin has potential as a novel therapeutic agent for myeloid leukemia [22]. Plumbagin exerted anticancer activity by the induction of apoptosis on non-small cell lung carcinoma cells by modulating the pro-survival and pro-apoptotic signaling pathway [23].

Semecarpus anacardium

Taxonomical description

Kingdom: Plantae

Family: Anacardiaceae

Order: Sapindales

Genus: *Semecarpus*

Species: *anacardium*

Morphological Description:

It is a moderate-sized deciduous tree grows upto 3500 ft height [24]. Leaves are simple, alternate, oblong, long, rounded at the apex coriaceous glabrous. The flowers are greenish white, in panicles. The nut is about 2.5 cm long, ovoid and smooth lustrous black. The fruit ripens appears ovoid or oblong drupe shining black. The bark is grey in color and exudes an irritant secretion on incising [25].

Phytochemistry

The most significant components of *S. anacardium* Linn. are bharbilanols, phenolic compounds [26], biflavonoids [27], Glycosides and sterols [28]. Other components isolated are Anacardoside [29], Semecarpetin [30], Semecarpuflavone [31], Anacarduflavone [32] etc.,

Traditional medicine

In traditional medicine, *Semecarpus anacardium* is used to treat ailments including skin infections and non-infectious conditions of the skin. It also been used to treat ringworms, psoriasis, eczema and corns.

Seeds are generally boiled in milk and the milk is consumed to treat several ailments. Externally, the oil is applied on wounds to prevent pus formation, better healing of wounds in glandular swellings and filariasis [33]. Traditional ayurvedic practitioners used this herb to cure reproductive complaints (low sperm count and painful menstruation). It is also effective for nervous complaints (forgetfulness, dementia and psychological trauma). The herb has been used to support digestion and improve the appetite [34].

Pharmacological activity

Antihyperlipidaemic activity

The cardiac activity of *Semecarpus anacardium* (SA) generally reduces the tissue and serum hyperlipidemia by the inhibition of intestinal cholesterol absorption coupled with peripheral disposal thus possessing anti-atherosclerotic activity [35].

CNS activity

Farooq *et al.* evaluated the therapeutic effect of SA nuts, extracted with milk, mainly for its locomotor and nootropic activities in experimental animal models. The extract showed a slight CNS depressant effect on 150 mg/kg [36].

Antiatherogenic effect

Atherosclerosis is the developmental imbalance between the pro-oxidants and antioxidants. To prevent such condition, antioxidant therapy is beneficial. Sharma *et al.* demonstrated the cardiac activity of SA, as it generally reduces the serum and tissue hyperlipidemia level by the

inhibition of intestinal cholesterol absorption. This coupled with peripheral disposal thus possessing anti-atherosclerotic activity. It is possible that the beneficial antiatherogenic effect may be related to its antioxidant, anticoagulant, hypolipidemic and platelet anti-aggregation properties [35].

Anti-inflammatory activity

The different solvent extracts of fruits of SA were tested to study the anti-inflammatory activity by carrageenan-induced paw edema in albino rats. The results showed significant anti-inflammatory activity comparable to aspirin (reference standard) [37].

Immunomodulatory activity

The immunomodulatory potency, antioxidative, glucose level restoring and mineral regulation properties of SA nut extract in hepatocellular carcinoma induced by aflatoxin B₁ in animals [38].

Antiarthritic activity

The antiarthritic activity was proved in the chloroform extract of *Semecarpus anacardium* nuts significantly reduced acute carrageenan-induced paw edema in rats and also active against the secondary lesions produced by adjuvant-induced arthritis [39].

Anticancer activity

The biochemical basis of anticarcinogenesis potency of *Semecarpus anacardium* was studied using Aflatoxin-B₁ induced hepatic carcinoma in rats [40]

Nephrotoxicity

Prabhu *et al.* studied the antimutagenic effect of SA under *in vivo* condition. The action of SA oil extract has definite beneficial role against mitomycin-C induced mutagenic role [41]. In other study, Krishnarajua *et al.* found that aqueous extracts of SA was found to be effective when screened for the cytotoxicity activity using brine shrimp lethality test [42].

Curculigo orchioides

Taxonomical description

Kingdom: Plantae
Family: Hypoxidaceae
Order: Asparagales
Genus: *Curculigo*
Species: *orchioides*

Morphological description

It is a perennial shrub having short or elongated fleshy roots. Leaves are 6 to 18 inch in length and half to one and half inch broad. It is speared shaped and bears stripes on it. The apex of the leaves is rooting. Flowers are 1 inch in length, half to two-third inch in diameter, shiny and are just above the ground. Fruit is half inch in length, capsulated, ovate in shape contains 1 to 4 seeds in it. Seeds are shiny, oblong in shape, one-eighth inch long, striped and sharp at the apex and base. Rhizome is 1 feet in length and is pulpy [43].

Phytochemistry

The rhizome has been reported to contain different types of compounds such as Curculigoside A_C [44], Curculigosaponins A_M [45], Orcinol glycoside [46], 2, 6-dimethoxy benzoic acid [47] and Orchiosides A and B [48], Tannin, Resin and alkaloid [49]. The secondary metabolites from the rootstock has been identified as 5, 7-dimethoxy glucopyranoside (flavones glycoside) and also numerous fatty acids from *C. orchioides* root oil [50]. They are palmitic, oleic, linolenic, linoleic, arachidic and benenic acid.

Curculigoside (phenolic glycoside) from the rhizomes and its structure has been elucidated as 5-hydroxy- 2-O-b-d-glucopyranosyl benzl 1, 2, 6-dimethoxy benzoate [51] (Misra *et al.*, 1984). The dried rhizomes of *Curculigo orchioides* yielded phenolic glycosides, curculigoside, orcinol-beta-D-glucoside, cycloartane saponins, curculigosaponin G and curculigosaponin I.

Traditional medicine

The plant has been named as safed musli in ayurvedic formulations. The juice of the plant tuber mixed with the juice of *Allium sativum* used to cure blindness and white spot on the eye ball. Rhizomes have been reported to be useful in asthma [52]. The plant also holds the reputation of being a demulcent, diuretic, tonic and aphrodisiac.

Rhizomes are prescribed in the treatment of piles, jaundice, asthma, diarrhea and gonorrhoea [43]. Rhizomes have been claimed for the anti diabetic properties in various studies [53]. It is claimed to be a medical cure for piles, asthma, jaundice, diarrhoea, colic, gonorrhoea and to be a aphrodisiac [54].

Pharmacological activity

Anticonvulsant and antioxidant

Ethanol extract of *C. orchioides* has been reported to have anticonvulsant, sedative and androgen-like effect and also adaptive effects (hypoxia and hyperthermia) [55]. Methanolic extract of *Curculigo orchioides* (MEC) was investigated using carbon tetrachloride (CCl₄)-intoxicated rat liver as the experimental model reported the antioxidant property [56].

Anti-inflammatory and Antihepatotoxic activity

The efficacy of methanolic extract of *C. orchioides* in combating oxidative stress by hepatic damage [57]. The study was performed by Sae-kang ku and his colleagues to examine the effects of *Curculigo orchioides* (curculiginis rhizome) on acute reflux esophagitis (RE) in rats that are induced by pylorus and forestomach ligation operation. Anti-inflammatory and protective effects of the extract of *C. orchioides* could decrease the severity of reflux esophagitis and control esophageal mucosal damage [58].

Estrogenic activity

Ethanol extract of rhizome possesses estrogenic activity as it showed a significant increases the vaginal cornification, uterine wet weight (p<0.01), uterine glycogen content (p<0.01) and a proliferation changes [59].

Anti osteoporotic activity

The ethanolic extract of the plant showed potential antiosteoporosis activity as it prevented bone loss in the trabecular bone of the tibia in ovariectomized rats [60].

Antiasthmatic activity

Ethanol extract of *Curculigo orchioides* showed effectiveness against histamine-induced contraction. *Curculigo orchioides* showed significant protection at lower doses while further increase in the dose level showed reduced activity [61].

Anti cancer activity

The anticancer activity of *C. orchioides* in indigenous system of medicine as a tonic and also reported as an ayurvedic food supplement against DMBA (dimethyl benz anthracene) induced mammary tumours [62].

Immunostimulant activity

The methanolic extract of the rhizome has been shown to enhance the phagocytic activity of macrophages. The pure glycoside fraction of the rhizome has been found to stimulate immune response by acting both on macrophages and lymphocytes [63]. Immunostimulant activity of *C. orchioides* has also demonstrated [64].

Asparagus racemosus

Taxonomical description

Kingdom: Plantae
Family: Asparagaceae
Order: Asparagales
Genus: *Asparagus L.*
Species: *racemosus*

Morphological description:

A. racemosus is plant with a woody stem has needle like leaves with small white flowers [65], Scandant, much branched spinous under shrub with tuberous, short root, stock bearing numerous fusiform tuberous roots, thick leaves reduced to minute chaffy scales & spines. Cladodes acicular 2-6 hate, falcate finely acuminate flower white, berries 7mm in diameter, globose, 1-seeded, red [66].

The air dried roots are tuberous, elongated and tapering at both the ends, up to 30-100 cm long. The fresh roots are fleshy and white in color; while on drying it become shrinked, longitudinal ridges appeared and the color turned to light brown. Outer surface of the fresh root is soft and contains epidermal hairs. Taste is mucilaginous, fracture brittle. Roots are cylindrical, fleshy tuberous straight or slightly curved, tapering towards the base and swollen in the middle, white buff colour, 5- 15cm in length 1-2 cm diameter [67].

Phytochemistry

Steroidal saponins, known as shatvarins. Shatvarin I-VI compounds are present. Shatvarin I is the chief glycoside with the combination of 3-glucose and rhamnose moieties attached to sarsapogenin [68]. Shatvarin V, Asparinins, Asparosides, Curillins,

Curillosides have also been reported [69]. Furan compound- Racemofuran [70]. Flavonoids- Glycosides, hyperoside, rutin and quercetin are present in flower and fruits [71]. Polycyclic alkaloid- Asparagine A, a cage type pyrrolizidine alkaloid [72].

The isolation and characterization of asparagine-polycyclic alkaloid [73], a new 9, 10-dihydrophenanthrene derivative named racemosol and kaempferol were also isolated from the ethanolic root extract of *A. racemosus* [74].

Pharmacological activity

The plant has several pharmacological properties like antioxidant, immunostimulant, antidyspepsia and antitussive effects [75].

Antiabortifacient activity

The Antiabortifacient activity of *A. racemosus* roots are recommended in cases of threatened abortions. This activity is due to Shatavarin-I [74]. The polycyclic alkaloid asparagine A is also reported to have an antioxytotic action, showing an antiabortifacient affect [76].

Immunomodulating property

Immunomodulating property of *A. racemosus* has been shown to protect the rat and mice against experimental induced abdominal sepsis [77].

Anticancer activity

The anticancer activity of shatavarins (containing shatavarin IV) isolated from the roots of *A. racemosus* (Wild) exhibited significant effect *in vitro* and *in vivo* in experimental models [78].

Antiulcer activity

A. racemosus was more effective in reducing gastric ulcer in indomethacin-treated gastric ulcerative rats [79].

Analgesic and antidiarrhoeal properties

The evaluation of analgesic and antidiarrhoeal properties of the ethanol extract of whole plant of *A. racemosus* carried out by Karmakar *et al.* The study showed that analgesic activity of the plant increased the mean latent period and decreased the frequency of defecation [80].

Notropic effect

A. racemosus root extract has been reported to generate positive ionotropic and chronotropic effect on frogs heart with lower doses and cardiac arrest with higher doses [81]. Alcoholic extract of root of *A. racemosus* has been shown to significantly reduce the enhanced levels of transaminase and alkaline phosphatase in CCl₄-induced hepatic damage in rats [82].

Tinospora cordifolia

Taxonomical description

Kingdom: Plantae

Family: Menispermaceae
Order: Ranunculales
Genus: *Tinospora Miers*
Species: *cordifolia*

Morphological description

It is an under-shrub, climbs up to 1-3 m high, with stout and creeping root stock. The root occurs in fascicle or clusters at the base of the stem with succulent and tuberous rootlets. The stem is scandent, woody, striate and climbing. The young stem is delicate, brittle and smooth. The spines are long, sub-recurved or straight [83, 84]. The flowers are simple or branched racemes of 3 cm long. The pedicel is slender and jointed in the middle. The anthers minute and purple. Perianth lobes white, fragrant and 3 mm in length. The berry is globular or obscurely, three lobbed, purple-reddish [85].

Phytochemistry

Three major groups of compounds include protoberberine alkaloids, terpenoids and polysaccharides [86]. Tinosporoside, cardioside, berberine and cordiofolioside were isolated from the plant leaves [87], 3(a, 4-dihydroxy-3-methoxybenzyl)-4-(4-hydroxy-3-methoxybenzyl) a lignin isolated [88].

Traditional medicine

Powder of *Terminalia chebula*, *Tinospora cordifolia* and *Trachyspermum ammi* in equal quantity is administered orally, once daily with salt for the treatment of cough. Decoction of these drugs is also to be taken for the treatment of cough by the people of Haryana. Paste of *T. cordifolia* and *Piper nigrum* is administered orally once daily for leucorrhoea. The inhabitants of Uttar Pradesh take the juice of stem orally with honey for the treatment of asthma [89]. The tribals of Bombay uses *T. cordifolia* as drug in the treatment of fever, jaundice, chronic diarrhoea and dysentery [90].

Pharmacological activity

Anticonvulsant activity

The anticonvulsant activity of Petroleum ether and ethanol extracts of *Tinospora cordifolia* (TC) was carried out using supramaximal electric shock method in rats. The Ethanolic extract of *Tinospora cordifolia* possesses significant anticonvulsant property [91].

Hypoglycaemic activity

The mechanism of anti-diabetic activity of TC may be through some peripheral machinery such as increasing the glycogen storage in the liver or decreasing the glucose release from the liver [92].

Anti-inflammatory activity

The decoction of *T. cordifolia* showed anti-inflammatory activity on carrageenin-induced hind paw

REFERENCES

1. Sarasa D, Sridhar S, Prabakaran E. Effect of an antidiabetic extract of *Trigonella foenum-graecum* on normal and alloxan induced diabetic mice. *Int J Pharm Pharm Sci*, 4(1), 2012, 63-65.

oedema in rats [93]. The effect of extract of stem of *T. cordifolia* was studied on the contractile response due to various agonists (such as histamine, 5-HT, bradykinin, prostaglandin E₁ and F_{2a}, cholinomimetics and KCl) on smooth muscles [94].

Antiulceration- property

The root of *T. cordifolia* was observed to induce a marked protective action against restrain stress induced ulcerization when compared to diazepam [95].

Anti-psychotropic activity

An herbal psychotropic preparation BR-16A containing *T. cordifolia* was investigated in short term memory paradigms in mice. The results suggest for possibly nootropic action of BR-16A involving cholinergic and GABAergic modulation [96].

Immunobiological activities

T. cordifolia has been studied for their effect on intra-abdominal sepsis to elucidate host defense mechanism to counter infective stress. The results indicate that *T. cordifolia* has immunomodulating properties [97].

Liver disorders

The drug was also studied against the hepatic damage induced by carbon tetra chloride-standard hepatotoxin. It was very effective in preventing fibrous alteration and inducing regeneration by paranchymal tissue [98].

Anti-HIV effects

TC has been shown to demonstrate a decrease in the persistent resistance of HIV virus thus improving the therapeutic effect. An anti-HIV effect of TC was revealed by reduction in eosinophil count, Hb count, stimulation of macrophages and B lymphocytes and polymorphonuclear leucocytes thus, revealing its promising role of application in management of the disease [99].

CONCLUSION

Herbal medicines are popularized as a solution for maintaining health and for the treatment for several diseases rather than allopathic medicine. The plants under the present study reviewed the phytochemicals responsible for several disease and these plants plays the crucial role in the preparation of several polyherbal formulations in siddha, Ayurveda and Unani medicines. The confidence in herbal medicines is backed by their long term usage and less toxicity but its toxicity plays a beneficial role in several diseases including cancer, arthritis etc., Furthermore, the combinations of these plants in controlling major diseases is to be carried out for the non-toxic herbal formulations.

2. Valeriote F, Moore RE, Patterson GML, Paul VJ, Scheuer PJ, Corbett T. Discovery of natural products from microalgae and marine organisms. In: Valeriote FA, Corbett TH, Baker LH, editors. *Anticancer Drug Discovery and Development: Natural Products and New Molecular Models*. Kluwer Academic Publishers; Boston/Dordrecht/London, 1994, 1–25.
3. Cordell GA, Beecher CWW, Pezzuto JM. Can ethnopharmacology contribute to the development anticancer drugs? *J. Ethnopharmacol*, 32, 1991, 117-133.
4. Ayyanara M, Ignacimuthu S. Ethnobotanical survey of medicinal plants commonly used by Kani tribals in Tirunelveli hills of Western Ghats in India. *J Ethnopharmacol*, 134, 2011,851-64
5. Geltz NR and Russell SD. Two dimension electrophoretic studies of the proteins and polypeptides in mature pollen grains and male germ unit of *Plumbago zeylanica*. *Plant Physiol*, 88, 1988, 764-9.
6. Schlauer J. New data relating to the evolution and phylogeny of some carnivores plant families. *Carnivores plants newsletter*, 26,1997,34-8
7. Jiangsu. New Medical College. *Zhongyao Dictionary (Encyclopedia of Chinese Materia Medica)*. Scientific & Technological Press, Shanghai, 1997, 711–712.
8. Modi J. *Textbook of Medicinal Jurisprudence and Toxicology*. Pripati Pvt Ltd, Bombay, India, 191, 595-596.
9. Anonymous. *The wealth of India: A dictionary of Indian Raw materials and Industrial products*, CSIR, New Delhi, 1989, 163-164.
10. Gunaherath GM, Gunatilaka AA. Studies on medicinal and related plants of srilanka,part 18,structure of a new naphthoquinone from *Plumbago zeylanica*. *J chem soc perkin trans*, 1, 1998, 407-10.
11. Lin LC, Yang LL, Chou CJ. Cytotoxic naphthoquinone and plumbaic acid glucosides from *Plumbago zeylanica*. *Phytochem*, 62, 2003, 619-22.
12. Amatya S, Ghimire U, Tuladhar SM. Isolation of aromatic sulphone from *Plumbago zeylanica*. *Pak J Sci Ind Res*, 50, 2007, 184-5.
13. Rahman MS, Anwar MN. Fungitoxic and cytotoxic activity of a novel compound 1,2 benzenedicarboxylic acid, diisooctyl ester of *Plumbago zeylanica* Linn. *Asian J Microbiol Biotechnol Environ Sci*, 8, 2006, 461-464.
14. Dinba B, Saha S. Chemical constituents of *Plumbago zeylanica* aerial parts and *Thevetia neriifolia* roots. *J Indian Chem Soc*, 67, 1990, 88-9.
15. Lin LC and Chou CJ. Meroterpenes and C-glucosylflavonoids from the aerial parts of *Plumbago zeylanica*. *Chinese Pharm J*, 2003, 5577-81.
16. Chowdhury AK, Chakder SK, Khan AK. Isolation and characterization of chemical constituents of *Plumbago zeylanica* root. *J Bangladesh Acad Sci*, 5, 1981, 71-4.
17. Bopaiah CP, Pradhan N. Central nervous system stimulatory action from the root extract of *Plumbago zeylanica* in rats. *Phytother Res*, 15, 2001, 153-6.
18. Kumar G. Sharmila Banu G, Maheswaran R, Rema S, Rajasekara Pandian M, Murugesan AG. On blood glucose and plasma antioxidant status in STZ Diabetic rats. *J nat rem*, 7(1), 2007, 66–71.
19. Dai Y, Hou LF, Chan YP, Cheng L, But PP. Inhibition of immediate allergic reactions by ethanol extract from *Plumbago zeylanica* stems. *Biol Pharm Bull*, 27(3), 2004, 429-32.
20. Sharma I, Gusain D, Dixit VP. Hypolipidaemic and antiatherosclerotic effects of plumbagin in rats. *Indian J Physiol Pharmacol*, 35(1), 1991, 10-14.
21. Mishra BB, Gour JK, Kishore N, Singh RK, Tripathi V, Tiwari VK, An antileishmanial prenyloxy - naphthoquinone from roots of *Plumbago zeylanica*. *Nat Prod Res*, 27(4-5), 2013, 480-5.
22. Xu KH, Lu DP. Plumbagin induces ROS-mediated apoptosis in human promyelocytic leukemia cells in vivo. *Leuk Res*, 34(5), 2013, 658-65.
23. Gomathinayagam R, Sowmyalakshmi S, Mardhatillah F, Kumar R, Akbarsha MA, Damodaran C. Anticancer mechanism of plumbagin, a natural compound, on non-small cell lung cancer cells. *Anticancer Res*, 28(2A), 2008, 785-92.
24. Kirtikar KR, Basu BD. *Indian medicinal plants Vol. 3*. Dehradun, India: International Booksellers and Publishers, 1975, 667.
25. Bhitre MJ, Patil S, Kataria M, Anwikar S, Kadri H. Anti-inflammatory activity of the of fruits *Semecarpus anacardium* Linn. *Asian J Chem*, 20, 2008, 2047–50.
26. Mathur HN, Agarwal JS. Phenolic modified resin of oil varnishes. *J Sci Indian Res*, 12, 1953, 411.
27. Ishatulla K, Ansari WH, Rahman W, Okigawa M, Kawanon N. Bioflavanoids from *Semecarpus anacardium* linn. *Indian J Chem*, 15, 1977, 617–22.
28. Govindachary TR, Joshi BS, Kamal VM. Phenolic constituents of *Semecarpus anacardium*. *Indian J Chem*, 9, 1971, 1044.
29. Majumdar SH, Kulkarni SB, Chakraborty GS. Medicinal potentials of *Semecarpus anacardium* nut: A review. *J Herb Med Toxicol*, 2, 2008, 9–13.
30. Murthy SS. Semicarpetin A biflavonoid from *Semecarpus anacardium*. *Phytochem*, 27, 1988, 3020–2.
31. Murthy S. A bioflavonoid from *Semecarpus anacardium*. *Phytochem*, 22, 1983, 1518-1520.
32. Murthy SS. New biflavonoid from *Semecarpus anacardium* linn. *Clin Acta Turcica*, 20, 1992, 30.
33. Paras Jain, Sharma HP. A Potential Ethnomedicinal Plant: *Semecarpus anacardium* Linn. A review. *Int. J. Res. Pharm. Chem*, 3(3), 2013, 564-572.

34. Kumar G, Sharmila Banu G, Maheswaran R, Rema S, Rajasekara Pandian M, Murugesan AG. On blood glucose and plasma antioxidant status in STZ Diabetic rats. *J. Nat. Remedies*, 7(1), 2007, 66 – 71.
35. Sharma A, Mathur R, Dixit VP. Hypocholesterolemic activity of nut shell extract of *Semecarpus anacardium* (Bhilawa) in cholesterol fed rabbits. *Indian J Exp Biol*, 33, 1995, 444–8.
36. Farooq SM, Alla TR, Rao NV, Prasad K, Shalam K, Satyanarayana S. A study on CNS effect of nut milk extract of *Semecarpus anacardium*. *Pharmacologyonline*, 1, 2007, 49–63.
37. Bhitre MJ, Patil S, Kataria M, Anwikar S, Kadri H. Antiinflammatory activity of the fruits of *Semecarpus anacardium* Linn. *Asian J Chem*, 20, 2008, 2047–50.
38. Premalatha B, Sachdanandam P. Potency of *Semecarpus anacardium* Linn. Nut milk extract against aflatoxin B induced hepatocarcinogenesis: Reflection on microsomal biotransformation. *Pharmacol Res*, 42, 2000, 161–6.
39. Saraf MN, Ghooi RB and Patwardhan BK. Studies on the mechanism of action of *Semecarpus anacardium* in rheumatoid arthritis. *J ethnopharmacol*, 25(2), 1989, 159.
40. Premalatha B, Sachdanandam P. Effect of *Semecarpus anacardium* nut extract against aflatoxin B -induced 1 hepatocellular carcinoma. *Fitoterapia*, 70, 1999, 484-492.
41. Prabhu D, Rajwani LS, Desai PV. The antimutagenic effect of *Semecarpus anacardium* under *in vivo* condition. *Asian J Chem*, 12, 2005, 13–6.
42. Krishnarajua AV, Rao TV, Sundararajua D, Vanisreeb M. Assessment of bioactivity of Indian medicinal plants using brine shrimp (*Artemia salina*) lethality assay. *Int J Appl Sci Eng*, 3, 2005, 125–34.
43. Chopra RN, Nayarand SL, Chopra IC. Glossary of Indian medicinal plants. Publication and information directorate, CSIR, New Delhi, 1956, 84.
44. Fu DX, Lei GQ, Chen XW, Chen JK, Zhou TS. Curculigoside C, A new phenolic glucoside rhizomes of *Curculigo orchioides*. *Acta Bot. Sin*, 46, 2004, 621–624.
45. Xu JR, Xu RS. Cycloartane-type saponin and their glycosides from *Curculigo orchioides*. *Phytochem*, 31, 1992, 2455–2459.
46. Wu Q, Fu DX, Hou AJ, Lei GQ, Liu ZJ, Chen JK, Zhou TS. Antioxidative phenols and phenolic glycosides from *Curculigo orchioides*. *Chem Pharm Bull*, 53, 2005, 1065–1067.
47. Chen CX, Ni W and Mei WL. The glycosides from *Curculigo orchioides*. *Acta Bot. Yunnan*, 21, 1999, 521–524.
48. Gupta M, Achari B and Pal BC. Glucosides from *Curculigo orchioides*. *Phytochem*, 66, 2005, 659–663.
49. Anon. The Ayurvedic Pharmacopoeia of India, 1st Ed., Vol. 4. Government of India, New Delhi, 2004, 122.
50. Mehta BK, Bokadia MM, Mehta SC. Study of root oil compound fatty acids of *Curculigo orchioides* roots. *Indian Drugs*, 18(3), 1980, 109-110.
51. Misra TN, Singh RS, Tripathi DM. Aliphatic compounds from *Curculigo orchioides* rhizomes. *Phytochem*, 23 (10), 1984(b), 2369-2371.
52. Atal CK, Kapoor BM. Cultivation and utilization of medicinal plants. Jammu-Tawi, India: Regional Research Laboratory, Council of Scientific and Industrial Research publication, 1977, 451.
53. Parrotta JA. Healing plants of peninsular India, New Delhi, Kalyani publishers, 1977, 306.
54. Kirtikar KR, Basu BD. Indian medicinal plants. Vol II. Leader Press. Allahabad, India, 1935, 2469.
55. Chen QS, Chen WR, Yang SY. Pharmacologic study of *Curculigo orchioides* Gaertn. *Zhongguo Zhong Yao Za Zh*, 14(10), 1989, 618-620.
56. Onkar P, Bangar J, Karodi R. Evaluation of Antioxidant activity of traditional formulation Giloy satva and hydroalcoholic extract of the *Curculigo orchioides* gaertn. *J Appl Pharm Sci*, 2(6), 2012, 209-213.
57. Venukumar MR, Latha MS. Antioxidant activity of *Curculigo orchioides* in carbon tetrachloride induced Hepatopathy in rats. *Indian J Clin Biochem*, 17 (2), 2002, 80-87.
58. Sae-Kang Ku, Jae-Soo Kim, Young-Bae Seo, *et al*. Effect of *Curculigo orchioides* on Reflux Esophagitis by Suppressing Proinflammatory Cytokines. *Am J Chin Med*, 40(6), 2012, 1241–1255
59. Vijayanarayanaa K, Rodrigues RS, Chandrasekhar KS, Subrahmanyam EV. Evaluation of estrogenic activity of alcoholic extract of rhizomes of *Curculigo orchioides*. *J Ethnopharmacol*, 114(2), 2007, 241-245.
60. Cao DP, Zheng YN, Qin LP, Han T, Zhang H, Rahman K, Zhang QY. *Curculigo orchioides*, a traditional Chinese medicinal plant, prevents bone loss in ovariectomized rats. *Maturitas*, 59(4), 2008, 373-80.
61. Pandit P, Singh A, Bafna AR, Kadam PV, Patil MJ. Evaluation of Antiasthmatic activity of *Curculigo orchioides* Gaertn. Rhizomes. *Indian J Pharm Sci*, 70(4), 2008, 440-4.
62. Tandon M, Shukla YN. Phytoconstitutes of *Asparagus adscendens*, *Chlorophytum arundinaceum* and *Curculigo orchioides*: A Review. *Curr Res Med Arom Plants*, 17, 1995, 42-50.
63. Lakshmi V, Pandey K, Puri A, Saxena RP, Saxena KC. Immunostimulant principles from *Curculigo orchioides*. *J Ethnopharmacol*, 89(2-3), 2003, 181-4.
64. Samanta SK. Modulation of male infertility by ayurvedic drugs. International Seminar on Traditional Medicine, Calcutta, 7 - 9, November, 1992, 127.
65. Aviva Romm. Botanical Medicine for woman's Health. Churchill Livingstone, 1st ed., 2010, 553.
66. Sharma SC, Yelne MB, Dennis TJ. Database on medicinal plants used in Ayurveda, CCRAS, New delhi, 2010, 418.
67. Jarald EE, Jarald ES. Textbook of Pharmacognosy & Phytochemistry. 1st Ed, New Delhi, 2007, 33-34.

68. Joshi JDS. Chemistry of ayurvedic crude drugs: Part VIII: Shatavari: 2. Structure elucidation of bioactive shatavarin I and other glycosides. *Indian J Chem B*, 27(1), 1988, 12-16.
69. Nair AGR, Subramanian SS. Occurrence of diosgenin in *Asparagus racemosus*. *Curr Sci*, 17, 1969, 414.
70. Wiboonpun N, Phuwapraisirisan P, Tip-pyang S. Identification of antioxidant compound from *Asparagus racemosus*. *Phytother. Res*, 8(9), 2010, 771-773.
71. Sharma SC. Constituents of the fruits of *Asparagus racemosus* Willd. *Pharmazie*, 36(10), 1981, 709.
72. Sekine TN, Fukasawa Structure of asparagamine A, a novel polycyclic alkaloid from *Asparagus racemosus*. *Chem Pharm Bull*, 42(6), 1994a, 1360-1362.
73. Sekine T, Fukasawa N, Kashiwagi Y, Ruangrunsi N, Murakoshi I. Structure of asparagamine A: A novel polycyclic alkaloid from *Asparagus racemosus*. *Chem Pharm Bull*, 42, 1994, 1360-2.
74. Sekine TN. TIFFNal Structure and relative stereochemistry of a new polycyclic alkaloid, asparagamine A, showing anti-oxytocin activity, isolated from *Asparagus racemosus*. *J Chem Soc*, 1, 1995, 391-393.
75. Bopana N and Saxena S. *Asparagus racemosus* ethnopharmacological evaluation and conservation needs. *J Ethnopharmacol*, 110(1), 2007, 1-15.
76. Dev S. Ancient-modern concordance in Ayurvedic plants: Some examples. *Environmental Health Perspectives*, 107(10), 1999, 783-789.
77. Dahanukar S, Thatte U, Pai N, Mose PB, Karandikar SM. Protective effect of *Asparagus racemosus* against induced abdominal sepsis. *Indian Drugs*, 24, 1986, 125-8.
78. Shankar K, Mitra, Neswi S, Prakash, and Ramachandran Sundaram. Shatavarins (containing Shatavarin IV) with anticancer activity from the roots of *Asparagus racemosus*. *Indian J Pharmacol*, 44(6), 2012, 732-736.
79. Bhatnagar M, Sisodia SS, Bhatnagar R. Antiulcer and antioxidant activity of *Asparagus racemosus* Willd and *Withania somnifera* Dunal in rats. *Ann N Y Acad Sci*, 1056, 2005, 261-78.
80. Karmakar SK, Sadhu SK, Biswas A, Chowdhury MC. Shill, Das J. Cytotoxicity, Analgesic and Antidiarrhoeal Activities of *Asparagus racemosus*. *J Appl Sci*, 12, 2012, 581-586.
81. Roy RN, Bhagwager S, Chavan SR, Dutta NK. Preliminary pharmacological studies on extracts of Root of *Asparagus racemosus* (Satavari), Willd, N.O. Liliaceae. *J Res Ind Med*, 6, 1971, 132-8.
82. Muruganadan S, Garg H, Lal J, Chandra S, Kumar D. Studies on the immunostimulant and antihepatotoxic activities of *Asparagus racemosus* root extract. *J Med Arom PI Sci*, 22, 2002, 49-52.
83. Kanitkar UK, Dange PS, Pendse GS. Pharmacognostic studies on a hitherto unrecorded form of *Asparagus racemosus* Willd. *J. Res. Indian Med*, 3, 1969, 123-37.
84. Sankarasubramanian. Introduction to natural drug. *Curr Sci*, 38, 1968, 414.
85. Sabin P, Gaitonde BB, Jetmalani M. Effects of alcoholic extracts of *Asparagus racemosus* on mammary glands of rats. *Ind J Exp Biol*, 6, 1968, 55-57.
86. Chintalwar G, Jain A, Sipahimalani A, Banerji A, Sumariwalla P, Ramakrishnan R and Sainis K. An immunologically active arabinogalactan from *Tinospora cordifolia*. *Phytochem*, 52(6), 1999, 1089-1093.
87. Backett AH, Stenlake JB. Practical Pharmaceutical Chemistry, 2, 1989, 86-96.
88. Hanuman J B, Bhatt RK, Sabata BK. A natural phenolic lignan from *Tinospora cordifolia* Miers. *J Chem Soc Perkin Trans*, 1, (1986b), 1181.
89. Anonymous. An Appraisal of Tribal-Folk Medicine. Central Council for Research in Ayurveda & Siddha, New Delhi, 1999.
90. Shah GL. Some Economically important plants of Salsette Island near Bombay. *J Econ Tax Bot*, 5, 1984, 753.
91. Raama Murthy J, Meera R, Venkataraman S, Bhojraj T satpute, Chidambaranathan N, Devi P. Phytochemical investigation and anticonvulsant activity of leaves of *Tinospora cordifolia* miers. *Int J Pharm Sc*, 2(2), 2010, 522-527.
92. Puranik N, Kammar KF, Sheela devi. Anti-diabetic activity of *Tinospora cordifolia* (Willd.) in streptozotocin diabetic rats; does it act like sulfonylureas? *Turk J Med Sci*, 40(2), 2010, 265-270.
93. Sharma AK, Singh RH. Screening of anti-inflammatory activity of certain indigenous drugs on carrageenin induced hind paw oedema in Rats. *Bull Medico Ethnobot Res*, 1(2), 1980, 12.
94. Patel SR, Goyal RK, Shah DS. Studies on the pharmacological effects of *Tinospora cordifolia*. *J Res Ind Med*, 13(2), 1977, 46.
95. Sarma DNK, Khosa RL, Chaurasia JPN, Sahai M. Antiulcer Activity of *Tinospora cordifolia* Meirs and *Centella asiatica* Linn. Extracts. *Phytother Res*, 9, 1995, 589.
96. Kulkarni SK, Verma A. An Herbal Preparation Improves Learning and Memory Performance in Mice. *Indian Drugs*, 30, 1993, 97.
97. Thatte UM, Chhabria S, Karandikar SM, Dahanukar SA. Immunotherapeutic modification of E.coli induced abdominal sepsis and mortality in mice by Indian medicinal plants. *Indian Drugs*, 25, 1987, 95-7.
98. Rege NN, Dahannkar SA, Karandikar SM. Hepatoprotective effects of *Tinospora cordifolia* against *CCl4* – induced liver damage. *Indian Drugs*, 21, 1984, 544-5.
99. Kalikar MV, Thawani VR, Varadpande UK, Sontakke SD, Singh RP, Khiyani RK. Immunomodulatory effect of *Tinospora cordifolia* extract in human immuno-deficiency virus positive patients. *Indian J Pharmacol*, 40, 2008, 107-10.