

A Review on Anthelmintic Plants

Ravindra G Mali and Anita A Mehta*

Department of Pharmacology

L. M. College of Pharmacy, Navrangpura, Ahmedabad-380 009, Gujarat, India

*Correspondent author, E-mail: dranitalmcp@rediffmail.com; Phone: 09275161224 (Mob.)

Received 27 July 2007; Accepted 30 October 2007

Abstract

Modern synthetic medicines are very effective in curing diseases but also cause a number of side effects. Crude drugs are less efficient with respect to cure of diseases but are relatively free from side effects. Parasites have been of concern to the medical field for centuries and the helminths considered causing considerable problems for human beings and animals. A large number of medicinal plants are claimed to possess anthelmintic property in traditional systems of medicine and are also utilized by ethnic groups worldwide. Following the folk claims, several medicinal plants have been scrutinized for this activity using various *in vitro* and *in vivo* methods. The present review summarizes some important pharmacological and preliminary studies on medicinal plants, products thereof and isolated principles from them, which can be investigated further to achieve lead molecules in the search of novel herbal drugs.

Keywords: Anthelmintic activity, Earthworms, Tapeworms, Hookworms, Essential oil, Medicinal Plants.

IPC code; Int. cl.⁸— A61K 36/00, A61P 33/10

high percentage of cure with a single therapeutic dose, free from toxicity to the host and should be cost effective. None of the synthetic drug available meets this requirement. Even most common drugs like Piperazine salts have been shown to have side effects like nausea, intestinal disturbances and giddiness³. Resistance of the parasites to existing drugs⁴ and their high cost warrants the search for newer anthelmintic molecules. The origin of many effective drugs is found in the traditional medicine practices and in view of this several researchers have undertaken studies to evaluate folklore medicinal plants for their proclaimed anthelmintic efficacy⁵.

Most of the screenings reported are *in vitro* studies using some worm samples like Indian earthworm *Pheretima posthuma*, *Ascardia galli*, *Ascaris lumbricoids*, etc. Adult Indian earthworm, *Pheretima posthuma* has been used as test worm in most of the anthelmintic screenings, as it shows anatomical and physiological resemblance with the intestinal roundworm parasite of human beings⁶⁻⁹. Because of easy availability, earthworms and *Ascardia galli* worms are used as suitable models for screening of anthelmintic drug¹⁰⁻¹⁹. These *in vitro* screenings are important as they give basis for further *in vivo* studies.

In the present review, various screening procedure and attempts made

Introduction

Helminthic infections are among the most common infections in human beings, affecting a large proportion of the world's population. In developing countries they pose a large threat to public health and contribute to the prevalence of anaemia, malnutrition, eosinophilia and pneumonia. Although the majority of infections due to worms are generally limited to tropical countries, they can occur to travelers, who have visited those areas and some of them can be developed in temperate climates¹. The helminthes which infect the intestine are cestodes e.g. Tapeworms (*Taenia solium*), nematodes e.g. hookworm (*Ancylostoma duodenale*), roundworm (*Ascaris lumbricoids*) and trematodes or flukes (*Schistosoma mansoni* and *Schistosoma hematobolium*). The diseases originated

from parasitic infections causing severe morbidity include lymphatic filariasis, onchocerciasis and schistosomiasis. These infections can affect most populations in endemic areas with major economic and social consequences. Helminthes also affect millions of livestock resulting in considerable economic losses in domestic and farm yard animals. Because of limited availability and affordability of modern medicines most of the world's population depends to a greater extent on traditional medical remedies. The traditional medicines hold a great promise as source of easily available effective anthelmintic agents to the people, particularly in tropical developing countries, including India. It is in this context that the people consume several plants or plant-derived preparations to cure helminthic infections². Ideally an anthelmintic agent should have broad spectrum of action,



Piliostigma thonningii



Butea monosperma



Cucurbita maxima



Trachyspermum ammi



Punica granatum



Capparis decidua



Capparis spinosa



Anacardium occidentale



Mimusops elengi bark



Cleome icosandra

by the researchers to evaluate the efficacy of plant species to discover new possible anthelmintic molecule(s) and to establish their possible mechanism (s) of action have been discussed.

***Ocimum sanctum* Linn.**
(Family-Lamiaceae)

Commonly known as Sacred Basil (*Tulsi*) contains volatile oil of which the chief constituents are Eugenol (about 51 %), β -caryophyllene (37%) and number of sesquiterpenes and monoterpenes²⁰. The essential oil and Eugenol showed

potent *in vitro* anthelmintic activity against *Caenorhabditis elegans* (Nematode). During experiment various concentrations of essential oil and Eugenol were tested using Levamisole as reference standard. Both exhibited ED₅₀ of 237.9 and 62.1 μ g/ml, respectively. Eugenol has been suggested as the putative anthelmintic principle²¹.

***Piliostigma thonningii* (Schum.) Milne-Redh.** (Family-Caesalpiniaceae)

Stem bark of this plant is used in traditional practices to treat dysentery,

snake-bite, toothache and as an anthelmintic²². Following the traditional claim, ethanol extract of the plant was screened for anthelmintic activity which exhibited a potent dose dependant activity against *Ascardia galli* infected cockrels by stimulating the neuromuscular junction principally and the ganglion to a lesser degree²³. In another study, an active principle D-3-O-Methylchiroinositol was isolated by bioassay-guided chromatographic separation technique from methanolic extract of stem bark of the plant and screened for anthelmintic activity by larval paralysis using Levamisole as a reference drug. Third stage larvae of *Haemonchus contortus* from faecal samples of infected lambs were used in the study. D-3-O-Methylchiroinositol induced approximately 60% larval paralysis within 24h at 4.4 mg/ml concentration²⁴.

***Melia azedarach* Linn.**

(Family-Meliaceae)

A native tree of Persia, India and China, this plant has long been recognized as an insecticidal and medicinal plant all over the world. The ethanolic extract of drupes was tested for its anthelmintic activity against the tapeworm *Taenia solium* (Cestoda) and the earthworm *Pheretima posthuma* (Annelida) using Piperazine phosphate as the standard drug. The extract was found active against both the tapeworm and the earthworm tested. Also the activity was better against tapeworm *Taenia solium*, than that of Piperazine phosphate¹³.

***Punica granatum* Linn.**

(Family-Punicaceae)

Locally known as Anar, is

cultivated in all parts of India. The root and stem bark of the plant is used as astringent and anthelmintic in the indigenous system of medicine. The alcoholic extract of its stem bark was evaluated for its proclaimed anthelmintic potential. The activity was found dose dependant, inhibiting transformation of eggs to filariform larvae of *Haemonchus contortus*²⁵. In clinical studies, the plant showed efficacy against nematodiasis in calves²⁶. The stem bark is reported to contain an alkaloid, Pelletierine.

***Moghania vestita* Kuntze syn.**

Flemingia vestita Benth.

(Family-Fabaceae)

A leguminous root crop commonly found in the North eastern regions of India. Its fleshy tuberous roots along with the peel are consumed raw by the tribal people of India to cure intestinal helminth infections. In a preliminary study the crude extract of the whole root tubers of this plant was reported to be effective against *Ascaris suum*, *in vitro*²⁷. Tegumental alterations and deformity were also observed in digenean flukes treated with the crude peel extract of the roots²⁸. Further in one study, root-tuber extract (50 mg/ml) and genistein (0.5 mg/ml), an active principle isolated from the root-tuber peel were tested against live parasites (Nematode: *Ascaris suum* from pigs, *A. lumbricoids* from humans, *Ascardia galli* and *Heterakis gallinarum* from domestic fowl; Cestode: *Railletina echinobothrida* from domestic fowl; Trematode: *Paramphistomum spp.* from cattle). The crude extract and genistein revealed complete immobilization of the trematode and cestode but not against cuticle-covered

nematodes. The treated parasites also showed structural alteration in their tegumental architecture. The activity of the peel extract and genistein is attributed to the changes induced in the tegumental integrity of the parasite¹⁸.

***Mimusops elengi* Linn.**

(Family-Sapotaceae)

In indigenous system of medicine the bark of *M. elengi* Linn. is reported to possess various therapeutic properties as cardiotoxic, stomachic, alexipharmic, anthelmintic and astringent²⁹. Various active principles such as taraxerol, taraxerone, ursolic acid, betulinic acid, α -spinosterol, β -sitosterol, alkaloid isoretronecyl tiglate and mixture of triterpenoid saponins have been reported from its stem bark. Crude alcoholic extract and its various fractions were evaluated for their anthelmintic potential using *Pheretima posthuma* (Annelida) and *Ascardia galli* (Nematode) as test worms. The crude alcoholic extract and its ethyl acetate and n-butanol fractions significantly demonstrated paralysis and also caused death of worms especially at higher concentration of 100 mg/ml as compared to standard reference Piperazine citrate (10mg/ml). The activity was attributed to the presence of polyphenolic compounds and tannins in the stem bark¹⁹. Tannins are believed to interfere with energy generation in helminth parasites by uncoupling oxidative phosphorylation and could bind to glycoprotein on the cuticle of the parasite, thereby may cause death³⁰.

***Calotropis procera* (Ait.) R. Br.**

(Family-Asclepiadaceae)

In traditional system of medicine

it is recommended for treatment of rheumatism, lupus, eczema, asthma, leprosy and syphilis³¹. The latex has been shown to possess anthelmintic activity against *Haemonchus contortus* infection in Najdi sheep in which it decreased the egg production and the number of worms in the abomasum and showed *in vitro* larvicidal activity³². Both fresh and aqueous extracts of dried latex of the plant were evaluated for their anthelmintic potential using earthworms as test worms. Both the extracts exhibited a dose-dependant inhibition of spontaneous motility and evoked responses to pin-prick. With higher doses (100 mg/ml of aqueous extract and 100 % fresh latex) the effects were comparable with that of 3% Piperazine. The study suggested that it might be effective against parasitic infections of both animals and humans caused by *Ostertagia*, *Nematodirus*, *Dictyocaulis*, *Taenia*, *Ascaris* and *Fasciola*¹⁴. The flowers of the plant were also evaluated for anthelmintic activity in comparison with levamisole through *in vitro* and *in vivo* studies and found to possess good activity against nematodes³³.

***Neolamarckia cadamba* (Roxb.) Bosser** (Family-Rubiaceae)

An ornamental plant traditionally used in stomatitis, eye inflammation and as an anthelmintic. It is reported to contain triterpenoids, alkaloids and saponins³⁴. Aqueous and ethanolic extracts of mature stem-bark of the plant were screened for anthelmintic potential against earthworms, tapeworms and roundworms using Albendazole as reference drug. The ethanolic extract was found potent than aqueous extract and activity was

comparable with the standard drug used in the study³⁵.

***Capparis decidua* Edgew.** (Family-Capparidaceae)

It is a struggling, glabrous shrub distributed throughout greater parts of India. In traditional system of medicine root bark is documented to be useful in asthma, cough, rheumatism, gout and helminthes infections. The anthelmintic activity of ethanolic extract of root bark of *C. decidua* Edgew. was evaluated against adult Indian earthworm *Pheretima posthuma* (Annelida) because of its anatomical and physiological similarity with roundworm parasite. The activity was found dose dependant, comparable with Piperazine citrate (10 mg/ml) at the higher concentration of 100 mg/ml of the extract used in the study³⁶. The root bark is reported to contain spermidine alkaloids³⁷. In another study ethanolic extract of root bark of *C. spinosa* Linn. was evaluated for its anthelmintic potential and has shown good anthelmintic activity against earthworm *Pheretima posthuma*. The activity was found dose dependant¹⁵.

***Xylopia aethiopica* A. Rich.** (Family-Annonaceae)

The seeds of this plant are used commonly in Nigeria by traditional herbalists to control gastrointestinal helminth parasites. To verify the claim, anthelmintic effect of the crude methanol extract of seeds was evaluated in rats experimentally infected with the rat hookworm *Nippostrongylus brasiliensis*. The plant exhibited activity at doses between 1.2 and 2.0g/kg as measured by reduction in worm counts at necropsy³⁸.

Tannins, flavonoids or terpenoids present in the crude extract of *X. aethiopica* A. Rich. were claimed to be responsible for its activity³⁹.

***Butea monosperma* (Lam.) Kuntze** syn. *B. frondosa* Koenig ex Roxb. (Family-Fabaceae)

The seeds are known to possess anthelmintic activity and their efficacy has been reported against ascarids⁴⁰, stomach worms of the sheep⁴¹, *Ascardia galli*⁴². In one study, isolated constituent palasonin, a lactone (C₁₆ H₂₂ O₆) from seeds was experimentally evaluated and found to possess significant anthelmintic property⁴³. Palasonin was also screened and found to be effective against *Ascaris lumbricoids*⁴⁴ and *Fasciola hepatica*⁴⁵. In another study, the biochemical mechanism of anthelmintic action of palasonin has been investigated on *Ascardia galli*. Palasonin inhibited the glucose uptake and depleted the glycogen content in the presence of glucose indicating that palasonin affects the energy generating mechanism of the parasite. It also significantly increased lactic acid suggesting inhibition of ATP production. The results indicated that palasonin may act via either inhibition of energy metabolism and/ or alteration in the motor activity of the parasite⁴⁶.

***Gynandropsis gynandra* (Linn.) Briq.** (Family-Capparidaceae)

Methanol extracts of leaves and stems of *G. gynandra* (Linn.) Briq. were investigated for their anthelmintic activity using *Fasciola gigantica* (liverfluke), *Taenia solium* (Tapeworm) and *Pheretima posthuma* (Earthworm) as test worms. Both the extracts exhibited

considerable activity in dose dependant manner and the order of sensitivity of the extracts to the worms were *P. posthuma* > *F. gigantea* > *T. solium*. The methanol extract of stem was found to be more active and showed significant activity compared to Piperazine citrate, standard reference drug used in the study⁴⁷.

***Evolvulus alsinoides* Linn.** (Family-Convulvaceae)

It is widely used in Ayurveda as powerful brain stimulant, aphrodisiac and anthelmintic. Ethanolic extract of the whole plant material was screened to validate its anthelmintic activity using adult Indian earthworm, *Pheretima posthuma* as model animal. The extract caused paralysis followed by death of the worms at all tested dose levels. At higher concentration of 100mg/ml the ethanolic extract was found to be more potent than the reference control Piperazine citrate¹².

***Centratherum anthelminticum* Kuntze** syn. *Vernonia anthelmintica* Willd. (Family-Asteraceae)

Commonly known as *Kalijiri* is reputed in Ayurvedic system of medicine as anthelmintic. Various extracts of seeds have been evaluated for their proclaimed anthelmintic activity by *in vivo* and *in vitro* methods. The study was conducted using *Fasciolopsis buski* and *Ascaris lumbricoides* and *Hymenolepis nana* as test worms. Alcoholic extract was found to possess maximum activity followed by ether extract while aqueous extract did not show this activity⁴⁸. The activity was compared to levamisole. *In vitro*, studies revealed higher anthelmintic effects of methanolic extract as compared to aqueous extract on live

Haemonchus contortus as evident from their mortality. *In vivo*, screening showed a maximum reduction (73.9%) in faecal egg counts per gram with crude aqueous extract, in sheep naturally infected with gastrointestinal nematodes⁴⁹.

***Carica papaya* Linn.** (Family-Caricaceae)

The anthelmintic potential of the aqueous extract of Papaya was evaluated using roundworm *Ascaris lumbricoides* and *Ascardia galli* (Nematodes) as test parasites⁵⁰. The phytoprinciple benzyl isothiocyanate was isolated from the extract as it is an active principle responsible for anthelmintic activity. The metabolic pathways in general and carbohydrate pathways in particular and neuromuscular coordination are the major target sites of action of anthelmintic compounds⁵¹. The compound benzyl isothiocyanate exerted its action by inhibiting energy metabolism and by affecting motor activity of the parasites, as developed by *in vitro* studies⁵². In another study, benzyl isothiocyanate isolated from *C. papaya* Linn. seed extract was tested for anthelmintic activity by viability assay using *Caenorhabditis elegans* and was claimed as the chief anthelmintic agent⁵³.

***Nigella sativa* Linn.** (Family-Ranunculaceae)

Commonly known as *Kala Jira*, is a reputed plant in Indian system of medicine for its usefulness in a variety of ailments and possesses carminative, digestive, astringent and diuretic properties⁵⁴. The anthelmintic potential of essential oil of *N. sativa* Linn. was evaluated against earthworms, tapeworms,

hookworms and nodular worms and exhibited fairly good activity against earthworms and tapeworms. The activity against hookworms and nodular worms was comparable with that of hexyl resorcinol⁵⁵. The main active principles of *N. sativa* are thymoquinone, dithymoquinone-cymene and α -pinene.

***Semecarpus anacardium* Linn. f.** (Family-Anacardiaceae)

It is found throughout the hotter parts of India and its nuts are commonly known as *Bhilawa*. The various concentrations of anacardic acid, extracted from the oil of nuts and its sodium salt were tested for anthelmintic activity. Both have been found to be potent anthelmintic agent than piperazine citrate with the same concentration⁵⁶.

***Piper longum* Linn.** (Family Piperaceae)

The essential oil from the fruits of *P. longum* Linn. was screened for the anthelmintic activity against *Ascaris lumbricoides*. The experiment revealed that its oil has a definite paralytic action on the nerve-muscular preparation of *Ascaris lumbricoides*. The activity of oil was found to be greater than the piperazine citrate used as standard in the study⁵⁷.

***Commiphora mukul* (Hook. ex Stocks) Engl.** (Family-Burseraceae)

Guggul is one of the noted drugs in Ayurveda and Unani system. In recent times its demand in therapeutics has been substantially increased. The alcoholic extract and essential oil obtained from oleo-gum resin of *C. mukul* (Hook. ex Stocks) Engl. has shown good anthelmintic activity against hookworms and tapeworms. The activity was

comparable to that of reference standards piperazine phosphate and hexyl resorcinol used in the study⁵⁸.

***Trachyspermum ammi* Linn.**

(Family-Apiaceae)

The seeds, commonly known as *Ajowan*, are used as diuretic, analgesic, anthelmintic and for the treatment of asthma. In one study seed extract of the plant was screened for its anthelmintic property in sheep and it produced significant results⁵⁹. The crude aqueous and methanolic extracts of seeds of *T. ammi* Linn. were also evaluated for the ovicidal efficacy by egg hatch test (EHT) on *Haemonchus contortus* ova. Lethal concentrations (LC₅₀) values were found to be 0.1698 and 0.1828 mg/ml, respectively⁶⁰.

***Ficus insipida* Willd.**

(Family-Moraceae)

The latex of some *Ficus* spp. including *F. insipida* Willd. has been traditionally used as vermifuge in Central and South America. It has been observed that anthelmintic activity is due to a proteolytic fraction called ficin and confirmed by clinical trials⁶¹. In another study the anthelmintic activity of the latex of this species was investigated in NIH mice (dose 4ml/kg/day), naturally infected with *Syphacia obvelata*, *Aspicularis tetraptea* and *Vampirolepis nana*. But due to high acute toxicity with haemorrhagic enteritis and weak anthelmintic efficacy, the use of latex is not recommended⁶². A study on re-evaluation of risks with the use of *F. insipida* latex as a traditional anthelmintic remedy has also been conducted and declared unsafe for

human use⁶³.

***Cucurbita maxima* Duch.**

(Family-Cucurbitaceae)

Seeds of the plant are reputed in Ayurvedic system of medicine as an anthelmintic especially against tape worms. The aqueous, alcoholic and ethereal extracts of the seeds were tested *in vivo* and *in vitro* on trematodes, cestodes and nematodes. The order of decreasing potency of the extracts in the *in vitro* studies was aqueous, alcoholic and ether extract. The kymographic studies suggested that the seed extracts act by bringing about a decrease in the motility leading to temporary paralysis⁶⁴.

***Nicotiana tabacum* Linn.**

(Family-Solanaceae)

In vitro and *in vivo* anthelmintic activity of *N. tabacum* Linn. leaves was studied to rationalize its traditional use. Live *Haemonchus contortus* were used to assess the *in vitro* anthelmintic effect of a crude aqueous extract and methanol extract. For the *in vivo* studies both the extracts were administered in increasing doses (1.0-3.0g/kg) to sheep naturally infected with mixed species of gastrointestinal nematodes. The results of the study showed that both the extracts possess dose-dependant anthelmintic activity, justifying the use of plant in traditional system of medicine⁶⁵.

***Cleome icosandra* Linn. syn.**

C. viscosa Linn. (Family Capparidaceae)

Commonly known as *Hul-Hul*, is an annual common weed found all over the plains of India and throughout the

tropics of the world. In Ayurvedic system of medicine this plant is believed to have several medicinal properties such as stomachic, laxative, diuretic, anthelmintic and considered useful in skin diseases, itching, ulcers, leprosy, malarial fevers⁶⁶. The crude alcohol and aqueous extracts of the seeds of *C. viscosa* Linn. were investigated for their anthelmintic activity against *Pheretima posthuma* and *Ascaridia galli*. Various concentrations (10-100mg/ml) of each extract were tested in the bioassay, which involved determination of time of paralysis and time of death of the worms. Both the extracts exhibited considerable anthelmintic activity in dose dependant manner. The most significant activity was observed at highest concentration of 100 mg/ml against both types of worms⁶⁷.

***Cannabis sativa* Linn.**

(Family-Cannabinaceae)

The efficacy of crude extract of its leaves on the motility and morphology of *Fasciolopsis buski* was studied using scanning electron microscopy. *In vitro* treatment with 5, 10 and 20 mg/ml of crude extract in phosphate buffered saline caused paralysis of the worm followed by death and found to be more lethal than the commercial flukicide, Oxyclozanide⁶⁸.

***Trifolium repens* Linn.**

(Family-Fabaceae)

In folk medicine of the *Naga* tribes of India it is used as a deworming remedy. The anticestodal activity of *T. repens* Linn. was evaluated using experimental *Hymenolepis diminuta* infections in albino rats. Aerial shoot extract of the plant at the concentrations 200 and 500 mg/kg reduced the mean

fecal egg counts of *H. diminuta* by 47.72 and 54.59% and worm recovery rate by 60 and 40%, respectively. Praziquantal, the standard cestocidal drug, reduced the mean fecal egg count by 65.90% and worm recovery rate by 26.67%^(Ref. 69).

***Strobilanthes discolor* T. Anders.** (Family-Acanthaceae)

Use of its leaves in the treatment of intestinal worm infections is a common ethnobotanical practice in the Naga tribes of North eastern part of India. The anticestodal efficacy of leaf extract was investigated using *Hymenolepis diminuta*, rat experimental model. The effects of leaf extract were adjudged by monitoring the eggs per gram of faeces (EPG) counts and percentage worm recovery rates following treatment with methanol leaf extract of this plant to different groups of rats harbouring *H. diminuta* infections. The leaf extract showed significant reductions in EPG counts as well as in recovery of surviving worms at autopsy. A notable result of the extract's efficacy was observed against the larval stages of parasite, where no single worm was recovered at its 800 mg/kg dose administered twice daily for 3 days. Effects of plant extract on adult stages were almost comparable with that of a standard drug Praziquantel. The study suggested that the leaf extract of *S. discolor* possesses significant anticestodal activity and supported its use in the folk medicine⁷⁰.

***Acacia auriculaeformis* A. Cunn.** (Family-Mimosaceae)

The triterpenoid saponins isolated from funicles of this plant (Acaciaside A and Acaciaside B) were

tested for *in vitro* and *in vivo* activity against *Microfilariae* and adult *Setaria cervi*. The mixture of two saponins has shown good cestocidal activity⁷¹.

Miscellaneous studies

Some workers have reported anthelmintic activity of essential oils obtained from plants. In one study, the essential oil of *Piper betle* Linn. has revealed anthelmintic activity against the earthworm *Pheretima posthuma*, *in vitro*⁷². The anthelmintic activity of the essential oil obtained from *P. betle* cv. 'Sagar Bangla' was also tested against tapeworms and was found to be better than the standard piperazine phosphate and the activity against hookworm was found to be greater than the reference drug Hexyl resorcinol⁷³. Seed oils of *Gynandropsis gynandra* (Linn.) Briq., *Impatiens balsamina* Linn., *Celastrus paniculata* Willd., *Embelia ribes* Burm. f. and *Mucuna prurita* Hook. were investigated for their anthelmintic property against *Pheretima posthuma*. Three concentrations (10, 50 and 100 mg/ml) of each oil were studied for bioassay. *Embelia ribes* Burm. f. seed oil showed the best activity as compared to Piperazine citrate (10mg/ml) included as reference standard⁷⁴. The essential oils of *Anacardium occidentale* Linn. and *Callistemon viminalis* (Soland.) Cheel have been found to possess anthelmintic activity against earthworms and tapeworms. The activity was found to be better than reference standard Piperazine phosphate. The oils were also found to be effective against hookworms as compared to Hexyl resorcinol⁷⁵. The essential oil of *Gardenia resinifera* Roth. syn. *G. lucida* Roxb. was also

evaluated and showed better efficacy against *Taenia solium* at higher concentration⁷⁶.

The *in vitro* anthelmintic activity of the aqueous and alcoholic extracts of *Melia azedarach* Linn., *Ananas comosus* (Linn.) Merr. syn. *A. sativus* Schult. f., *Embelia ribes* Burm. f. and *Mucuna prurita* Hook. were evaluated against *Taenia canina* and *Phamphistomum cervi*. *M. prurita* was found more active against the trematodes⁷⁷. Ellagitannins and complex tannins isolated from the stem bark of *Quercus petraea* (Matt.) Liebl. showed good anthelmintic activity⁷⁸. In one study, different parts of ten indigenous medicinal plants were screened for their *in vitro* anthelmintic activity against *Ascaridia galli* worms of birds. Preparations from *Sapindus trifoliatus* Linn. and *Momordica charantia* Linn. were found more effective than Piperazine hexahydrate¹⁷. The active principle, solamargine, isolated from the ripe berries of *Solanum viarum* Dunal was found effective against *Microfilariae* and adults of *S. cervi*⁷¹. In one study, the ethanolic extract of *Centella asiatica* (Linn.) Urban was evaluated for antifilarial activity on canine dirofilariasis (*Dirofilaria immitis*)⁷⁹.

Conclusion

Ancient classical literature and ethnomedical surveys described the use of plants in traditional system of medicines for the treatment of helminthic infections. This traditional medical wisdom is excellent proof of clinical efficacy and safety of medicinal plants. Present report is a survey of literature

indicating the screenings of crude plant extracts, essential oils and isolated active principles for *in vitro* and *in vivo* anthelmintic studies to substantiate the folk claim. To conclude, in future studies, there is need for thorough phytochemical, clinical and possible studies on molecular mechanism of action. At the same time efforts should be made to standardize the plant extracts with good anthelmintic activity and formulate best alternative herbal preparations to replace or complement the synthetic drugs which are currently in use.

References

- Bundy DAP, Immunoepidemiology of intestinal helminthic infection, *Trans Royal Soc Trop Med Hygiene*, 1994, **8**, 259-261.
- Satyavati GV, Use of plant drugs in Indian Traditional System of Medicine and their relevance to primary health care, *In: Economic and Medicinal Plant Research by Farnworth NR and Wagner H (Eds)*, Academic Press Ltd, London, 1990, pp.190-210.
- Liu LX and Weller PF, An update on antiparasitic drugs, *N Engl J Med*, 1996, **334**, 1178.
- Walter PJ and Prichard KK, Chemotherapy of parasitic infections, *In: W.C. Campbell and L.S. Rew (Eds)*, Plenum, New York, 1985, pp. 278-539.
- Temjenmongla and Yadav A, Anticestodal efficacy of folklore medicinal plants of Naga tribes in North-East India, *Afr J Trad Cam*, 2005, **2**(2)129-133.
- Vidyarthi RD, A Textbook of Zoology, S. Chand and Co, New Delhi, 14th Edn, 1967, pp. 329-370.
- Vigar Z, Atlas of Medical Parasitology, P.G. Publishing House, Singapore, 2nd Edition, 1984, p.216.
- Thorn GW, Adams RD, Braunwald E, Isselbacher KJ and Petersdorf RG, Harrison's Principals of Internal Medicine, McGraw Hill Co, New York, 1977, p. 1088.
- Chatterjee KD, Parasitology, Protozoology and Helminthology, Guha Ray Sree Saraswathy Press Ltd, Calcutta, 1967, pp.168-169.
- Sollmann T, Anthelmintics: Their efficiency as tested on earthworms, *J Pharmacol Exp Ther*, 1918, **12**, 129-170.
- Jain ML and Jain SR, Therapeutic utility of *Ocimum basilicum* var. *album*, *Planta Med*, 1972, **22**, 66-70.
- Dash GK, Suresh P, Sahu SK, Kar DM, Ganapaty S and Panda SB, Evaluation of *Evolvulus alsinoides* Linn. for anthelmintic and antimicrobial activities, *J Nat Rem*, 2002, **2**(2), 182-185.
- Szewezuk VD, Mongelli ER and Pomilio AB, Antiparasitic activity of *Melia azedarach* growing in Argentina, *Mol Med Chem*, 2003, **1**, 54-57.
- Shivkar YM and Kumar VL, Anthelmintic activity of latex *Calotropis procera*, *Pharm Biol*, 2003, **41**(4), 263-265.
- Mali RG, Mahajan S and Patil KS, Anthelmintic activity of root bark of *Capparis spinosa*, *Indian J Nat Prod*, 2005, **21**(4), 50-51.
- Kaushik RK, Katiyar JC and Sen AB, Studies on the mode of the action of anthelmintics with *Ascaridia galli* as a test parasite, *Indian J Med Res*, 1974, **62**, 1367-75.
- Lal J, Chandra S, Raviprakash V and Sabir M, *In vitro* anthelmintic action of some indigenous medicinal plants on *Ascaridia galli* worms, *Indian J Physiol Pharmacol*, 1976 **20**(2), 64-68.
- Tandon V, Pal P, Roy B, Rao HSP and Reddy KS, *In vitro* anthelmintic activity of root-tuber extract of *Flemingia vestita*, an indigenous plant in Shillong, India, *Parasitol Res*, 1997, **83**, 492-498.
- Mali RG, Mahajan SG and Mehta AA, *In vitro* anthelmintic activity of stem bark of *Mimusops elengi* Linn, *Pharmacog Mag*, 2007, **3**(10), 73-76.
- Handa SS and Kapoor VK, Pharmacognosy, Vallabh Prakashan, New Delhi, 1988, pp.240-245.
- Asha MK, Prashant D, Murali B, Padmaja R and Amit A, Anthelmintic activity of essential oil of *Ocimum sanctum* and Eugenol, *Fitoterapia*, 2001, **72**, 669-670.
- Lewis WH and Elvin-Lewis MPF, Medical Botany, John Wiley and Sons, New York, 1979, pp. 236-261.
- Asuzu IU and Onu OU, Anthelmintic activity of the ethanolic extract of *Piliostigma thonningii* bark in *Ascaridia galli* infected chickens, *Fitoterapia*, 1994, **65**(4), 291-297.
- Asuzu IU, Gray AI and Waterman PG, The anthelmintic activity of D-3-O-methylchiroinositol isolated from *Piliostigma thonningii* stem bark, *Fitoterapia*, 1999, **70**, 77-79.
- Prakash V, Singhal KC and Gupta RR, Anthelmintic activity of *Punica granatum* and *Artemisia silversiana*, *Indian J Pharmacol*, 1980, **12**, 62-65.
- Pradhan KD, Thakur DK and Sudhan NA, Therapeutic efficacy of *P. granatum* and *C. maxima* against clinical cases of nematodiasis in calves, *Indian J Ind Med*, 1992, **9**(1), 53-54.
- Yadav AK, Tandon V and Rao HSP, *In vitro* anthelmintic activity of fresh tuber extract of *Flemingia vestita* against *Ascaris suum*, *Fitoterapia*, 1992, **63**, 395-398.
- Roy B and Tandon V, Effect of root-tuber extract of *Flemingia vestita*, a leguminous plant, on *Artyfechinostomum sufratyfex* and *Fasciolopsis buski*: a scanning electron microscopy study, *Parasitol Res*, 1996, **82**, 248-252.
- Kirtikar KR and Basu BD, Indian Medicinal

- Plants, M/S Bishen Singh Mahendra Pal Singh, Dehra Dun, 2nd Edition, 1935, pp.1494-1496.
30. Thompson DP and Geary TG, The structure and function of helminth surfaces. *In: Biochemistry and Molecular Biology of Parasites*, Marr, JJ (Ed), Academic Press, New York, 1st Edn, 1995, pp. 203-232.
31. The Wealth of India—A Dictionary Indian Raw Materials Vol. 3, Revised series, Publication and Information Directorate, CSIR, New Delhi, 1992, pp.78-84.
32. Al-Qarawi AA, Mahmoud OM, Sobaih, Haroun EM and Adam SE, A preliminary study on the activity of *Calotropis procera* latex against *Haemonchus contortus* infection in Najdi sheep, *Vet Res Commun*, 2001, **25**, 61-70.
33. Iqbal Z, Lateef M, Jabbar A, Muhammad G and Khan MN, Anthelmintic activity of *Calotropis procera* Ait. flowers in sheep, *J Ethnopharmacol*, 2005, **102**(2), 256-261.
34. Kaushik P and Dhiman A, Medicinal Plants and Raw Drugs of India, Orient Longman, Kottakal, 1999, pp.304-310.
35. Gunasekaran R, Senthilkumar KL and Gopalkrishnan S, Anthelmintic activity of bark of *Neolamarckia cadamba*, *Indian J Nat Prod*, 2006, **22**(1), 11-13.
36. Mali RG, Hundiware JC, Sonawane RS, Patil RN and Hatapakki BC, Evaluation of *Capparis decidua* for anthelmintic and antimicrobial activities, *Indian J Nat Prod*, 2004, **20**(4), 10-13.
37. Ahmed VU, Ismail N, Arif S and Amber A, Two new N-acetylated spermidine alkaloids from *Capparis decidua*, *J Nat Prod*, 1992, **55**(10), 1509-1512.
38. Suleiman MM, Mamman M, Aliu YO and Ajanusi JO, Anthelmintic activity of the crude methanol extract of *Xylopiya aethiopica* against *Nippostrongylus brasiliensis* in rats, *Vet Arhiv*, 2005, **75**(6), 487-495.
39. Lahlou M, Potential of *Origanum compactum* as a cercaricide in Morocco, *Ann Trop Med Parasitol*, 2002, **96**, 587-593.
40. Ramanan MV, *Butea frondosa* seeds in round worm infestation, *Antiseptic*, 1960, **57**, 927-928.
41. Garg LC and Mehta RK, *In vitro* studies on anthelmintic activity of *Butea frondosa* and *Embllica ribes*, *J Anim Husb Res*, 1958, **3**, 28-32.
42. Satyanarayanrao V and Krishnaiah KS, Note on comparative efficacy of some indigenous anthelmintics against *A. galli* infection in chicks, *Indian J Anim Sci*, 1982, **52**, 485-486.
43. Raj RK and Kurup PA, Isolation and characterization of palasonin, an anthelmintic principle from the seeds of *Butea frondosa*, *Indian J Chem*, 1967, **5**, 86-87.
44. Rao KS, Raviprakash V, Chandra S and Sabir M, Anthelmintic activity of *Butea frondosa* against *Ascaris lumbricoids*, *Indian J Physiol Pharmacol*, 1977, **21**, 250-253.
45. Sabir M, Lal J, Raviprakash V, Chandra S and Rao KS, Anthelmintic effect of *Butea frondosa* seeds, *Proc Decennial Conf Indian Pharmacol Soc*, 1977, 103.
46. Kumar D, Mishra SK, Tandan SK and Tripathi HC, Possible mechanism of anthelmintic action of palasonin on *Ascardia galli*, *Indian J Pharmacol*, 1995, **27**(3), 161-166.
47. Ajaiyeoba EO, Onocha PA and Olarenwaju OT, *In vitro* anthelmintic properties of *Buchholzia coriacea* and *Gynandropsis gynandra* extracts, *Pharm Biol*, 2001, **39**(3), 217-220.
48. Singh S, Ansari NA, Srivastava MC, Sharma MK and Singh SN, Anthelmintic activity of *Vernonia anthelmintica*, *Indian Drugs*, 1985, **22**(10), 508-511.
49. Iqbal Z, Lateef M, Jabbar A, Muhammad G and Khan MN, Anthelmintic activity of *Vernonia anthel* seeds against *Trichostrongylid* nematodes of sheep, *Pharm Biol*, 2006, **44**(8), 563-567.
50. Dhar RN, Garg LC and Pathak RD, Anthelmintic activity of *Carica papaya* seeds, *Indian J Pharm*, 1965, **27**(12), 335-336.
51. Sharma S, Treatment of helminth diseases—Challenges and achievements. *In: Progress in Drug Research*, Jucker, E (Ed), Birdhauser Verlag, Boston, 1987, pp.69-100.
52. Kumar D, Mishra SK, Tandan SK and Tripathi HC, Mechanism of anthelmintic action of benzyl isothiocyanate, *Fitoterapia*, 1991, **62** (5), 403-410.
53. Kermanshah R, Brian E, Rosenfeld J, Summers PS and Sorger GJ, Benzyl isothiocyanate is the chief or sole anthelmintic in papaya seed extracts, *Phytochemistry*, 2001, **57**(3), 427-435.
54. Nadkarni AK, Indian Materia Medica, Popular Book Depot, Bombay, 1960, pp. 584.
55. Agrawal R, Kharya MD and Shrivastava R, Antimicrobial and anthelmintic activities of the essential oil of *Nigella sativa* Linn, *Indian J Exp Biol*, 1979, **17**, 1264-1265.
56. Chattopadhyaya MK and Khare RL, Isolation of anacardic acid from *Semecarpus anacardium* Linn. and study of its anthelmintic activity, *Indian J Pharm*, 1969, **31**(4), 104-105.
57. D'Cruz JL, Nimbkar AY and Kokate CK, Evaluation of fruits of *Piper longum* Linn. and leaves of *Adhatoda vasica* Nees. for anthelmintic activity, *Indian Drugs*, 1980, **17**(4), 99-101.
58. Kakrani HK and Kalyani GA, Anthelmintic activity of the essential oil of *Commiphora mukul*, *Fitoterapia*, 1984, **55**(4), 232-234.
59. Lateef M, Iqbal Z, Akhtar MS, Jabbar A, Khan MN and Gilani AH, Preliminary screening of *Trachyspermum ammi* (L) seeds for anthelmintic activity in sheep, *Trop Anim Health Prod*, 2006, **38**(6), 491-496.
60. Jabbar A, Iqbal Z and Khan MN, *In vitro* anthelmintic activity of *Trachyspermum ammi* seeds, *Pharmacog Mag*, 2006, **2**(6), 126-128.

61. Phillips O, *Ficus insipida*: Ethnobotany and ecology of an Amazonian anthelmintic, *Econ Bot*, 1990, **44**(4), 534-536.
62. De Morin A, Borba HR, Carauta JP, Lopes D and Kaplan MA, Anthelmintic activity of the latex of *Ficus* species, *J Ethnopharmacol*, 1999, **64**(3), 255-258.
63. Hansson A, Zelada JC and Noriega HP, Reevaluation of risks with the use of *Ficus insipida* latex as a traditional anthelmintic remedy in the Amazon, *J Ethnopharmacol*, 2005, **98**(3), 251-257.
64. Srivastava MC and Singh SW, Anthelmintic activity of *Cucurbita maxima* (Kaddu) seeds, *Indian J Med Res*, 1967, **55**(6), 629-632.
65. Iqbal Z, Lateef M, Jabbar A, Muhammad G and Gilani HA, *In vitro* and *in vivo* anthelmintic activity of *Nicotiana tobacum* Linn. leaves against gastrointestinal nematodes of sheep, *Phytother Res*, 2006, **20**(1), 46-48.
66. Chatterjee A and Prakash SC, *The Treatise on Indian Medicinal Plants*, Council for Scientific and Industrial Research, New Delhi, 2nd Edn, 1991, pp.155-160.
67. Mali RG, Mahajan SG and Mehta AA, *In vitro* anthelmintic screening of *Cleome viscosa* extract for anthelmintic activity, *Pharma Biol*, 2007, **45**(10), 766-768.
68. Roy B and Tandon V, *In vitro* flucicidal effect of leaf extract of *Cannabis sativa* Linn. on the trematode *Fasciolopsis buski*, *Indian J Exp Biol*, 1997, **35**(1), 80-82.
69. Tangpu V, Temjenmongla K and Yadav A, Anticestodal activity of *Trifolium repens* extract, *Pharm Biol*, 2004, **42**(8), 656-658.
70. Tangpu V, Temjenmongla K and Yadav A, Anticestodal property of *Strobilanthes discolor*: An experimental study in *Hymenolepis diminuta*—rat model, *J Ethnopharmacol*, 2006, **105**(3), 459-463.
71. Sinha Babu SP, Saponins and its possible role in the control of helminth parasites. *In: Recent Progress in Medicinal Plants*, Sharma SK, Govil JN and Singh VK (Editors) Studium Press LLC, USA, Vol. 10, 2005, pp. 405- 418.
72. Ali SM and Mehta RK, Preliminary pharmacological and anthelmintic studies of the essential oil of *Piper betle* Linn., *Indian J Pharm*, 1970, **32**(5), 132-133.
73. Garg SS and Jain R, Biological activity of the essential oil of *Piper betle* Linn. cv.' Sagar Bangla', *J Essent Oil Res*, 1992, **4**(6), 601-606.
74. Jalalpure SS, Alagawadi KR, Mahajanshetti CS, Shah BN, Singh V and Patil JK, *In vitro* anthelmintic property of various seed oils against *Pheretima posthuma*, *Indian J Pharma Sci*, 2007, **69**(1), 158-160.
75. Garg SC and Kasera HL, *In vitro* anthelmintic activity of the essential oil of *Anacardium occidentale*, *Indian Perfum*, 1982, **26**, 239-240.
76. Girgune JB, Jain NK and Garg BD, Antimicrobial and anthelmintic activity of essential oil from *Gardenia lucida* Roxb., *Indian Perfum*, 1979, **23**(3&4), 213-215.
77. Neogi NC, Baliga PAC and Srivastava RK, Anthelmintic activity of some indigenous drugs, *Indian J Pharm*, 1964, **26**, 37-39.
78. Konig MH, Scholz H, Hartmann R, Lehmann W and Rimple H, Ellagitannins and complex tannins from *Quercus petraea* bark, *J Nat Prod*, 1994, **57**(10), 1411-1415.
79. Chakraborty T, Sinha Babu and Sukul NC, Preliminary evidence of antifilarial effect of *Centella asiatica* on canine dirofilariasis, *Fitoterapia*, 1996, **2**, 110-112.