



INSECT PESTS AND DISEASES OF MEDICINAL AND AROMATIC PLANTS AND THEIR MANAGEMENT

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Medicinal and Aromatic Plants (MAPs) are the rich and major source for plant secondary metabolites. Huge world human population especially in developing nations mainly depend on traditional system of medicine for the treatment of various human ailments. At the moment traditional medicine derived from different MAPs have gained more importance and have become one of the useful ways to support herbal system of medicine in most of the countries like India, Japan, China, Indonesia, Pakistan, Bangladesh, Sri Lanka, Thailand, Malaysia etc. Similarly aroma compounds from botanical sources hold a promising field by being increasingly used in perfumery, nutraceuticals, food and flavour industries due to the growing awareness in common masses about the risks involved in synthetic components in parallel products. Accordingly, nearly more than one lakh such secondary metabolites are now known to occur in about 50,000 plant species and almost 4,000 new secondary metabolite are being discovered every year from a variety of plant species (Verpoorte et al. 1999; Gomez-Galera et al. 2007, Pandey, 2017). Since last several decades, these natural plant products have been utilized for human healthcare in the form of drugs, antioxidants, flavors, fragrances, dyes, pesticides, pheromones etc. During the last two decades, however, the use of synthetic drugs led to a decline in the use of plant-derived compounds, so that at one time it was believed that the synthetic drugs would perhaps completely replace the use of traditional plant-derived medicines. However, in recent years, a resurgence of the use of herbal drugs has once

again been witnessed prominently, firstly because the synthetic drugs have been found to be hazardous in many cases, and secondly because there is growing awareness that the plant-derived medicines have negligible side effects that are so common in the case of synthetic drugs. During COVID-19 outbreak there is an intensive search for plant derived molecules in order to find a cure for this pandemic. In such a situation plant based anti-malaria drug Chloroquine is playing a significant role to cure this deadly disease. Consequently, the current global herbal drug market has reached a level of more than US \$62 billion, which is expected to grow to US \$5 trillion by the year 2050 (Joshi et al. 2004). The world market for herbal medicines including herbal products and raw materials is actually growing at an annual rate of 10–15%. This is an indication of a possible growing demand for plant-derived drugs in coming years. Despite this, although the area under cultivation of medicinal and aromatic plants have been slowly increasing due to increasing demand for herbal drugs. The use of medicinal plant species for human healthcare can not be avoided due to the following reasons: (1) low concentration of desired molecules in specific plant tissues; (2) medicinal crops being low priority crops; (3) technical difficulty in cultivation of these crops; (4) over-harvesting of naturally occurring herbal plants; (5) destructive collection techniques; and (6) conversion of habitats of medicinal plants to crop-based agriculture. The business of medicinal and aromatic plant industries could be enhanced by ensuring the disease free plants

Table 1. Insect pests of *Mentha* spp

Common name	Scientific name	Order	Family	Pest Status	References
Bihar hairy caterpillar	<i>Spilosoma obliqua</i>	Lepidoptera	Arctiidae	Major	Singh, 1999
Soybean Semilooper	<i>Thysanoplusia orichalcea</i>	Lepidoptera	Noctuidae	Major	Sagar & Ramji, 1991
Tobacco caterpillar	<i>Spodoptera litura</i>	Lepidoptera	Noctuidae	Major	Kattimani, 2000
Variegated cutworm*	<i>Peridroma saucia</i>	Lepidoptera	Noctuidae	Major	Berry and Shields, 1980
Black cutworm	<i>Agrotis ipsilon</i>	Lepidoptera	Noctuidae	Major	Singh, 1999
Mentha leaf roller	<i>Syngamia abruptalis</i>	Lepidoptera	Pyralidae	Major	Kareem <i>et al.</i> 1960; Sagar and Reddy, 1987; Singh, 1999
Mint root borer*	<i>Fumibotys fumalis</i>	Lepidoptera	Crambidae	Major	Morris, 2007
Cutworm	<i>Spodoptera exigua</i>	Lepidoptera	Noctuidae	Major	Morris, 2007; Singh, 1999
Red pumpkin beetle	<i>Aulucophora foveollis</i>	Coleoptera	Chrysomelidae	Major	Singh and Gupta, 1970
Mint flea beetle*	<i>Longitarsus ferrugineus</i>	Coleoptera	Chrysomelidae	Major	Morris, 1990
Cotton whitefly	<i>Bemisia tabaci</i>	Hemiptera	Aleyrodidae	Major	Singh, 1999
Green peach aphid	<i>Myzus persicae</i>	Hemiptera	Aphididae	Minor	Neubauer <i>et al.</i> , 1974
Aphids	<i>Aphis affinis</i>	Hemiptera	Aphididae	Major	Singh and Sagar, 1981; Singh, 1999
Mint aphid	<i>Ovatus crataegarius</i>	Hemiptera	Aphididae	Major	Leonard, 1980
Citrus Mealybug	<i>Planococcus citri</i>	Hemiptera	Pseudococcidae	Minor	Ahmed and Abd-Rabou, 2010
Lace bug	<i>Cochlochila bullita</i>	Hemiptera	Tingidae	Minor	Samuel, 1939; Sharga, 1953; Singh, 1999
Two-spotted spider mite	<i>Tetranychus urticae</i>	Trombidiformes	Tetranychidae	Major	Morris, 2007
Termites	<i>Odontotermes obesus</i>	Isoptera	Termitidae	Major	Singh, 1999

* Not present in India

and planting materials. Among the different plant pathogens, fungi, bacteria, viruses, phytoplasma, plant parasitic nematodes and large number of insects are the major are serious threat to the future prospects of medicinal and aromatic plants globally. The cultivation of various medicinal and aromatic plants has been severely affected by these pests and pathogens. Improvement of the yield and quality of plant secondary metabolites and products through disease management is one of the major tasks.

MINTS (*Mentha* spp): Mints are group of aromatic herb belonging to family Lamiaceae, which are considered to be the most important cash crop in Indo-Gangetic plains. Different types of mints, which are commercially cultivated in Indo-Gangetic plains, are: Menthol mint (*Mentha arvensis*), Peppermint (*Mentha piperita*), Spearmint (*Mentha spicata*), Scotch spearmint (*Mentha cardiaca*), Bergamot mint (*Mentha citrata*) and Garden mint (*Mentha viridis*). Among different

medicinal and aromatic plants, mints come in the front line because Indian farmers grow it as a bonus crop and it also fits well in the cropping system with other crops like paddy, wheat, potato, sugar cane, etc. Now it is cultivated on an area of more than 1,50,000 ha of land.

Insect pests of *Mentha* spp. (Table 1)

Bihar Hairy Caterpillar, *Spilosoma obliqua* Walker, (Lepidoptera: Arctiidae):

Distribution: Occurrence of *S. obliqua* has been reported from Bangladesh, Sri Lanka, Pakistan, Philippines and Myanmar. In India the occurrence in Northern and North Eastern province is common.

Host plants: *Spilosoma obliqua* is a polyphagous pest damaging many crops including *Mentha* spp. sesamum, sunflower, cowpea, jute, sunnhemp and other pulse crops.

Identification and Biology: The adult moths are light brown in colour. The female moth lays

Figure 1: *Spilosoma obliqua*

yellowish green eggs in clusters on the underside of leaves. The incubation period is 5-13 days. The larvae complete development in 4-8 weeks through 6-7 instars. The early three instars feed gregariously and later instars feed solitarily. The larvae are yellowish orange and covered with hairs (Fig.1). The full grown larvae spin a loose silken cocoon in which it pupates in plant debris or in the soil. The pupal duration of is 10-15 days. The adult moth longevity is 5-7 days. The total life cycle is completed in 6-12 weeks. The Bihar hairy caterpillar passes through three to four generations in a year.

Nature of Damage: The young larvae feed gregariously on the lower surface of leaves and apical the apical shoots. The damaged leaves by young caterpillars appear dusty white colour which can be easily recognized. Later the grownup larvae disperse to the entire field and start feeding voraciously on leaves. Under severe occurrence of the pest complete defoliation may occur.

Management

- Grow insect-tolerant cultivar “Sambhav” of *Mentha arvensis* (Khanuja *et al.* 2004).
- Hand collection and destruction of gregariously feeding larvae of the pest.
- Natural enemies, parasitoids (*Meteorus pilosomae* and *Protapantales obliquae*) and entomopathogens (*Bacillus thuringiensis*, *Entomophaga aulicae* and Nuclear Polyhedrosis Virus) play important role in the control of *Spilosoma obliqua* (Gotyal *et al.*,

Figure 2: Soybean semilooper, *Thysanoplusia orichalcea*

2019; Senthil *et al.* 2011).

- Spray insecticides, like quinalphos 25EC (2ml/l) or cypermethrin 10EC (1.3ml/l).
- Install light traps (1trap/acre) in the field monitoring and attracting the moths of the pest.

Soybean semilooper, *Thysanoplusia orichalcea* (Fabricius) (Lepidoptera: Noctuidae)

Distribution: *T. orichalcea* is widely distributed in the world. The pest is present in Sri Lanka, Africa, India, Europe, New Zealand and Australia.

Host plants: Soybean semilooper is polyphagous pest. The pest in addition to *Mentha* spp., feeds on plant belonging to families Chenopodiaceae, Cucurbitaceae, Compositae, Leguminosae, Cruciferae, and Linaceae (Singh and Rawat 1980; Sagar and Ramji 1991).

Identification and Biology

T. orichalcea moth is medium sized and brown in colour. The forewings of moth are brown with golden colour patch (Fig.2). The hindwings are grayish brown. The caterpillars are green in colour with white stripes on the body and full grown caterpillar measures about 45 mm in length. Caterpillars while moving form distinct looping action. Female moth lays 65-120 yellowish eggs on the lower surface of leaves singly. Incubation duration ranges



Figure 3: *Spodoptera litura*

between 3 to 4 days. The caterpillar passes through 6 larval instars and larval period ranges between 16-23 days. The full grown larva pupates in loose silken cocoon on the lower side of leaves. The pupal period varies from 6 to 9 days. Life cycle is completed in 30-35 days.

Nature of Damage: The freshly hatched larvae feed on the lower surface of leaves by scrapping the epidermal part of leaves. The grown up larvae feed voraciously on leaves and make holes. The pest is active during March to October (Sagar, 1985). Severe infestation affects the vegetative growth of the plant, thereby affecting the yield of *Mentha* spp.

Management

- Foliar application of *Bacillus thuringiensis* var. kurstaki, serotype H-39, 3B, Strain Z-52 (0.75–1.0 kg/ha) for the management semilooper.
- Install light traps in the field for monitoring and collection of adult moths.
- Conservation of natural enemies (*Apanteles ruficrus* and *A. glomeratus*) by minimizing use of broad spectrum pesticides (Sagar and Ramji, 1990).
- Foliar application of quinalphos 50EC (2ml/l) or monocrotophos 36SL (1.5ml/l) is effective for control of semilooper (Sagar, 1988; Tripathi, 2018).

Tobacco caterpillar, *Spodoptera litura* (Fabricius) (Lepidoptera: Noctuidae)

Distribution

Tobacco caterpillar is one of the most important insect pests of agricultural crops in the tropical

and temperate Asia, Australasia and the Pacific Islands (EPPO 2013).

Host plants: *Spodoptera litura* is a polyphagous pest damaging sunflower, cabbage, cotton, pulses, soybean, tobacco, castor, okra, groundnut etc. including *Mentha* spp.

Identification and Biology: The adult moth is medium sized dark brown with grayish white markings on forewings and hindwings are white in colour (Fig. 3). The female lays 300 spherical eggs in mass on the surface of leaves and covers them with scales. The incubation period ranges between 3 to 5 days. Full grown larvae are velvety black with yellow dorsal stripes and lateral white strips. Full grown larvae measures about 35-40 mm in length. The larva passes through 5 stages and the larval period is completed in 15-30 days. The full grown up larvae enter soil for pupation. The pupal duration lasts for 7-15 days. Life cycle is completed within 32-60 days. The pest completes 8 generations in a year.

Nature of Damage: Freshly emerged caterpillars feed gregariously on the leaves from the low surface by scrapping the leaf lamina. Later on the larvae feed voraciously on leaves by making irregular holes in solitary phase. The larvae remain hidden in cracks, crevices and plant debris in soil during daytime.

Management

- Hand collection and destruction of egg

masses and gregariously feeding larvae of *S. litura*.

- Castor or sunflower plants can be grown as trap crops with main crop to attract female moths of *S. litura* for egg laying.
- Foliar application of SINPV (*Spodoptera litura* nuclear polyhedrosis virus) at 250 LE/ha for *S. litura*.
- Foliar application of *Bacillus thuringiensis* var. kurstaki, serotype H-39, 3B, Strain Z-52 (0.75–1.0 kg/ha) to control larvae of *S. litura*.
- Spray 5% neem seed kernel extract (NSKE) during the early stages of crop.
- Foliar application of chlorpyrifos 50EC + cypermethrin 5EC (1ml/litre) or quinalphos 25EC (2ml/l).
- Release of egg parasitoids *Trichogramma chilonis* and *Telenomus remus* four times at 50,000/ha at weekly interval against *S. litura*.
- Use pheromone traps (5 traps/ha) for monitoring of the pest.

Black cutworm, *Agrotis ipsilon* (Hufn.) (Lepidoptera: Noctuidae)

Distribution : *Agrotis ipsilon* is well distributed in Asia, North Africa, Europe, America and Oceania.

Host plants: A part from *Mentha* spp. the pest are also known to feed on the other crops such as potato, barley, oats, pulses, peas, beetroot, cabbage, tobacco and many other cultivated crops (Khan, 1976).

Identification and Biology : The moth is stout and has wavy lines and spots on brownish forewings whereas; hindwings of adults are off-white with dark veins (Fig.4). The larva is dark brown in colour (50 mm long) with a distinct yellow dot on each segment down the centre of back. The female lays about 300 eggs in groups of 30 on the under surface of leaves or moist soil. The egg, larval and pupal periods is respectively 2-13, 10-30 and 10-30 days. Larva moults six times to become pupa. The total life cycle lasts from 30 to 68 depending on weather conditions. The pest completes four



Figure 4: Black cutworm, *Agrotis ipsilon*

generations in a year.

Nature of Damage: Larvae remain hidden during day time in cracks in the soil, become active at dusk, feed on leaves and also cut the tender, stems of young and growing plants and thus cause retardation in growth and reduction in yield.

Management

- Deep ploughing exposes the larval and pupal stages to predators.
- Collect and destroy larvae mechanically as they may be curled near the base of young plants.
- Install bird perches to increase the activity of birds in the field.
- Use poison baits (2 g malathion 5% dust with 1 kg of wheat bran) or *Bt* mixed bait (2 g of *Bacillus thuringiensis* formulation with 1 kg of wheat bran) @ 10kg bait/ha in field one week before planting.
- Keep piles of weeds in the crop field as the larvae hide inside weeds to remain protected from day light. Collect and destroy the hidden larvae during morning hours.
- Soil application of granular phorate 10G (1.5 kg a.i. ha) before planting reduces cutworm damage (Szezeponek and Mazur, 2006; Rita and Animesh, 2011) or Before planting broadcast chlorpyrifos 50EC treated sand (3 litre per 10 kg sand) @ 10 kg treated sand/ha in the field.
- Release egg parasitoids, *Trichogramma* sp. at weekly interval to parasitize cutworm eggs.
- Release entomopathogenic nematodes in moist soil which kills the cutworms living underground (Yuksel *et al.* 2018).

Cutworm, *Spodoptera exigua* (Hbn.)



Figure 5: Cutworm, *Spodoptera exigua*

(Lepidoptera: Noctuidae)

Distribution

Spodoptera exigua is native to Southeast Asia and currently found in Africa, Southern Europe, North America, Japan and Australia.

Host plants: *S. exigua* has a wide host range, occurring as a serious pest of asparagus, cabbage, pepper, tomato, lettuce, celery, strawberry, eggplant, sugar beet, alfalfa, *Mentha* spp. and cotton

Identification and Biology: The adult moths' are medium sized, with wing span measuring 25-30 mm. The forewings are grayish brown with irregular banding pattern and a light yellow coloured spot (Fig.5). Hind wings are grayish white colour. The early instars larvae are pale green in colour. The later instar larvae are brownish green with yellowish white stripe laterally and yellowish pink colour ventrally. The larvae hide in soil during day time and feed on leaves at night and become full grown in 10-16 days. The full grown larvae pupate in the soil in earthen cocoons and emerge as adults in 7 to 11 days. The female moth lays cluster of eggs on the lower portions of young plants.

Nature of Damage: Larvae feed on the foliage of the mint plants. Young caterpillars feed gregariously and skeletonize the foliage. As they mature, caterpillars become solitary and eat large irregular holes in foliage.



Figure 6. *Syngamia abruptalis*

Management: Described under *Agrotis ipsilon*.

Mentha leaf roller, *Syngamia abruptalis* Walker (Lepidoptera: Pyralidae)

Distribution

It occurs in the tropics of the Old World from Africa to Australia.

Host plants: *S. abruptalis* has a host range restricted to *Mentha* spp. and *Ocimum sanctum* (Sagar and Reddy 1985).

Identification and Biology: The adult moths are light yellowish brown in colour with small size. The female moth lays 12-74 eggs on the lower surface of leaves singly. The incubation period varies 3 to 5 days. Larval duration ranges between 18 to 25 days with five larval instars. Pupal period lasts 5-7 days and 6-10 adult longevity.

Nature of Damage : Early instar larvae feeds on the undersurface of leaves and removes the parenchyma making very small patches of injury without forming webbing of leaves (Fig.6). The later growth stages of larvae forms webbing by joining the edges of upper and lower leaves with silken threads. Due to webbing the growth of plant is hindered (Kareem *et al.* 1960; Sandhu and Bhala 1975; Sagar and Reddy 1987).

Management

- Cypermethrin (50 gm a.i./ha) and Quinalphos



Figure 7. Cotton whitefly, *Bemisia tabaci*

(0.05%) found effective in controlling the *Syngamia abruptalis* (Sagar, 1983,1984).

Cotton whitefly, *Bemisia tabaci* Gennadius (Hemiptera: Aleyrodidae)

Distribution: The whitefly, *Bemisia tabaci* (Gennadius) (Hemiptera: Aleyrodidae) is a devastating pest of different crops throughout the tropical and subtropical regions of the world (Oliveira *et al.* 2001).

Host plants: *B. tabaci* has been recorded from more than 900 different plant species (Attique *et al.* 2003; Greathead 1986, GISD, 2005).

Identification and Biology: Adults are winged, they are 1.0-1.5 mm long and their yellowish bodies are slightly dusted with white waxy powder (Fig.7). They have two pairs of pure white wings and have prominent long hind wings. The wings are held over the body like a tent. The adult males are slightly smaller in size than the females. The nymphs are flattened, oval-shaped, and greenish-yellow in color. The first instar nymphs are mobile and have antennae, eyes, and three pairs of well developed legs. The legs and antennae are atrophied during the next three instars and they are immobile during the remaining nymphal stages. The last nymphal stage has red eyes. This stage is sometimes referred to puparium. The females lay stalked eggs singly on the underside of the leaves, with a range between

80-110 eggs per female. The incubation period is 3-5 days. The nymphs feed on cell sap and grow into three stages to form the pupae within 14-28 days. In 2-8 days, the pupae change into whiteflies. The total lifecycle completed in 14-100 days depending on weather conditions.

Nature of Damage: Nymphs and adult whiteflies suck the cell sap from the lower side leaves, thus lowering the vitality of the plants. The whiteflies excrete honeydew on which growth of sooty mould takes place, which results in blackening of leaves, consequently affecting photosynthesis. In case of very high incidence, there is total blackening of crop resulting in drying of leaves and ultimately total failure of the crop. Whiteflies are also a vector of number of viral diseases.

Management

- Grow resistant cultivars of *Mentha arvensis* (MA-92-194 and Kosi), *M. piperita* (MPS-1), *M. cardiaca* (MCAS-2 and MCAM-19), *M. citrata* (Kiran and Arka) and *M. spicata* (Neerkalka and MSS-5) which suffer less infestation of whiteflies (Singh *et al.*, 2004).
- Removal the weed plants in and around the mint cultivated fields may in decreasing the pest population.
- Yellow sticky traps are effective for monitoring and managing the whiteflies.
- Conserve the naturally occurring natural enemies, namely, predators: *Serangium parcesetosum*, *Cheilomenes sexmaculata*, *Coccinella septempunctata*, *Brumoides suturalis*, chrysopid *Chrysoperla zastrowi sillemi* and parasitoid, *Encarsia lutea* play important role in controlling *B. tabaci* (Kedar *et al.* 2014; Kedar *et al.* 2017).
- Grow sorghum or maize as barrier crop in may be planted to reduce the pest population (Hilje *et al.* 2001).
- During initial stages foliar application based on neem oil, castor oil, cotton seed oil etc. must be followed.
- Foliar application of entomopathogenic fungus, *Verticillium lecanii* (10gms/lit) against *B. tabaci*.

Aphids, *Aphis affinis* Del Guercio (Hemiptera: Aphididae)

Distribution

Aphid, *Aphis affinis* is found in Southern Europe, Russia, the Middle East, India and Pakistan.

Host plants: This pest is monophagous feeds on plant belonging to **Lamiaceae family** (*Mentha viridis* and *Ocimum basilicum*).

Identification and Biology: They are small to medium sized soft-bodied insects, dark grey-green to almost black in colour. These aphids are dusted with light wax powder with clear intersegmental lines. The adult female multiplies parthenogenetically and viviparously.

Nature of Damage: The nymphs and adults are usually clustered on stem apices and tender shoots and upper leaves of *Mentha* spp. Nymphs and adults suck the plant sap due to which the growth of the plants is affected. Such stressed plants are more susceptible to other pests. Aphids excrete honey dew on which growth of sooty mould fungus takes place, which results in blackening of leaves consequently affecting the photosynthesis process. Aphids are also vector of viral diseases which can be more damaging than the feeding aphids (Singh and Sagar 1981).

Management

- Use of yellow sticky traps for monitoring and mass trapping of aphids.
- Conservation of natural enemies by minimizing use of broad spectrum insecticides.
- *Coccinella septempunctata*, *Coccinella transversalis*, *Cheilomenes sexmaculata*, *Brumoides suturalis* and *Propylea dissecta* reported biocontrol agents of *A. affinis* infesting Japanese mint (Singh and Bali, 1993).
- Foliar application of oxydemeton-methyl 25EC (2ml/l) or dimethoate 30EC (1.5ml/l) can be used for control *A. affinis* (Sagar and Singh, 1981).

Red Pumpkin Beetle, *Aulacophora foveicollis* (Lucas), (Coleoptera: Chrysomelidae)

Distribution: Red Pumpkin Beetle is widely distributed in Asia, Africa, Australia and South Europe. In India, it occurs throughout the country.

Host plants: *Aulacophora foveicollis* is a major pest of cucurbitaceous crops. Damage of this pest was also reported on *Mentha arvensis* (Singh and Gupta 1970).

Identification and Biology; The adult beetle is oblong in shape deep orange and ventral side is black in colour (Fig.8). The adult beetle measures about 5-8 mm in length and 3.5-3.75 mm in width. Adult female beetle lays brownish elongate eggs singly or in batches of 8 to 9 in the moist soil near the base of host plant. A beetle may lay 150 to 300 eggs. The incubation period is 5 to 8 days. The grubs become full grown in 13-25 days and pupate in soil. The pupal period ranges from 7 to 17 days. In a year there may be 5 to 8 generations of the insect. The adult beetles hibernate in soil or in dry weeds during November to February.

Nature of Damage: The adult beetle feeds on the leaves. The beetle bite holes on the leaves, causing death or retardation of growth. Damage during seedling stage is often devastating. The grubs of this pest feed on the



Figure 8: Red Pumpkin Beetle, *Aulacophora foveicollis*



Figure 9:Termite, *Odontotermes obesus*

roots and sometimes on stem also (Singh and Gupta 1970).

Management

- Hand collection and destruction of adult beetles.
- Foliar application of 0.1% malathion 50EC (Singh and Gupta, 1970) or trichlorfon* 50EC (500ml/ha) or trichlorfon* 5% dust (500gm/ha) or diclorvos* 76EC (1.25ml/l) is effective against this pest. (*The use these insecticides is completely banned with effect from 31st December 2020).

White ants/Termites, *Odontotermes obesus* (Ramb.) (Isoptera: Termitidae)

Distribution: *Odontotermes obesus* is a widespread species in India, Bangladesh and Pakistan.

Identification: Termites are social insects; live in termitaria, in distinct castes, workers, soldiers, kings and queen. Worker caste causes damage to plants. Workers are small and have soft, white body and brown head.

Nature of Damage: Termites damage plants soon after transplanting and continue till late growing stage. Termites bore in to root and stem. The leaves of the damaged plant droop down (Fig.9). Such plants, if pulled, are easily uprooted and termite cuts or galleries are seen.

The incidence of termite is common during drought period.

Management

- Field application of oil cakes of *Azadirachta indica*, *Mahua longifolia* and *Ricinus communis* is effective for termite management.
- Soil application of chlorpyriphos 20EC is very effective.
- Avoid use of partially decomposed farm yard manure (FYM).
- Irrigating field at short period of intervals also reduces termite infestation.
- Destroy the termitaria around field and kill the termite queen.
- Spot application of granular insecticides (phorate 10G) if termite infestation is in patches.
- Entomopathogenic nematodes effective against termites (Rathour *et al.* 2014).

Fungal diseases of mints (Table 2)

Rust: Rust is an important disease which is caused by *Puccinia menthae* Pers. (Shukla *et al.* 1999). This fungus infest leaves and twigs of *Mentha* species and produces postules on upper and lower leaf surface, are in redish brown and yellow orange in colour

Host range: *Mentha arvensis* and *M. spicata* are the major host.



Figure 10. Rust of *Mentha arvensis* caused by *Puccinia menthae*

Diseases occurrence and distribution: Due to infection of rust fungi the defoliation in mints occurred and caused severe economic loss due to reduction in oil yield. This disease is prevalent in all mint growing area of India (Ganguli and Pandotra, 1962; Shukla et al. 1999) whereas the fungus *Puccinia menthae* has also been reported from Indiana, USA (Stone and Green, 1967). In tarai region of Uttar Pradesh the dual infection of *P. menthae* and *Alternaria alternata* have been reported (Shukla et al. 1999). The young shoots infected by rust fungi are usually twisted and most of the times broken off at the infection site (Fig. 10).

Diagnostic symptoms: In menthol mint typical symptoms of rust pustules develop on lower surface of leaves between April and June. In general, there are three types of pustules found

on the infected area. First types of pustules are circular, yellowish brown in colour whereas second is orange in colour and third type of pustules are of variable shape and size, dotted tan in colour.

Disease cycle: These fungi generally persist in the form of ureidiospores on runner and plant debris of the host plant. The rust pustules possess enough ureidiospores which under favourable condition infect the host plant. The rust fungi also formed telia at later stage which is thick walled teliospores and survives in adverse conditions.

Management

Use of disease free materials is the major preventive measures of this disease. Treatment of planting materials with hot water and use of resistant variety is common method of this disease management. Elimination of old mint rust infected mint plant also help to prevent the disease spreading and also reduce the rust inoculum. Application of few fungicides like mancozeb, propiconazole, Diclobutazol and nickel chloride offer reliable protection against rust disease (Krall 1977, Margina and Zheljzakov 1994). Suab and Nagy (1972) also reported that application of denitroamine and Krezonit-E (DNOC) help to decrease the disease incidence. Shukla et al. (2008) screened different varieties/genotypes for multiple disease resistance of menthol mint, himalaya, kalka and Damroo were found to be

Table 2. Major fungal diseases of *Mentha* spp.

Sr. No.	Disease	Causal Organism	References
1	Alternaria leaf spot	<i>Alternaria tenuis</i> <i>A.alternata</i>	Ganguli and Pandotra, 1962 Srivastava and Srivastava, 1971
2	Cercospora leaf spot	<i>Cercospora menthicola</i>	Shukla et al. 1988
3	Corynespora leaf spot	<i>Corynespora cassicola</i>	Sattar et al. 1981
4	Curvularia leaf spot	<i>Curvularia verruciformis</i> , <i>C.trifoli</i>	Thakur et al. 1974
5	Alternaria blight	<i>Alternaria alternata</i> .	Shukla et al. 2000
6	Rhizoctonia blight	<i>Rhizoctonia solani</i>	Sharma and Munjal, 1978
7	Anthraxnose disease	<i>Sphaceloma menthae</i>	Bains, 1938
8	Root rot	<i>Thielavia basicola</i> <i>Rhizoctonia bataticola</i> <i>Sclerotinia roffsii</i> <i>Rhizoctonia solani</i> <i>Sclerotinia sclerotiorum</i>	Sattar and Husain, 1976 Husain and Janardhnan, 1965 Pandotra and Shastry, 1968 Singh, 1991
9	Stem rot	<i>Botryodiplodia theobromae</i>	Shukla et al. 2001
10	Wilt	<i>Fusarium oxysporum</i>	Sattar and Husain, 1980

resistant to rust fungi whereas genotypes MAH-3, S-10-11-45 and S-13-2-125 were moderately resistant to rust, The variety Kalka was resistant to rust, leaf spot and leaf blight, but was found susceptible to powdery mildew, and stem blackening and rot.

Viral diseases of mints: Viruses are obligate parasites; that is, they require a living host in order to grow and multiply. Therefore they are considered intracellular (inside cells) pathogenic particles that infect other living organisms. In general the virus particles are very small and can be seen only with the help of an electron microscope. Most of the plant viruses are either rod or isometric (polyhedral) in shape. Generally TMV, potato virus Y (PVY) as well as cucumber mosaic virus (CMV) are very good examples of a short rigid rod-shaped, a long flexuous rod-shaped, and an isometric virus, respectively. For viral infection there are need of injury for their entrance in plant cell and injury may occur naturally in plants like lateral branching. The transmission of plant viruses may be through agronomic or horticultural practices, or fungal, nematode, insects etc.

Several viruses belonging to more than twelve genera have been reported to cause diseases in various species of *Mentha* spp. World-wide, the important genera include: *Alfamovirus*, *Begomovirus*, *Cheravirus*, *Closterovirus*, *Cucumovirus*, *Cytorhabdovirus*, *Hordeivirus*,



Figure 11: Mosaic on leaves of *M. gracillis*

Nepovirus, *Potexvirus*, *Tobamovirus*, *Tospovirus*, *Vitivirus*, etc. However, there are only few reports of viral diseases on mints from India which may be attributed that this crop has come in regular and mass cultivation here, lately.

A mosaic disease of *M. arvensis* was suspected to be caused by virus, was reported as early as in 1965 in India, but the causal virus could not be characterized (Nene *et al.* 1965). The genus *Begomovirus* belongs to the family *Geminiviridae*. A light yellow mosaic disease has been reported from Lucknow on *Mentha spicata* cv. *viridis* (Samad *et al.* 2009). The symptoms include mosaic, yellowing, leaf curling, crinkling, and retarded growth that induced drastic reduction in herb yield. An isolate of *Tomato leaf curl Pakistan virus* (*Begomovirus*) was identified from infected plants.

Tobacco mosaic virus belongs to genus

Table 3. Major viral diseases of mints in India

Sr. No	Name of Virus/Diseases	Genus	Host plant	Transmission	Place	References
1.	Mosaic Disease of <i>M. arvensis</i>	--	<i>M. arvensis</i>	--	India	Nene <i>et al.</i> 1965
2.	Tobacco mosaic virus	<i>Tobamovirus</i>	<i>M. cardiaca</i>	Mechanical	India	Samad <i>et al.</i> 1994
3.	Tobacco mosaic virus	<i>Tobamovirus</i>	<i>M. gracillis</i>	Mechanical	India	Samad <i>et al.</i> 2000
4.	Tomato leaf curl Pakistan virus	<i>Begomovirus</i>	<i>M. spicata</i>	whitefly	India	Samad <i>et al.</i> 2009
5.	Mosaic and leaf deformation disease virus	--	<i>M. piperita</i>	--	India	Zaim (2011, unpublished)

Tobamovirus. The virus was first reported in Scotch spearmint (*M. cardiaca* Baker) in 1994 in India (Samad et al. 1994). Subsequently an isolate of TMV was isolated from Scotch spearmint (*M. gracilis* Sole) (Samad et al. 2000). The disease symptoms included green mosaic with vein banding, deformation of leaves, and retarded growth (Fig. 11). The virus particles were rod shaped 300x18 nm in size. Mol. Mass of capsid protein was found to be 17000 ± 500 Daltons. The virus showed close serological relationship with TMV-UI and Brinjal necrotic mosaic virus

Another mild mosaic symptom has been recorded on peppermint (*M. piperita* L.). Initially terminal leaves of the diseased plant showed mottling, curling and finally mild mosaic symptom which spread in whole plant giving sick appearance, associated with heavy stunting and occasional necrosis of the leaves. Flowering was delayed with small size flower. The DNA from the diseased leaves amplified and identified as the causal virus could be a member of family Geminiviridae (Zaim 2011 unpublished).

Nematode diseases of mints

Menthol mint (*Mentha arvensis* L.): Menthol mint (*Mentha arvensis* L.), an important aromatic herb, is widely cultivated for its essential oil, which is a potential source of natural menthol, menthyl acetate, menthone and terpenes. Different constituents of menthol mint oil are extensively used in pharmaceutical, perfumery, food and cosmetic industries all over the world. In India the commercial cultivation of menthol mint suffers a great yield loss due to root-knot disease caused by *Meloidogyne* spp. Both the species of root knot nematode (*Meloidogyne incognita* & *M. javanica*) occur together but the frequency of *M. incognita* is greater than *M. javanica* (Pandey et al. 1992). In the root rhizosphere other plant parasitic nematodes viz. *Tylenchus* sp., *Tylenchorhynchus vulgaris*, *Hoplolaimus indicus*, *Helicotylenchus dihystra*, *Pratylenchus thornei*,

Rotylenchulus reniformis, *Aphelenchoides* sp., *Longidorus pisi*, and *Xiphinema* sp. were reported (Pandey et al. 1992, Pandey 1997a). In 1938, Buhner for the first time reported that *M. arvensis* as host of *Meloidogyne* species. For the first time Pandey (1989) reported the root-knot disease of menthol mint (*Mentha arvensis*) and Bergamot mint (*M. citrata*). Two major nematode species viz. *M. incognita* and *M. javanica* were found to be associated with menthol mint. Introduction of *M. arvensis* as a commercial crop into sandy loam soil in India showed that a threat was posed by the root-knot nematode, *Meloidogyne incognita* (Kofoid et White) Chitw. (Haseeb and Pandey 1989). As this species is sedentary endoparasitic, its spread is facilitated by infected suckers/roots that are used in the vegetative propagation of the crop (Pandey 1998a, b).

Distribution: Root-knot disease was first observed in Lucknow and subsequently found in Pantnagar, Rampur, Moradabad, Badaun, Bareilly, Saharanpur. The incidence of disease varied from cultivar to cultivar and location to location. The most severe infestation was found in Lucknow, Barabanki, and Moradabad whereas least infestation was recorded at Pantnagar (Pandey, 1989).

Symptoms: Whole roots and suckers are invaded by root-knot nematode. Small to medium size of galls were noticed on *M. incognita* infested roots. Secondary roots of suckers are also highly infested with *M. incognita* and most of the times whitish colour egg masses of *M. incognita* were seen on the root system. Heavily root knot infested menthol mints plants are stunted poorly tillered and have chlorotic leaves. Infested suckers serve as major source of infection and means of dissemination. Root knot nematode continue to develop after the crop has matured and been harvested.

Other host: Several weeds, which are present

in menthol mint growing areas, are also major source of root-knot nematodes

Host-parasite relationship: Experiments have been carried out to determine the pathogenic potentiality of different plant parasitic nematodes on menthol mint (*Mentha arvensis* L.). The pathogenic potentiality of *M. incognita* was observed on all cultivars of *Mentha arvensis*. With an increase in load of inoculum there had been a significant decrease in oil yield, fresh and dry weight of plant and rate of photosynthesis. The reduction in different growth parameters was found to be directly correlated with inoculum load of nematode. Root-knot nematode (*M. incognita*) caused 25-30% oil yield reduction in menthol mint. The quality of mint oil was also adversely affected due to nematode infection (Pandey *et al.* 1992).

MANAGEMENT

To manage plant parasitic nematodes in different medicinal and aromatic plants through ecofriendly way is one of the difficult tasks because of different nature of parasitism. Studies conducted by Pandey *et al.* (1997, 1998) on interaction between *M. incognita* and AM fungi indicated that root-knot nematode, *M. incognita* multiplies well in absence of AM fungi and significantly reduced plant growth/ yield. Root-knot nematode infection was drastically impaired when plants were inoculated with three AM fungi simultaneously as compared to their alone inoculation. It was concluded with this experiment that AM fungi besides improving plant growth biomass could effectively inhibit nematode infection (Pandey *et al.* 1998).

Successful control of root-knot disease has been achieved with carbofuran (@ 3kg a.i. /ha (Pandey 1995). Addition of neem cake (@500kg/ha) was also found encouraging. Significant success both in pots as well as in fields with regard to management of *M. incognita* with bio-agents, organic matter and

integration of both has also been obtained (Pandey, 2000). Pandey and Patra (2001) screened large number of mint accessions against root-knot nematode, *M. incognita* and reported moderate to high degree of resistance in 11 accessions of mint species. Development of some varieties, which could be propagated through seeds, would also help in breaking the disease cycle by avoidance of infested suckers. Management of the nematode by integration of various means would probably be the best method in fighting nematode problems. Current research is directed towards this goal. Pandey (2005) conducted field trial to determine the efficacy of *Trichoderma harzianum* isolate U, *Glomus aggregatum*, oil seed cakes of neem (*Azadirachta indica*), and mustard (*Brassica campestris*) in the management of *Meloidogyne incognita* and their impact on yield of menthol mint (*Mentha arvensis*) cv. Kosi. Significant reductions in nematode populations and root-knot indices were noticed in plots receiving oil seed cakes and bio agents, whose effects were equal to that of carbofuran. Application of oil seed cakes and *T. harzianum* significantly enhanced crop yield. The late transplanted mint technology developed at Central Institute of Medicinal and Aromatic Plants Lucknow, which allows farmers to have non-host crop like wheat, mustard etc. has greatly benefited the farmer in fighting root-knot nematode menace to some extent. Further the higher temperature prevailing during the transplanted cropping season (April-July) also checks the nematode population buildup and infection of menthol mint crop.

STRATEGIES TO PREVENT SPREAD OF MINT NEMATODES

Most plant-parasitic nematodes, which affect the crop yield, occur in the soil or plant roots. Prevention of spreading of phytonematodes from nematode infested field to noninfested field is an important part as the nematodes

which damage mints are generally endoparasitic in nature and they can easily spread through suckers / runners (propagating materials) from infested planting sites to noninfested planting sites. Therefore it is essential and utmost important to check their spread in new areas/ fields. We should take following prevention measures:

- Use of certified suckers as planting material
- Use of nematode free soil for suckers multiplication
- Cleaning soil from equipment before moving between fields
- Prevention of animal movement from infested to uninfected fields
- Eliminating important weed and alternate hosts of phytonematodes

Peppermint (*Mentha piperita* L.)

:Peppermint (*Mentha piperita* L.) Belonging to family Lamiaceae is also known as black mint, candy mint or “Mitcham” yielded peppermint oil by the steam distillation of herbs. The major oil component of peppermint oil is menthol, menthone, isomenthone, menthyl acetate and neomenthol. Oil of these mint species is used in pharmaceutical, flavour and food industries. The oil possesses antibacterial, antiviral, antiparasitic, antifungal as well as nematicidal properties. Peppermint is commercially cultivated in USA, Russia, Bulgaria, Italy, Morocco and India. In India it is cultivated on 4300 ha of land and oil production is about 830 tonnes annually. The world requirement is approximate 8000 tonnes per year and USA is the largest producer and user of peppermint oil. In India the major cultivated states are, Uttar Pradesh, Himachal Pradesh, and Uttarakhand. Buhner (1938) reported *M. piperita* as host of *Meloidogyne* species. Horner and Jenson (1954) reported in western Oregon, USA the occurrence of *M. hapla* on different mint species. Species of root lesion nematode, *Pratylenchus* like *P. minyus*, *P. penetrans* and *P. thornei* associated with, *M. piperita* var. Mitcham (Skotland and Menzies, 1957;

Pandey, 1997b). The name “root lesion” nematode has been derived from the characteristic symptoms produced due to this nematode infection in roots. De Man described the first species of *Pratylenchus* in 1880 under the name of *Tylenchus pratensis*. Filipjev established the genus *Pratylenchus* in 1936. and more than 50 species of *Pratylenchus* have been reported all over the world.

Major causal organism: *Pratylenchus thornei*, *P. penetrans*, *P. scribeneri*, *P. minyus*

Symptoms: This mint species is highly susceptible to *Pratylenchus thornei*, *P. penetrans*, *P. scribeneri*, *P. minyus* (Skotland and Menzies, 1957; Faulkner, 1962; 1964 Pandey, 1997c). *Pratylenchus* species is migratory endoparasitic nematode and attack the root cortex. Roots develop dark lesions and an overall brown color. The lesions first develop on younger roots but may be found anywhere on the roots. The lesions enlarge, coalesce and ultimately girdle whole root which appear constricted this follows secondary attack by fungi and bacteria. The affected area may be sloughed off and the overall root system is reduced. Loss of the epidermis and cortex decreased root growth nutrient and water uptake. Under stress, plants are yellow, become stunted, and have reduced yields. The extent of damage depends on the phenolic content of the cells. The oxidation of these phenols causes the characteristic browning of the lesions. The aerial symptoms always resembled like due to malnutrition or water or deficiency. In the initial stages, the infestation can be noticed as small patches of stunted growth of mint and yellowing of leaves. These stunting increases every year and may formed a large patch. Leaves may become yellowish brown and wilt during April-May. The root system of mint species become very poor when severely infested with *P. thornei* and can be pulled out from the field very easily (Pandey, 1997b).

Host parasite relationship:The root lesion

nematode is second most important pest of medicinal and aromatic crops. *Pratylenchus* gain importance due to its ubiquitous distribution, although certain species have climatic preferences, wide host range render the host more vulnerable to secondary attack by other pathogens due to extensive necrosis. The nematode migration is both inter and intra cellular and feeding is restricted to cortical region. Bergeson and Green (1979) reported that all cultivars of peppermint grown in Indiana are highly susceptible to *P. penetrans*. The damage becomes many folds when *Pratylenchus* species interact with plant pathogenic fungi in peppermint and spearmint. Interactive effect of *P. penetrans* and *Verticillium albo atrum* on yield of peppermint was established by Bergeson (1963). Similar result was reported by Faulkner (1965) with *P. minyus* and *V. dahliae* (kleb) f. *menthae*. Presence of nematode species increases both incidence and severity of disease. Faulkner and Bolander (1969) have studied the disease severity of peppermint at different temperature when inoculated peppermint plant with *V. dahliae* f.sp. *menthae* and *Pratylenchus minyus*. The 27°C temperature favours the disease development whereas optimum wilting occurs at 24°C temperature, higher temperature did not favour the nematode multiplication as least nematode reproduction was reported at 30°C. Faulkner *et al.* (1970) reported that *P. minyus* not only help to the *V. dahliae* for infection but also alter the physiology and biochemistry of *M. piperita* plant. Esmenjaud *et al.* (1990) found that three-peppermint cultivar proved to be good host of *Pratylenchoides lauticauda*. Studies carried out in CIMAP reveals that the population of *P. thornei* was very high in roots and the root rhizosphere of rhizospheric stunted peppermint plants. Pathogenicity of *P. thornei* was established on this mint species (Pandey 1997b). *Pratylenchus* spp. frequently caused wounds in roots through which other pathogenic organism such as fungi and bacteria enter the root tissues. The interaction of these agents results in the formation of lesions that

finally destroy the whole root tissues.

Management

Information pertaining to the management studies of root-lesion nematode on *M. piperita* is very meager. Pinkerton *et al.* (1988) successfully managed the population of *P. penetrans* on peppermint through the use of oxamyl, aldicarb and carbofuran. Marrifield and Ingham (1992) used ethoprop@6.6kg a.i./ha for the control of *P. penetrans* in *Mentha piperita*, whereas Ingham (1992) applied same nematicide @6.6, 13.2 or 26.4 kg a.i./ha for the management of lesion nematode in peppermint. They concluded their findings and reported that ethoprop @6.6kg a.i./ha is very effective to reduce the root lesion nematode population in soil at significant level. On control aspect several experiment were carried out with herbicides, pesticides, and cultural methods with variety of plant parasitic nematodes on peppermint (Faulkner and Skotland 1963,, Jatala and Jensen 1974, Gokte and Mathur 1980, Pinkerton and Jensen 1983, Jensen and Horner 1959, Pandey 1993). The non-chemical methods are getting more importance nowadays due to hazardous effect of chemical pesticides. In general the management strategies can be concluded as follows

- I. The use of resistant crops and varieties provides the best option for farmers to control *Pratylenchus* as these crops will limit multiplication of the nematode and reduce losses to following crops.
- II. Some crops and varieties vary in their resistance/susceptibility and tolerance to either *P. penetrans* or *P. thornei*.
- III. To limit yield losses from *Pratylenchus*, soil assays to identify the species and nematode levels present are highly recommended. With this information, the likely risks can be assessed, and management decisions made, especially for variety resistance and tolerance of crops to be used in rotations.
- IV. Good soil nutrition (particularly nitrogen, phosphorus and zinc) can reduce the effects of

Pratylenchus on infested peppermint plant.

Spearmint (*Mentha spicata* Huds.)

Distillation of spearmint herbs yielded spearmint oil, which is major source of carvone and limonene. Oil is mainly used in flavouring industries. Species of root lesion nematode, *Pratylenchus* like *P. minyus*, *P. penetrans* and *P. thornei* associated with *M. spicata* var. native (Skotland and Menzies, 1957).

Causal Organism: *Pratylenchus thornei*, *P. penetrans*, *P. scribeneri*,

Host parasite relationship: Bergeson (1963) for the first time established the pathogenicity of *P. penetrans* on *M. spicata* and reported a reduction of foliage and root/ stolon growth up to 34 and 66% respectively. The population of *P. penetrans* increases from 30X to 80X after 8 months. Bergeson and Green (1979) showed the herb reduction of spearmint in Indiana due to *P. penetrans*. Further they reported that all cultivars of spearmint grown in Indiana are highly susceptible to *P. penetrans*. Rhoades (1983) isolated the population of *Belonolaimus longicaudatus*, *Dolichodorus heterocephalus*, *Pratylenchus scribeneri* and *Paratrichodorus christie* from the stunted plant rhizosphere of *M. spicata* and pathogenic potentiality of these nematode has also been tested. Similarly Inserra and Rhoades (1989) reported damage potential of *Belonolaimus longicaudatus*, *Dolichodorus heterocephalus*, *Pratylenchus scribeneri* and *Paratrichodorus christie* on *M. spicata* and suggested the use of non-volatile nematicide for lowering the population of these nematodes and increasing yield of the plant. Studies carried out in CIMAP reveals that the population of *P. thornei* was high in the root and rhizosphere of chlorotic spearmint plant (Pandey 1997b). Pathogenicity of *P. thornei* was established on spearmint (Pandey 1997b). The damage becomes several folds when *Pratylenchus* species interact with plant pathogenic fungi in spearmint crops.

Management: Least experiment has been carried out on the management aspect of lesion nematode in spearmint. Rhoades (1984) studied the effect of carbofuran, fenamiphos and oxamyl @5.6 and 11.2 kg/ha and terbufos@11.2kg/ha for the control of *P. scribeneri* infesting *M. spicata*. The nematode population decreased significantly but terbufos best followed the affectivity of fenamiphos. Similarly Ingham *et al.* (1992) used ethoprop@6.6, 13.2,26.4 kg a.i./ha in their experimentation for the control of *P. penetrans* in *M. spicata*. They reported ethoprop @6.6kg a.i/ha was highly effective to reduce the population of *P. penetrans* in spearmint field. Some of the experiments on control aspect were also carried out with herbicides, pesticides, and cultural methods with variety of plant parasitic nematodes on spearmint (Jatala and Jenson 1974; Lisetskaya 1985, Emenjaud *et al.* 1989, Gokte and Mathur 1980, Pandey 1993).

Nematodes associated with Scotch spearmint, Bergamot mint and Gardenmint Scotsch spearmint (*Mentha cardiaca* (GF Gray) baker, Bergamot mint (*Mentha citrata* Ehrh.) and Gardenmint (*M. viridis* L.) are affected by various plant parasitic nematodes. Large number of plant parasitic nematodes is associated with different mint species other than root knot and root lesion nematodes. Horner and Jenson (1954) reported the association of *M. hapla*, *Pratylenchus* species, *Aphelenchoides* sp. and *Longidorus* sp. with these three mint species and established pathogenicity of *M. hapla* on *M. cardiaca*. *Aphelenchoides olesistus* and *A. ritzemabosi* associated with *M. piperita* and *M. spicata* (Goodey, 1940). Skotland and Menzies (1957) reported the association of *Tylenchorhynchus capitatus*, *Trichodorus* sp. with *M. cardiaca* var. native in the Yakima valley and in Columbia basin. They further reported the population of *P. hamatus* was associated with *M. cardiaca* var. Scotch, *M. piperita* var. mitcham and *M. spicata* var. native. Population

of *Longidorus elongatus* (de man) Thorne & Swanger associated with peppermint cultivation in USA (Horner and Jenson, 1954). The population of *L. elongatus* was controlled with the use of oxamyl on peppermint (Jatala and Jenson, 1974).

In India the most important nematode pathogen of Scotch spearmint and Garden mint is *M. incognita* while bergamot mint is affected by *P. thornei* (Pandey, 1997b). Among these three mint species *M. cardiaca* and *M. viridis* are

most susceptible while *M. citrata* is considered to be tolerant to *M. incognita* and *M. javanica* (Pandey, 1998a). Most plant-parasitic nematodes, which affect the crop yield, occur in the soil or plant roots. Prevention of spreading of plant parasitic nematodes from nematode infested field to non-infested field is an important part as the nematodes which damage mints are generally endoparasitic in nature and they can easily spread through suckers (propagating materials) from infested

Table 4. Insect pests of Patchouli

Common name	Scientific name	Order	Family	Pest Status	References
Weevil	<i>Blosyrus</i> sp.	Coleoptera	Curculionidae	Major	Triveni, 2017
Ash weevil	<i>Mylocerus viridanus</i>	Coleoptera	Curculionidae	Major	Triveni, 2017
Weevil	<i>Apion</i> sp.	Coleoptera	Curculionidae	Minor	Triveni, 2017
Scarab beetle	<i>Maladera</i> sp.	Coleoptera	Scarabaeidae	Minor	Triveni, 2017
Beetle	<i>Longitarsus</i> sp.	Coleoptera	Chrysomelidae	Minor	Triveni, 2017
Bug	<i>Aspilocoryphus dixnoui</i>	Hemiptera	Lygaeidae	Minor	Triveni, 2017
Leaf hopper	<i>Neodartus acocephaloides</i>	Hemiptera	Cicadellidae	Minor	Triveni, 2017
	<i>Kolla ceylonica</i>	Hemiptera	Cicadellidae	Minor	Triveni, 2017
	<i>Nirvana pallida</i>	Hemiptera	Cicadellidae	Minor	Triveni, 2017
Aphid	<i>Aphis gossypii</i>	Hemiptera	Aphididae	Major	Triveni, 2017
Bug	<i>Pachypeltis</i> sp.	Hemiptera	Miridae	Major	Ramani and Bhumannavar, 1990
Yellow scale of cotton	<i>Cerococcus hibisci</i>	Hemiptera	Cerococcidae	Minor	Umesha <i>et al.</i> , 1998
Thrips	<i>Bathrips melanicornis</i>	Thysanoptera	Thripidae	Minor	Sanjta and Chauhan, 2018
	<i>Thrips carthami</i>	Thysanoptera	Thripidae	Minor	Sanjta and Chauhan, 2018
	<i>Anaphothrips sudanensis</i>	Thysanoptera	Thripidae	Minor	Sanjta and Chauhan, 2018
	<i>Aeolothrips</i> sp.	Thysanoptera	Aeolothripidae	Minor	Sanjta and Chauhan, 2018
	<i>Haplothrips</i> sp.	Thysanoptera	Phlaeothripidae	Minor	Sanjta and Chauhan, 2018
Soybean leaf folder	<i>Nacoleia vulgaris</i>	Lepidoptera	Pyralidae	Minor	Triveni, 2017
Leaf webber	<i>Pronomis profusalis</i>	Lepidoptera	Pyralidae	Minor	Ramani and Bhumannavar, 1990
Leaf roller	<i>Psara stultalis</i>	Lepidoptera	Pyralidae	Minor	Ramani and Bhumannavar, 1990
Semilooper	<i>Thysanoplusia orichalcea</i>	Lepidoptera	Noctuidae	Minor	Triveni, 2017
Grasshopper	<i>Attractomorpha</i> sp.	Orthoptera	Acrididae	Major	Triveni, 2017
	<i>Acrida exaltata</i>	Orthoptera	Acrididae	Major	Triveni, 2017
	<i>Cantantops innotabilis</i>	Orthoptera	Acrididae	Major	Triveni, 2017
Mite	<i>Tetranychus</i> spp.	Trombidiformes	Tetranychidae	Major	Triveni, 2017
	<i>Brevipalpus phoenicoides</i>	Trombidiformes	Tenuipalpidae	Minor	Ramani and Bhumannavar, 1990



Fig.12. *Attractomorpha* sp

planting sites to no infested planting sites. Therefore it is essential and utmost important to use of nematode free planting material would check the spread of pathogen.

Patchouli (*Pogostemon patchouli* Pellet.), (Syn. *P. cablin* Benth.): Patchouli is an important aromatic plant, belonging to family Lamiaceae is the major source of commercial patchouli oil. In the perfumery industries still there is no synthetic chemical which could replace the patchouli oil. It is native of the Phillipines and cultivated mainly in Malaysia, Jawa, Brazil, West Indies, Singapore, Indonesia, China and India. It is a perennial aromatic herb that yields fragrant leaves containing very sweet smelling oil, attains a height of about 1m – 1.2 meter. The chemical components of patchouli oil are β -patchoulene, α -guaiene, caryophyllene, α -patchoulene, seychellene, α -bulnesene, norpatchoulenol, patchouli alcohol and pogostol. It is propagated by stem cuttings. The commercial oil of patchouli is obtained by steam distillation of the shade dried herbage. The oil of patchouli has a strong fixative property, which helps to prevent the rapid evaporation of perfume and thereby promotes tenacity. The oil is generally blended with other essential oils. It is used in a wide range of toilet soaps, scents, body lotions, etc. At very low concentration, the oil is extensively used to add flavour to major food products including alcoholic and non alcoholic



Figure 13. *Aphis gossypii*

beverages, frozen dairy desserts, candy etc. It also posses medicinal properties like helpful in relieving from all forms of depression, anxiety and stress related conditions. The other therapeutic properties of patchouli oil are antiseptic, aphrodisiac, astringent, deodorant, diuretic, febrifuge, fungicide, sedative and also used in several other pharmaceutical preparation. It is used as general tonic and a stimulant and helps the digestive system (Sarwar *et al.* 1982). In India this plant is cultivated in state of Assam, Madhya Pradesh, Tamil Nadu, Kerala and Karnataka. This crop suffers from fungal, viral and nematode diseases.

Insect pests of Patchouli: Triveni (2017) documented 16 species of pest species on patchouli (*Pogostemon cablin*) which included 10 species of defoliators belonging to Coleoptera, Lepidoptera and Orthoptera orders and 6 species of sucking pests (Hemiptera, Acarina) (Table 3). Among the different species of pests observed, *Attractomorpha* sp. (Fig.12), *Cantantops innotabilis*, *Acrida exaltata*, *Blosyrus* sp., *Myllocerus viridanus*, *Aphis gossypii* and *Tetranychus* spp. were major pests of patchouli. Both the nymphs and adults of grass hoppers, namely, *Attractomorpha* sp., *C. innotabilis* and *A. exaltata* act as defoliators, which feeds on the leaves of patchouli and showing nibbling



Fig. 14. Alternaria Blight of *Pogostemon cablin*

symptoms (Triveni 2017). Adult weevils, *Blosyrus* sp., and *M. viridanus* feeds on the leaf margin of patchouli (Triveni, 2017). Sanjta and Chauhan (2018) recorded infestation of five species of thrips, viz. *Bathrips melanicornis*, *Thrips carthami*, *Anaphothrips sudanensis*, *Aeolothrips* sp. and *Haplothrips* sp. on patchouli.

Cotton aphid, *Aphis gossypii* Glover (Hemiptera: Aphididae) has a worldwide distribution. The nymphs and adults are yellowish in colour (Fig.13). The alate as well as apterous females multiply parthenogenetically and viviparously. The female give birth to 8-22 nymphs which becomes adults in 7-9 days. Nymphs and adults of *A. gossypii* infest the tender shoots and the under surface of leaves in very large numbers and suck the sap. Severe infestation results in downward curling of leaves. Black sooty mould develops on the honey dew of the aphids which falls on leaves (Triveni, 2017).

Wilt disease: It is most important disease of several medicinal and aromatic plants but in patchouli is very serious disease caused by *Fusarium solani*.

Symptoms: The prevalence of this disease is mainly during kharif season. The infected plants are characterized by gradual yellowing and drying of leaves followed by premature death. The plants show discoloration of roots and collar regions of fully grown plants. Severe

disintegration of secondary roots surface is also observed. Singh and Angadi (1990) reported that *Fusarium solani* wilted plants of Patchouli symptoms were characterized by blackening of roots at collar regions of fully grown plants. Severe disintegration of secondary roots surface was also observed. Similarly yellowing of leaves were found in Alternaria blight of patchouli (Fig.14).

Host range: Leath and Kendall (1978) conducted experiments on host range of *F. solani* and showed the susceptibility of white clover and *Lotus corniculatus* to root rot pathogen. Cock foot and *Coronilla varia* were the most resistant while lucerne, red clover, subterranean clover and *Coronilla globosa* showed moderate reaction. Nielson and Mayer (1979) used *F. solani* causing root rot of stored sweet potatoes for studying the reaction with other crops. The results revealed that roots of the Cv. Jewel were more susceptible than those of jersey orange, georgia red, continental, nugget, fruits of apple, cucumber, egg plant, squash, tomato and potato which were susceptible. Moubasher *et al.* (1984) observed that *F. solani* and *F. oxysporum* were most common species infecting cotton, pea, wheat and tomato. Out of 16 different hosts treated for determining the host range of *F. solani*, black gram, green gram, bengal gram, tomato, brinjal, ladies finger, potato and guava were found to be susceptible. However, maize, peanut, linseed, sunflower, pigeonpea, soybean and safflower were free from infection (Kore and Patil, 1985). Patel (1991) reported considerable variation among 13 isolates of *F. solani*. Sharma and Agnihotri (1972) recorded morphological and pathogenic variation among three isolates of *F. orthocerus* App. & Wr.

Management

Essential oils of *Thymus vulgaris* L., *Lavandula angustifolia*, *Mentha piperita*, Neem oil, *Allium sativum* were tested in the

laboratory against *F. solani*. *T. vulgaris* was effective from a minimum concentration of 50 ppm. Mint and Lavender oils were only effective at higher concentrations (400-800 ppm). In the field, neem oil gave good protection against *F. solani* (Aulerio and Zambonelli 1997). Shivapuri *et al.* (1997) investigated that fungitoxic properties of different plant extracts against *F. solani* and other pathogenic fungi. Ethanol extract of *Allium cepa* L., *Allium sativum*, *Lantana camara* L., *Polyalthia longifolia* Benth and Hook, *Tagetes erecta* L., *Vinca rosea* L. and *Withania somnifera* showed fungitoxic effect against *F. solani*. The inhibitory effects of essential oils extracted from the genus Eucalyptus, *O.basilicum*, *Prosopis cineraria* and *Pongamia pinnata* were evaluated against *F. solani*. Among these oils extracted from the Eucalyptus species markedly inhibited fungal growth (Rai *et al.* 1999). The effects of aqueous leaf extract of 5 medicinal plants (*Strychnos nux-vomica*, *Calotropis procera*, *Azadiracta indica*, A. Juss, *Ocimum sanctum* and *Allium sativum*) on the spore germination of five species of *Fusarium* viz. *F. solani*, *F. moniliformae*, *F. equiseti*, *F. acuminatum* and *F. oxysporum* were examined. Spore germination of all the fungi tested was completely inhibited by 100% aqueous extract of *A. indica* while only 20% of germination of spores was observed with treatment of 100% aqueous extract of *A. sativum* (Tripathi *et al.* 1999). Leaf extracts of *L. camara* followed by *A. indica*, and *Acalypha indica* were found to be equally effective in inhibiting the growth of *F. solani* *in vitro*. Leaf extracts of *L. camara* has been reported to exhibit maximum toxicity against spore germination of *F. solani* (Mamatha and Ravishankar Rai 2004).

Sarhan (1989) observed that an isolate of *B. subtilis* antagonistic to *F. solani* in culture was found effective against root rot of Fabae beans in green house studies. Bacterium applied as a seed treatment significantly reduced seed

colonization and root rot. *Bacillus subtilis* inhibited the growth of *F. solani* on potato dextrose agar and resulted in significant spore reduction due to production of toxic metabolites. Patel (1991) found that *T. harzianum* initially showed two mm inhibition zone of *Fusarium* sp. and later it overgrew the colony of pathogen. Fluorescent Pseudomonads (FPs) have received much attention in the last few years for their ability to suppress plant diseases especially those caused by soil-borne plant pathogens (Thomashow and Weller, 1990). Fluorescent Pseudomonads with potential antagonistic activity and *Trichoderma harzianum* strain THA were found to be highly effective and inhibited the growth of *F. solani* in dual culture (Zapata *et al.* 2001). Twelve isolates of *Trichoderma* obtained during survey from soils in dry root-rot affected acid lime gardens of Andhra Pradesh were evaluated for their variation in phenotypic characters, growth rate and antagonistic potential against *F. solani*. Out of 12 isolates, only two isolates (T2 and T4) showed very fast growth rates and the antagonistic potential (Kavitha *et al.* 2004).

Several techniques for evaluating fungicides have been described from time to time. Poisoned food technique is the most commonly practiced method for evaluating fungicides under laboratory conditions (Flack 1907). Six fungicides were evaluated in laboratory against mycelial growth and conidial germination of *Fusarium solani* the cause of decay in colocasia and yam. carbendazim was found to be the best fungicide among the tested, in disease control (Mishra and Rath 1987). Kapoor and Kumar (1991) reported that carbendazim and benomyl (500 mg a.i. /ml) were found to be most toxic to *Fusarium solani*. Etebarian (1992) found that iprodione and carbendazim totally inhibited fungal growth of *F. solani* at 10 ppm and 100 ppm concentrations. Wahid *et al.* (1995) concluded that Derosol (carbendazim) and benlate (benomyl) at 10 ppm completely

Table 5. Major fungal diseases of *Pogostemon cablin*

Disease	Causal Organism	References
Alternaria blight	<i>A.alternata</i>	Parameshwaran <i>et al.</i> 1987
Leaf spot	<i>Corynespora cassiicola</i>	Chen <i>et al.</i> 2010

inhibited the growth of *F. solani* which was isolated from soybean seed. Topsin-M and vitavax gave 100 percent inhibition at 50 ppm.

Integration of biological and chemical control seems to be a promising way of controlling many pathogens with minimum interference in the biological equilibrium in soil (Papavizas, 1973). Integration of chemicals and bioagents (*Trichoderma* Sp.) has been the subject of intense research during recent years (Chet, 1991).

Certain strains of *Trichoderma hamatum* and *T. pseudokoingii* are reported to be adapted to excess soil moisture, whereas *T. viride* and *T. polysporum* are restricted to low temperature conditions. *T. harzianum* is most commonly distributed in warm climate region, whereas *T. harzianum* *T. koningii* are reported to be distributed under diverse environmental conditions. Species of *Trichoderma* have been shown to be effective against wide range of plant pathogens, such as *Botrytis* sp., *Fusarium* sp., *R.solani*, *Sclerotium rolfisii*, etc. Various mechanism of parasitism by *Trichoderma* sp. on different plant pathogenic fungi are reported to be varied such as inhibition by contact action, parasitism, hyphal interaction, coiling, penetration and lysis of hyphal cell, antibiosis and production of toxic metabolites etc. (Hriday 1991). Hartman and Fletcher (1991) reported *Trichoderma harzianum* to be



Figure 15. Yellow mosaic symptoms on *Pogostemon* efficient against *Fusarium* root rot of tomato. **Viral Diseases** (Table 6)

Patchouli is known to be infected by a number of viruses such as *Patchouli mosaic virus* (PaMV), *Tobacco necrosis virus* (TNV), *Patchouli mild mosaic virus* (PaMMV), *Patchouli mottle virus* (PaMoV), *Patchouli yellow mosaic virus* (PaYMV) and *Peanut stripe virus* (PStV) distributed all over world belonging different genera of the viruses.

Mosaic disease of *P. cablin* (Blanco) Benth. is widespread In India and has been reported by different workers. However the causal virus (es) was not fully characterized. A yellow mosaic disease has been reported on *P. cablin* (Blanco) Benth. From Lucknow (Zaim *et al.* 1999). The typical yellow mosaic appears when temperature is high during April to June

Table 5. Major viral diseases on *Pogostemon cablin* (Blanco) Benth. in India

S. No.	Viral Diseases	Genus	Transmission	References
1.	Yellow mosaic	--	Whitefly	Sastry and Vasentha kumar, 1981
2.	Patchouli mosaic	Potyvirus	Aphid & Mechanical	Subba Rao 1986
3.	Mixed infection of Patchouli	Potyvirus and Tobamovirus	Mechanical	Zaim <i>et al.</i> 1999
4.	Peanut stripe virus (PstV)	Potyvirus	--	Singh <i>et al.</i> 2009

(Fig. 15). These symptoms generally are masked during winter season. The Lucknow isolate of virus which is tentatively designated as patchouli yellow mosaic virus (PaYMV) was transmitted by sap and aphids. Host range was found to be restricted. The virus incited typical line pattern on the leaves of *Nicotiana benthamiana*, which is diagnostic host for this virus. Purified virus preparation showed flexuous rod shaped virus particles about 805 nm in length. The virus was detected by RT-PCR using potyvirus specific degenerate primers.

Nematode diseases : Root-knot nematode (*Meloidogyne incognita*, *M. javanica*, *M. hapla*) become most important constraint for the successful cultivation of patchouli in India.

Symptoms: Root-knot infested plants are weak and grow slowly. Heavy galling on root system with root knot nematode on patchouli results in stunting, wilting, defoliation and chlorosis of the plant. Sometimes root galls are very small or the surrounding galls coalesce to form large one up to 2-5 cm large even more. Infection of root-knot nematodes occurs when plants are in their early stage of development.

Causal organism and Host Parasite relationship: Root-knot diseases of patchouli caused by *Meloidogyne incognita*, *M. javanica*, *M. hapla*. Krishna Prasad and Reddy (1979, 1984) carried out the pathogenicity experiment and reported the threshold level of *M. incognita* on *P. cablin* was 4 larvae/g soil. They reported 46.7 and 52% loss in fresh and dry weight of patchouli by *M. incognita*. They further reported that the multiplication of *M. incognita* was more in sandy soil on patchouli than clay soil. Dijwanti and Momota (1991) reported the association of *M. incognita*, *M. hapla*, and *Pratylenchus brachyurus* with *P. cablin* in West Jawa.

Symptoms of damage: Root knot infested



Figure 16. Root-knot nematode infested field of patchouli

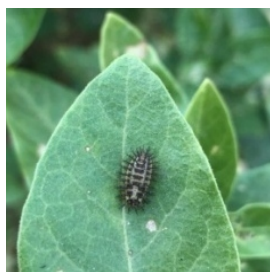
plants are weak and grow slowly. Heavy galling on root system with root knot nematode on patchouli results in stunting, wilting, defoliation and chlorosis of the plant (Fig. 16). Sometimes root galls are very small or the surrounding galls coalesce to form large one up to 2-5 cm large even more. Infection of root-knot nematodes occurs when plants are in their early stage of development.

Disease Management:

A large number of experiments were conducted to manage phytoparasitic nematode on patchouli. Krishnaprasad (1978) used aldicarb-sulfone, aldicarb-fensulphothion, carbofuran, cyerolan AC-92 and AC-100 @3.6 and 10kg a.i./ ha as pre and post inoculation treatment. Good control of *M. incognita* on patchouli was obtained by these chemicals. Sarwar *et al.* (1982) also conducted some experiment with fensulphothion, carbofuran, aldicarb, nemagon, metham sodium, fenamiphos to control root-knot nematode *M. incognita* on patchouli. Carbofuran was one of the most effective chemicals for *M. incognita* control. In other experiment effect of different oil seed cakes such as neem, pongamia and castor cake on plant growth and nematode population development was studied. Neem oil seed cake @4 tonnes / ha proved better than other oil seed



Eggs



Grub



Pupa



Adult

cakes for increasing growth and yield of the crop and reducing the *M. incognita* population. *Mucuna purita* as rotational crop with patchouli was found to be effective for reducing root-knot nematode population. Summer fallowing also proved to be a good method for reducing *M. incognita* population below threshold level. Kumar and Nanjan (1984) applied aldicarb, carbofuran or phorate @2-3 kg a.i. / ha to manage the *Helicotylenchus dihystra* on *P. patchouli*. Significant control of spiral nematode on this crop was observed and increase yield of crop was detected in all the treatment.

Ashwagandha (*Withania somnifera* (L.) Dunal): Ashwagandha (*Withania somnifera* L. Dunal), also known as Indian ginseng, belongs to the family Solanaceae, is an important ancient medicinal plant, used in the Indian traditional systems of medicine, ayurveda and unani. It is a widespread species having distributed from the southern mediterranean area to the different parts of Africa, India, Isreal, Jordan, Egypt, Sudan, Iran, Afghanistan and Pakistan. It grows in dry parts in subtropical regions. In India it is commercially cultivated in the states of Rajasthan, Punjab, Haryana, Uttar Pradesh, ujarat, Maharashtra, Tamil Nadu and Madhya Pradesh. The estimated annual production of ashwagandha roots in India is more than 2000 tonnes, against annual requirement of about 7000 tonnes (Patra *et al.* 2004). Ashwagandha is regarded as the treasure of biochemical constituents serving as remedy for many health problems. Roots contain several pyrazole alkaloids like withasomnine, withaferin A, withanolides,

withaninol and steroidal lactones, starch and reducing sugars. Withaferin is the chief constituent (0.13 to 0.31%) having bacteriostatic and antitumerous properties. It is considered to be one of the best rejuvenating agents in Ayurveda (Farooqi *et al.* 2003). Its roots, seeds and leaves are used in ayurvedic and unani medicines. Roots are used for treatment of rheumatic pain, inflammation of joints, nervous disorders and epilepsy, and also used as tonic for hiccup, cold, cough and female disorders. The leaves are also useful for carbuncles, inflammation, swellings and conjunctivitis.

Insect pests of Ashwagandha, *Withania somnifera* (Table 7)

Spotted leaf eating beetle/Hadda beetle, *Henosepilachna vigintioctopunctata* (Fabricius) (Coleoptera: Coccinellidae)

Distribution: *Henosepilachna vigintioctopunctata* (Synonym: *Epilachna vigintioctopunctata* L.) is widely distributed in Southeast Asian countries, Korea, Australia, Sri Lanka, China, Japan including India (Kapur 1950, Richards 1983).

Host plants : The spotted leaf beetle is considered is a polyphagous pest and is considered as voracious foliage feeder of various cultivated and wild plants belonging to the families solanaceae (potato, brinjal, tomato, tobacco, *Datura* spp., *Physalis* spp., *Solanum nigrum*) and cucurbitaceae (melon, gourds, cucumber etc.) in different parts of India (Mohanasundaram and Uthamaswamy, 1973; Ghosh and Senapati 2001, Venkatesh 2006; Varma and Anandhi 2008). Spotted leaf beetle has been recorded as one of the major

Table 7. Insect pests of Ashwagandha, *Withania somnifera*

Common name	Scientific name	Order	Family	Pest Status	References
Epilachna beetle	<i>Henosepilachna vigintioctopunctata</i>	Coleoptera	Coccinellidae	Major	Mathur and Srivastava, 1964; Kumar <i>et al.</i> 2009; Chaudhary, 2013; Nirmal <i>et al.</i> , 2015; Rehaman <i>et al.</i> , 2018; Kumar <i>et al.</i> , 2009 b
Fruit borer	<i>Helicoverpa armigera</i>	Lepidoptera	Noctuidae	Major	Hanumanthaswamy <i>et al.</i> , 1994; Kumar <i>et al.</i> 2009 a; Kumar <i>et al.</i> , 2009 b; Chaudhary, 2013; Nirmal <i>et al.</i> , 2015; Rehaman <i>et al.</i> , 2018
Oleander hawk-moth	<i>Deilephila nerii</i>	Lepidoptera	Sphingidae	Minor	Hanumanthaswamy <i>et al.</i> , 1994; Nirmal <i>et al.</i> , 2015; Rehaman <i>et al.</i> , 2018
Metallic blue	<i>Corynodes peregrines</i>	Coleoptera	Chrysomelidae	Minor	Chaudhary, 2013
Snout beetle	<i>Cyrtozemia dispar</i>	Coleoptera	Curculionidae	Minor	Chaudhary, 2013
Weevils	<i>Myllocerus viridanus</i>	Coleoptera	Curculionidae	Minor	Hanumanthaswamy <i>et al.</i> 1994
	<i>Myllocerus discolor</i>	Coleoptera	Curculionidae	Minor	Ramanna <i>et al.</i> , 2010
	<i>Blosyrus inaequalis</i>	Coleoptera	Curculionidae	Minor	Hanumanthaswamy <i>et al.</i> , 1994
Semi looper	<i>Hyposidra successaria</i>	Lepidoptera	Geometridae	Minor	Chaudhary, 2013
Til hawk moth	<i>Acherontia styx</i>	Lepidoptera	Sphingidae	Minor	Manjoo, 2006
Grass hopper	<i>Crotogonus trachypterus</i>	Orthoptera	Acrididae	Minor	Chaudhary, 2013
	<i>Acrida exaltata</i>	Orthoptera	Acrididae	Minor	Hanumanthaswamy <i>et al.</i> , 1994
	<i>Trilophida annulata</i>	Orthoptera	Tettigoniidae	Minor	Hanumanthaswamy <i>et al.</i> , 1994
Solenopsis mealybug	<i>Phenacoccus solenopsis</i>	Hemiptera	Pseudococcidae	Major	Chaudhary, 2013; Sharma and Pati, 2013; Kedar and Saini; 2015
Eggplant mealybug	<i>Coccidohystrix insolitus</i>	Hemiptera	Pseudococcidae	Major	Hanumanthaswamy <i>et al.</i> , 1994
Striped mealybug	<i>Ferrisia virgata</i>	Hemiptera	Pseudococcidae	Minor	Ramanna <i>et al.</i> , 2010
Citrus mealybug,	<i>Planococcus citri</i>	Hemiptera	Pseudococcidae	Minor	Attia and Awadallah, 2016
Mango mealybug	<i>Drosicha mangiferae</i>	Hemiptera	Margarodidae	Major	Sharma <i>et al.</i> , 2014 Bhagat, 2004; Kumar <i>et al.</i> , 2009 b
Aphids	<i>Myzus persicae</i>	Hemiptera	Aphididae	Major	Kumar <i>et al.</i> , 2009a; Kumar <i>et al.</i> , 2010
	<i>Aphis gossypii</i>	Hemiptera	Aphididae	Minor	Chaudhary, 2013; Rohini <i>et al.</i> , 2018
Tree hopper	<i>Oxyrachis tarandus</i>	Hemiptera	Membracidae	Minor	Ramanna <i>et al.</i> , 2010; Sharma and Pati, 2011
	<i>Tricentrus bicolor</i>	Hemiptera	Membracidae	Major	Kumar <i>et al.</i> 2009; Kumar <i>et al.</i> , 2010
	<i>Tricentrus congestus</i>	Hemiptera	Membracidae	Minor	Mitra and Biswas, 2002
	<i>Otinotus oneratus</i>	Hemiptera	Membracidae	Minor	Mitra and Biswas, 2002; Chaudhary, 2013
	<i>Leptocentrus taurus</i>	Hemiptera	Membracidae	Minor	Chaudhary, 2013;
	<i>Leptocentrus substitutus</i>	Hemiptera	Membracidae	Minor	Mitra and Biswas, 2002; Rehaman <i>et al.</i> , 2018;
Lantana bug	<i>Orthezia insignis</i>	Hemiptera	Orthezidae	Minor	Hanumanthaswamy <i>et al.</i> , 1994
Jassids	<i>Amrasca biguttula biguttula</i>	Hemiptera	Cicadellidae	Minor	Chaudhary, 2013
	<i>Acomurella prolix</i>	Hemiptera	Cicadellidae	Minor	Chaudhary, 2013
	<i>Nephotettix virescens</i>	Hemiptera	Cicadellidae	Minor	Chaudhary, 2013
	<i>Balclutha incise</i>	Hemiptera	Cicadellidae	Minor	Chaudhary, 2013
	<i>Balclutha saltuella</i>	Hemiptera	Cicadellidae	Minor	Chaudhary, 2013
	<i>Penthimia sp.</i>	Hemiptera	Cicadellidae	Minor	Hanumanthaswamy <i>et al.</i> , 1994
Spittle bug	<i>Poophilus costalis</i>	Hemiptera	Aphrophoridae	Minor	Mitra and Biswas, 2002
Lygaeid bugs	<i>Graptostethus servus</i>	Hemiptera	Lygaeidae	Minor	Chaudhary, 2013
Seed bug	<i>Spilostethus pandurus</i>	Hemiptera	Lygaeidae	Minor	Chaudhary, 2013
Seed bug	<i>Spilostethus hospes</i>	Hemiptera	Lygaeidae	Minor	Mitra and Biswas, 2002; Manjoo, 2006; Rehaman <i>et al.</i> , 2018
Stink bug	<i>Aspongopus janus</i>	Hemiptera	Pentatomidae	Minor	Chaudhary, 2013;
	<i>Nezara viridula</i>	Hemiptera	Pentatomidae	Minor	Chaudhary, 2013; Nirmal <i>et al.</i> , 2015; Mitra and Biswas, 2002; Kumar <i>et al.</i> , 2009 b
	<i>Plautia fimbriata</i>	Hemiptera	Pentatomidae	Minor	Mitra and Biswas, 2002; Kumar <i>et al.</i> , 2009 b
	<i>Agonoscelis nubila</i>	Hemiptera	Pentatomidae	Minor	Mitra and Biswas, 2002
	<i>Acrosternum gramineum</i>	Hemiptera	Pentatomidae	Minor	Rehaman <i>et al.</i> , 2018

Black shield bug	<i>Dollicoris indicus</i>	Hemiptera	Pentatomidae	Minor	Rehaman <i>et al.</i> , 2018
Red cotton bug	<i>Dysdercus cingulatus</i>	Hemiptera	Pyrhocoridae	Minor	Sharma <i>et al.</i> , 2014
Coreid bug	<i>Acanthocoris sordidu</i>	Hemiptera	Coreidae	Minor	Manjoo, 2006
Greenhouse whitefly	<i>Trialearodes vaporariorum</i>	Hemiptera	Aleyrodidae	Minor	Kumar <i>et al.</i> , 2009 b
White grub	<i>Holotrichia serrata</i>	Coleoptera	Scarabaeidae	Major	Meshram, 2005
Pea nut Trash bug	<i>Elasmolomus pallens</i>	Hemiptera	Rhyarochromidae	Minor	Rehaman <i>et al.</i> , 2018
Red Spider mite	<i>Tetranychu urticae</i>	Trombidiformes	Tetranychidae	Major	Gupta and Karmakar, 2011; Chaudhary, 2013; Sharma and Pati, 2012; Rehaman <i>et al.</i> , 2018
Mite	<i>Tetranychus macfarlanei</i>	Trombidiformes	Tetranychidae	Major	Gupta and Karmakar, 2011
Oriental red mite	<i>Eutetranychus orientalis</i>	Trombidiformes	Tetranychidae	Major	Gupta and Karmakar, 2011
Broad mite	<i>Polyphagotarsonemus latus</i>	Trombidiformes	Tarsonemidae	Major	Gupta and Karmakar, 2011
Mite	<i>Brevipalpus phoenicis</i>	Trombidiformes	Tenuipalpidae	Major	Gupta and Karmakar, 2011

pest of *Withania somnifera* causing severe foliage damage to crop (Mathur and Srivastava, 1964; and Ramanna *et al.* 2010).

Identification and Biology: The brownish hemispherical beetle has 28 lack spots on elytra (Fig. 17). The female lays elongate, spindle-shaped yellowish eggs in groups of 10 to 20 on the under surface of leaves (Fig.17). The gravid females lay 562–1500 eggs in several batches during the ovipositional period. The incubation period is 2-5days. The yellowish spiny grubs become full grown in 10-30 days and pupate on the leaf or stem. The grub passes through four instars (Venkatesha, 2006). The first instars feed gregariously on the foliage. The pupa is yellowish with spines on the posterior part, the anterior portion being devoid of spines. The pupal period is about 5-9 days under optimal conditions. The total life cycle is completed in 17 to 50 days depending on weather conditions. Male and female longevity has been recorded as 27–108 days and 37–108 days, respectively (Venkatesha 2006).

Nature of Damage: The grubs scrape the epidermal layer of leaves and in severe infestation the epidermis is entirely removed (Mathur and Srivastava, 1964). The nature damage by adult beetle is same as of grubs.

Gram pod borer, *Helicoverpa armigera*

(Hübner) (Lepidoptera: Noctuidae)

Distribution: *Helicoverpa armigera* is an important polyphagous pest widely distributed in the tropics, subtropics and warmer temperate regions of the world including India (CABI, 2020). This species is a well known important pest of pulses and cotton in India.

Host Plants: *Helicoverpa armigera* is polyphagous in nature and feeds on various agricultural crops. The important host plants of *H. armigera* are gram, pigeon pea, tomato, maize, cotton, okra etc. The pest also feeds on many weed species during off season.

Identification and Biology: The moth is medium sized and with 30-38 mm wing span, V-shaped speck on the light brownish forewings and a dark border on the hindwings (Fig. 18). Sexual dimorphism is distinct, male moth is light brown and female is dark brown. The female moth lays 600-1000 spherical yellowish white eggs singly on the tender parts of plants. Incubation period varies from 3-10 days depending on the weather conditions. Caterpillar has six growth stages (instars) takes 18-25 days to fully develop. The full grown caterpillar measures about 35-45 mm in length and is greenish with dark grey lines laterally on the body. The egg, larval and pupal periods is 2-4, 18-25 and 6-21 days, respectively. The caterpillar passes through five to six instars.



Fig. 18. *Helicoverpa armigera*

Under optimal conditions the pest completes 11 generations in a year.

Nature of damage: The larvae feed on tender foliage, flowers and later instars damage the berries. They bore circular holes in berries of *Withania somnifera* keeping the head portion inside the hole to eat the inner content (Rehaman *et al.* 2018).

Solenopsis mealybug, *Phenacoccus solenopsis* Tinsley (Hemiptera: Pseudococcidae)

Distribution: *Phenacoccus solenopsis* was originally described from USA in 1898 (Tinsley 1898). *P. solenopsis* is distributed over a wide range of agro-ecological zones in more than 24 countries worldwide including India (Fand and Suroshe 2015).

Host plants: *P. solenopsis* is highly polyphagous in nature damaging over 202 plant species of 55 families across the globe comprising field crops, vegetables, ornamentals, fruit trees, medicinal and weeds. Most of the host plants belonging to plant families Malvaceae, Solanaceae, Asteraceae, Euphorbiaceae, Amaranthaceae and Cucurbitaceae. The pest causes severe damage to cotton followed by okra, brinjal, tomato, sesame, sunflower and China rose. The wild host plants supports carryover of the pest from one season to other season (Fand and Suroshe, 2015, Kedar and Saini 2015). The incidence of *P. solenopsis* as major pest on *Withania somnifera* has been recorded by several



workers in India (Vennilla *et al.* 2011; Sharma and Pati 2013, Kedar *et al.* 2013; Sahu *et al.* 2017).

Identification and Biology: The mealybug is cottony in appearance, small, oval, soft-bodied covered with white mealy wax (Fig.19). Adult females are wingless, 2.5-4.0 mm long with 9 segmented antennae and long flagellate dorsal setae. Short filaments around the body and dark stripes on the dorsal side are present. The anal filaments are about one fourth the length of the body. Females produce egg mass in an ovisac.



Figure 19. *Solenopsis mealybug, Phenacoccus solenopsis*

Males have one pair of wings, long antennae and absence of mouth parts with white wax filaments projecting posteriorly (Kedar *et al.*, 2013). The developmental period of first, second and third instars are 3-4, 5-8 and 4-5 days, respectively. Males have an additional pupal stage of 5-7 days. The total developmental period is 13-15 days for female and 18-19 days for male. Reproduction is parthenogenetic and ovoviviparous (Kedar *et al.* 2013). A female may produce 120-800 crawlers and duration of reproduction varies from 10 to 47 days. Life cycle of female mealybug is completed in 25-38 days and male mealybug in 17-27 days (Kedar *et al.* 2011). The mealybug completes 15 generations per year.

Nature of Damage: *P. solenopsis* nymphs and adults suck the sap and once established on a plant often remains there till the attacked parts get dried up, causing even death of the whole plant. When the attack is in the vegetative phase, the plants remain stunted; the attacked parts gradually dry up; and the plants bear few or no fruiting bodies. The pest secretes large amounts of honey dew which fall on the underlying plant parts and sooty mould develops on them causing their blackening. The development of sooty mould reduces the photosynthetic ability of the plants.

Eggplant mealybug, *Coccidohystrix insolitus* (Green), (Homoptera: Pseudococcidae)

Distribution : The eggplant mealybug, *Coccidohystrix insolita* (Green) is broadly distributed in the tropics and subtropics and well known as a agricultural and horticultural pest (Lit *et al.* 1998, Williams 2004, Williams and Watson 1988).

Host plants: *Coccidohystrix insolitus* is polyphagous and recorded on host plants belonging to families *Solanaceae*, *Fabaceae*, *Malvaceae*, *Amaranthaceae*, *Cucurbitaceae*,



Figure 20. Eggplant mealybug, *Coccidohystrix insolitus*

Acanthaceae, etc. (Ben-Dov 2020).

Nature of Damage: Nymphs and adults of mealybugs suck sap from the under surface of leaves and tender shoots (Fig.20). The attack results in yellowing and sometimes dying of plants. A heavy black sooty mould may develop on the honeydew like droplets secreted by mealybugs.

Red spider mite, *Tetranychus urticae* Koch (Trombidiformes: Tetranychidae)

Distribution : Red spider mite emerged as a serious pest of eggplant, tomato, French bean, okra, cucumber, rose and other field crops in South Asia, Southeast Asia, Africa, Europe, and Mediterranean countries. On *Withania somnifera* the damage of *T. urticae* has been reported by several workers from different states of India (Gupta and Karmakar 2011; Chaudhary 2013, Sharma and Pati 2012; Rehaman *et al.* 2018).

Identification and Biology: They are minute in size, and vary in color (green, greenish yellow, brown, or orange red) with two dark spots on the body (Fig. 21). Reddish spherical eggs are laid on an average 100 eggs by a female. After hatching it passes through a larval stage and two nymphal stages (protonymph and deutonymph) before becoming adult. Newly hatched larva is pale



Figure 21. *Tetranychus urticae* a) adult; b) nymphs; c) egg

white. At the deutonymph sex can be distinguished, males being smaller and females larger with darker colour. Life cycle is completed in 5 to 12 days and in a year about 40 generations are completed.

Nature of Damage:The incidence of mites is observed during the warm and dry periods of the season. It is generally found on the under surface of the leaves but, in heavy infestations, it may be found all over the leaf surface. Nymph and adult mites punctures the leaf tissue and oozing plant sap is sucked. The removal of plant sap with chlorophyll and other plants pigments results in the characteristic blotching of leaves. The affected leaves become mottled. Mite infested plants can be diagnosed from distance by characteristic mottling symptoms produced on the upper surface of the leaves. Under severe infestation, the mites move to the tip of the leaves or top of the plant and congregate using strands of silk to form a ball like mass, which will be blown by winds to new leaves or plants, in a process known as “ballooning”.

Green peach aphid, *Myzus persicae* (Sulzer) (Hemiptera: Aphididae)

Distribution : *Myzus persicae* is of Asian

origin distributed over Africa, America, the Caribbean, Europe, Oceania, Australia, Fiji, New Zealand etc. It occurs everywhere in the world except where there are extremes of temperature or humidity

Host plants :*Myzus persicae* is highly polyphagous on summer host plants. Infestation of green peach aphids has been reported hundreds of host plants in over 40 plant families including *Withania somnifera* (Blackman and Eastop 2007, Kumar *et al.* 2009).

Identification and Biology;Green peach aphid adults are about 2 mm long, greenish-yellow or pink, with red eyes. Adults may be winged or wingless (Fig.22). Winged aphids have a black head and thorax and a yellowish green abdomen with a large dark patch dorsally. The nymphs are similar to the wingless adult, except in size. Eggs are initially yellow or green but soon turn black. Nymphs are initially greenish but soon turn yellowish, greatly resembling viviparous (parthenogenetic, nymph-producing) adults (Margaritopoulos *et al.* 2002). In tropical countries, male do not exist. Females reproduce parthenogenetically. The life cycle is completed in about 12-18 days. There are about 20 generations of the pest per year.

Nature of Damage;Nymphs and adults of aphids feed on the young foliage of *W. somnifera* and stems during December to March in North Indian conditions (Kumar *et al.* 2009). They suck the cell sap from tender



Figure 22. *Myzus persicae*

shoots which often results in drying out shoots, causing wilts and distortion. Aphids produce honeydew which falls onto foliage and which leads to black sooty mould fungi development. The development of sooty mould reduces the photosynthetic ability of the plants. *M. persicae* is most important aphid virus vector. More than 100 plant viruses have been reported to be transmitted by *M. persicae*.

Management of insect pests of *Withania somnifera*:

Ayyar (1984) suggested hand collection and destruction of egg masses, larval and adult stages of epilachna beetle in the field could be useful for control the pest population effectively. Venkatesha (2006) reported 51.91 per cent population of *H. vigintioctopunctata* was naturally controlled by a larval parasitoid *Pediobius foveolatus* in Karnataka province of India. Chandrananth and Katti (2010) evaluated six insecticides under field conditions for management of epilachna beetle, and recommended dimethoate 30EC (1 ml/litre), fenvalerate 20EC (0.5 ml/litre) and quinalphos 25EC (2 ml/litre); whereas less effective insecticides included chlorpyrifos 20EC (2 ml/litre), endosulphan 35EC (2 ml/litre) and neem seed kernel extract 5% (NSKE 5%). They reported least epilachna beetle population (4.3 beetles/plant) in treated plot compared to 14.9 beetles/plant in untreated plot of *Withania somnifera*, with highest root yield of 4.60 q/ha (3.20 q/ha in the untreated control plot).

To check the fruit damage from feeding of *Helicoverpa armigera* on *Withania somnifera* Rehman and Pradeep (2016) evaluated organic insecticides in a field trial and found highest per cent reduction of *H. armigera* infestation with Ha NPV (250 LE), which was on par with nimbecidine (3 ml/L) followed by NSKE (5%) and neem oil (5 ml/L).

Ravikumar *et al.* (2008) reported soil application of NO (3%) in a mixture of farm yard manure (12.5 t/ha) and a commercial biofertilizer Azophos (containing a mixture of

azospirillum and phosphobacteria) at 2 kg/ha resulted in significant reduction in damage caused by mealybug, *C. insolitus* and epilachna beetle, *H. vigintioctopunctata*.

Mealybug, *Phenacoccus solenopsis* management primarily depends on the use of chemical insecticides. Commonly used insecticides for managing mealybug infestations included profenophos, monocrotophos, malathion, quinalphos, methyl demeton, triazophos, dichlorvos, acephate, acetamiprid, imidacloprid, thiamethoxam, alphamethrin, buprofezin, cypermethrin, deltamethrin, endosulfan and chlorpyrifos (Nagarare *et al.* 2011, Kumar *et al.* 2012). Biocontrol agents are successful means of suppressing invasive mealybugs providing a non-toxic, self-perpetuating control method. Several predators and parasitoids have been reported that feed on *P. solenopsis* (Fand *et al.* 2010; Kedar *et al.* 2011; Suroshe *et al.* 2013). Over 88 per cent natural parasitization of *P. solenopsis* population by solitary endoparasitoid *Aenasius arizonensis* (= *Aenasius bambawalei*) has been reported from different agroecosystems by several workers (Ram *et al.* 2009, Dhawan *et al.*, 2011, Kedar *et al.* 2012). For suppression of the *P. solenopsis* several findings revealed conservation of parasitoid, *A. arizonensis* prior to applying chemical insecticides (Ram *et al.* 2009). Nagarare *et al.* (2011) reported 45-60 per cent mortality of *P. solenopsis* caused by entomopathogenic fungi viz., *Verticillium lecanii* (Zimm.), *Beauveria bassiana* (Bals.) and *Metarhizium anisoplae* (Metchnikoff) under laboratory conditions.

Hirekurubar *et al.* (2018) evaluated the bio efficacy of different insecticides and acaricides viz., acetamiprid 20SP, thiamethoxam 25WG, spiromesifen 240SC, difenthiuron 50WP, azadirachtin 1500 ppm, NSKE 5 per cent, dimethoate 30 EC and dicofol 18.5EC against red spider mite, *Tetranychus urticae* infesting *Withania somnifera*. Significant reduction in red spider mite population was noticed in plots sprayed with acaricide spiromesifen 240SC (1

ml/l) followed by difenthiuron 50WP (0.75 g/l) and dicofol 18.5 EC @ 2.50 ml/l. Spiromesifen sprayed treatment registered maximum root and seed yield.

Myzus persicae (Hemiptera: Aphididae) is a major pest of young shoots (Kumar *et al.*, 2009). Meena *et al.* (2016) evaluated bio-efficacy of some botanical products against mixed population of aphids (*Myzus persicae*, *Hyadaphis coriandri* and *Aphis craccivora*) on coriander. The highest reduction in aphid population was recorded in the plots treated with organic salt (bio-product 5ml/l) followed by karel extract 10ml/l and tumba fruit extract 10ml/l.

Fungal diseases: Root-rot and wilt is one of the most important disease of this crop in the nurseries and commercial field ashwagandha growing areas of Uttar Pradesh. The diseases occur during April-May and caused 30-50% mortality of plants (Gupta *et al.* 2004). The disease is caused by *Fusarium solani*.

Diagnosis: Many fungi can cause root rots. Often, it is possible to identify which fungus is responsible either by observing the structure of the fungus in the roots using a microscope or by placing infected roots on artificial media or baits (apple, carrot, or potato pieces) and allowing the fungus to grow out where it can be detected and then identified.

Symptoms: The major symptoms of the disease is withering and drooping of plants whereas at advance stage it causes severe wilting leading to decomposing and death of underground parts. The cottony growth appeared around the infected roots under moist condition. The infected roots become soft and brown in colour. In addition to killing plants and thereby reducing the quantity of saleable crop, root rots can also slow or stop plant growth and thus suppress plant quality. Root-rotted plants are usually smaller, less vigorous, produce fewer and/or small leaves, flowers, and fruit than healthy plants of equal age. Flowering may be delayed when the plant's roots are

rotted. As a result, the crop quality is very uneven. Root rots must be managed early in the disease if the losses are to be avoided.

Symptoms of Root Rot

- Growth of infected plants slows as compared to healthy plants.
- Older leaves yellow and fall.
- Margins of leaves die.
- Roots appear dark brown or black and few or no white roots or root tips can be found when the root ball is washed free of soil
- Roots are limp and not brittle and crisp as is found in healthy plants of all types.
- When plants are pulled from the potting mix, the outer layer of cells strips off the roots leaving only the central strand of water conducting tissue.

Management

Many effective pesticides have been tested against soil borne pathogens but not considered as long term solution because of concerns about exposure risks, health and environmental hazards, expensiveness, residue persistence, development of resistance to pesticides and elimination of natural enemies. The disease root rot and wilt complex of ashwagandha is caused by soil borne pathogens. In nature, plants are rarely exposed to the influence of single pathogen. Fawcett (1931) recognized that "nature does not work with pure cultures" and that many plant diseases are influenced by associated organisms. Before any action is taken, a diagnosis must be made of the actual cause of the symptoms. If the damage is due to the activity of fungi, chemicals are sometimes available which can check the fungus and allow the plant to grow. The dead roots do not recover. New roots must grow. The fungus is usually not completely eradicated (killed) by chemicals. Some fungus usually remains alive although its growth is greatly slowed as long as the fungicide is in high enough concentration. Therefore, repeated applications of fungicides

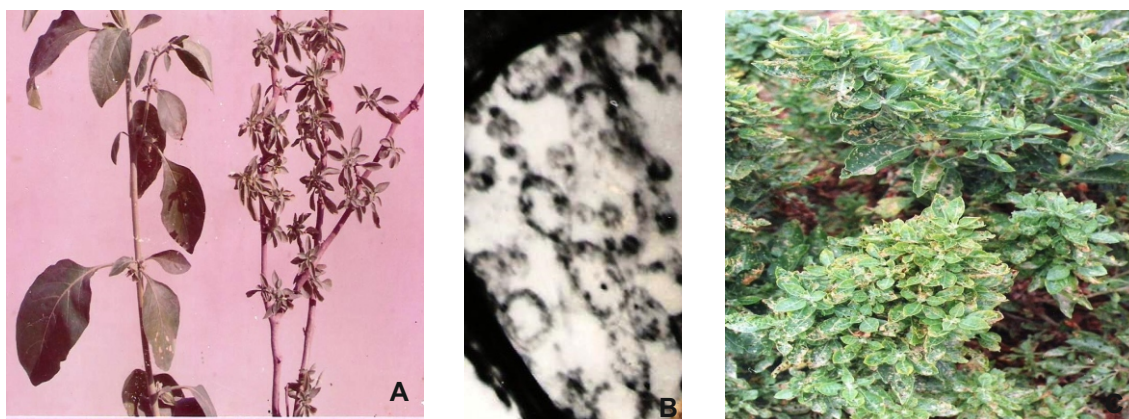


Fig.23. Phytoplasma infected plant along with healthy (A), electron microscopy of Phytoplasma infected *W. somnifera* (B), and field view of naturally infected plants (C).

are necessary.

Phytoplasma disease: In CIMAP, Lucknow the Witches broom disease was first observed in 1988 and more than 30% plants were affected by this disease.

Symptoms: This disease was reported for the first time from Lucknow (Zaim and Samad 1995). The succession of symptoms appearance include, little leaf, phyllody, dense clusters of highly proliferating branches with shortened internodes resulting witches'-broom and bushy appearance of infected plants (Fig 23 –A, B & C).

Causal organism: In subsequent studies *Withania* phytoplasma was found to be PCR amplified using universal primers, designed from the 16S rDNA sequences of phytoplasmas

Table 8. Major viral & phytoplasma Diseases on *Withania somnifera* (L.) Dunal in India

S. No	Name of Diseases	Causal agent	Host	References
1.	Witches-broom disease	Phytoplasma	<i>W. somnifera</i>	Zaim and Samad 1995
2.	Little leaf disease	Phytoplasma-Aster yellows group/Genus	<i>W. somnifera</i>	Khan <i>et.al</i> 2006
3.	Witches-broom disease	Phytoplasma-Clover proliferation group	<i>W. somnifera</i>	Samad <i>et.al.</i> 2006
4.	Tobacco leaf curl virus	--	<i>W. somnifera</i>	Pathak and Raychoudhuri , 1967
5.	A yellow mosaic disease	<i>Begomovirus</i>	<i>W. somnifera</i>	Baghel <i>et al.</i> 2010

and sequenced. Nucleotide sequence data and phylogenetic studies suggested that the *Withania* phytoplasma be assigned to the Aster yellows group (Khan *et al.* 2006). In another studies the *Withania* phytoplasma was detected by PCR using by universal phytoplasma primers P1/P6. Upon sequencing, the PCR product showed highest similarity with several isolates of the 16SrVI group of phytoplasmas and classified as a member of Clover proliferation group (16SrVI) (Samad *et al.* 2006). Other viral and phytoplasma diseases of *Withania somnifera* have been given in table 8.

Nematodes of Ashwagandha:

Root-knot disease of ashwagandha caused by the root-knot nematode *Meloidogyne incognita* (Kofoid and White) Chitwood, is very widespread affecting more than 80% plants in India (table 8). Race identification was done using host differential test and was identified as race-2 (Pandey and Kalra 2003). Nematodes were multiplied on roots of tomato and pathogenicity of *M. incognita* was confirmed on 10-day-old potted plant of *W. somnifera*. Large number of other phytoparasitic nematodes were found to be

Table 9. Loss in total alkaloids in healthy and root-knot infested roots of *Withania somnifera**

Treatments	Alkaloids (%)
Healthy root	0.32
<i>M. incognita</i> severely infested roots	0.11

associated in root rhizosphere of ashwagandha.

Symptoms of damage:The nematode infected plants typically show chlorotic, stunted, less branched with fewer and smaller leaves and poor response to fertilizer and irrigation. Such symptoms usually are not noticeable until severe damage to root system has been done by the nematodes. Roots of such plant were severely galled. When stem touches the soil it was also found to be infested with root-knot nematode *Meloidogyne incognita*. It was also noticed that root-knot nematode infected plants are more likely to be killed early with adverse effect of environment than healthy noninfested plants.

Control measures:Management studies were carried out (Pandey and Kalra, 2003) through the use of different organic materials and bioagent on growth/ yield of *W. somnifera* and nematode reproduction. In the experimentation neem compound, *Artemisia annua* marc, *Mentha* and *Murraya koenigii* distillates were found highly useful to suppress the root-knot development and *M. incognita* population on *W. somnifera*. Bio-agents however were found less effective as compared to organic materials on *W. somnifera*; may be due to host reactions towards the bio-agents. Integration of Vermicompost with *Trichoderma harzianum* and *Mentha* distillates with *Glomus aggregatum* were found nematode suppressive

and enhanced the growth of *W. somnifera* significantly. It was concluded that neem compound, *A. annua* marc, vermicompost and their integration with bio-agents can be an important way for nematode management, which could pave the way in future for integrated nematode management programme on agricultural and commercial crops.

Use of bio-organics (organic materials, bioagent) is a potential alternative to the environment detrimental chemical nematicides, which are generally used to restrict nematode infestation in agricultural crops. Pandey and Kalra (2011, In press) used different bio-organics viz. farm yard manures (Fym), cow urine, neem cake, vermicompost, *Trichoderma harzianum* separately and in dual combinations against root-knot disease of *Withania somnifera* and successful management of root-knot nematode on this crop were achieved.

Henbanes (*Hyoscyamus species*)

Henbanes (*Hyoscyamus muticus*, *H. niger* & *H. albus*) are important tropane alkaloid bearing plants belonging to family Solanaceae and one of the chief source of tropane alkaloids viz. hyoscine or scopolamine, hyoscyamine and atropine which are obtained from the dried leaves and other plant parts (Table 7). The hyoscine and its derivatives are used in pharmaceutical preparations, since they have

Table 10. Insect pests of Henbane

Common name	Scientific name	Order	Family	Pest Status	References
Henbane Flea Beetle	<i>Psylliodes hyoscyami</i>	Coleoptera	Chrysomelidae	Major	Parfentjev, 1921; Newton, 1934
Colorado potato beetle	<i>Leptinotarsa decemlineata</i>	Coleoptera	Chrysomelidae	Minor	Hsiao, 1978; Hubert, 1950
Three-lined potato beetle	<i>Lema trilineata daturaphila</i>	Coleoptera	Chrysomelidae	Minor	Kogan and Goeden, 1970
Bordered straw moth	<i>Heliothis peltigera</i>	Lepidoptera	Noctuidae	Minor	Zacher, 1921; Shchegolev, 1929; Printz, 1925
Green peach Aphid	<i>Myzus persicae</i>	Hemiptera	Aphididae	Major	Singh <i>et al.</i> , 1988
Potato psyllid	<i>Bactericera cockerelli</i>	Hemiptera	Triozidae	Minor	Knowlton and Thomas, 1934
Belladonna leaf-miner	<i>Pegomyia hyoscyami</i>	Diptera	Anthomyiidae	Minor	Cameron, 1914

anticholinergic, antiphasmodic and mydriatic properties. Leaves, seed and flowering tops of the plant are used in respiratory diseases and old cough. These plants originated in Egypt and spread to rest of the tropics. The two species of henbane i.e. black henbane (*H. niger*) and Egyptian henbane (*H. muticus*) are widely cultivated for pharmaceutical purposes.

Insect pests of Henbane

Incidence of insect pests in *Hyoscyamus* spp. is not as severe as in other medicinal and aromatic crops. However, a number of insect pests have been observed to damage the crop growth to variable level depending on climatic conditions and crop management practices (Table 9).

Henbane Flea Beetle, *Psylliodes hyoscyami* Linn. (Coleoptera: Chrysomelidae)

Distribution :Palaeartic, including the Middle East and North Africa. Widely scattered in England, South Wales and Scotland (West Lothian).

Host plants:*Hyoscyamus niger*, *Atropa Belladonna* (Van Emden 1924).

Identification and Biology:Adult is 2.9-3.5mm in length, blackish with blue-green or dark bronze reflection; may appear metallic green or bronze, sometimes coppery or blue-green. The egg is oval, yellowish white in colour with a deeper yellow equatorial band in the later stages. The full-fed larva is of the same general appearance and elongate form as other flea-beetle larvae. The body colour is white with the head dark brown, prothoracic shield, anal shield and legs light brown. The pupa is of the typical flea-beetle form. It is white in colour. The female lays eggs in the soil which hatch about in 2 weeks. The larval stages passed within the plant leaf-stalks mines. The prepupa constructs the usual earthen cell, lining the inside surface with a cementing secretion. The majority of the pupae lie between 5 and 7.5

Table 11. Major fungal and bacterial diseases of *Henbane* sp.

Disease	Causal Organism	References
Alternaria leaf spot	<i>Alternaria alternata</i>	Pandey and Nigam, 1985
Myrothecium leaf spot	<i>Myrothecium roridum</i>	Maharshi 1986
Soft rot	<i>Pseudomonas cichorii</i>	Sattar et al. 1981
Wilt	<i>Fusarium solani</i>	Thakur et al. 1974
Xanthomonas blight	<i>Xanthomonas compestris</i> pv. <i>turf</i>	Sattar and Alam, 1996

cm. deep. Pest is active during May to November (Newton 1934). Adults hibernating in the grass bordering the field during the winter. Generally, only one generation completed per year

Nature of Damage :Adults feed on leaves, larvae mine leaf-stalks (petioles) and leaf-blades. The central pith of the main stem is also invaded and some larvae also infest the tap root. In severe infestation the stem is completely hollowed out

Management

- Crop rotation
- Early harvesting in June month.

Fungal diseases (Table 11)

Leaf spot :This is an important disease and first time reported in the state of Jammu and Kashmir caused by *Ascochyta kashmiriana* (Ganguli and Pandotra, 1962) Leaf spot also caused by *Alternaria alternata* on Egyptian henbane (Kumar *et al.* 1984). This disease appeared during cultivation in the month of January, February and March.

Symptoms:On infected leaves several irregular brownish spots occurs and later on these spots enlarges in size and become big spots. The infected leaves generally defoliated prematurely and consist of spores. These spores are the major source of perpetuation of the disease.

Management:Spraying of Dithane M-45 (0.3%) or Kavach (0.2%) at an interval of 15 days minimizes the disease severity and incidence.

Viral Diseases: Diseases caused by different isolates of viruses on henbane belong very diverse genera. Among the first viruses reported to cause diseases on henbane are *Hyoscamus virus II*, *Hyoscamus virus III* and *Hyoscamus virus I*, (Hamilton, 1932). Later number of diseases had been reported to be caused by viruses belonging to genus; *Tobamovirus*, *Cucumovirus*, *Tospovirus*, *Potexvirus*, *Tobravirus* etc. (Table 11).

A cucumber mosaic virus isolate was found to be associated with green mosaic disease of *H. niger* L. Based on host range, symptomatology, mode of transmission, biological properties, electron microscopy and serology, it was found to be an isolate of cucumber mosaic virus. The disease still is prevalent. The virus was transmitted by aphid, *Myzus persicae* Sulz, the virus was isometric, 27 nm in diameter and it reacted with the antiserum of CMV9ATCC, PVAS-30)

Tomato spotted wilt virus (TSWV) belong to genus *Tospoviruses*. TSWV had been found to cause a mild mosaic disease in *Hyoscyamus muticus* L (Zaim, 1999). It had diagnostic host as *Vigna unguiculata* CV.152 which produced Chlorotic spots on inoculated leaves. Biological properties and electron microscopy of ultrathin sections of infected *H. muticus* leaves exhibited the typical Tospoviruses surrounded by endoplasmic reticulum inside cytoplasm. Purified virus preparation spherical virus particles about 87 nm in diameter.

A cucumber mosaic virus isolate was found to be associated with mottle crinkle and severe mosaic disease of Egyptian henbane (*Hyoscyamus muticus* L.). The virus has been characterized as an Indian isolate of cucumber mosaic virus (CMV) based on non-persistent transmission by aphid, presence of 28-nm isometric particles, capsid protein of 26 K and single-stranded tripartite RNA genome with a subgenomic RNA (RNA 4). There was no evidence of satellite RNA genome. The isolate showed a strong serological relationship with S

and A strains of CMV (CMV-S and CMV-A) in double diffusion test. A band of the 26 K capsid protein was also detected by Western blot analysis using antibodies specific to CMV-S.

Nematodes of Henbane:

Large number of plant parasitic nematodes viz. *Tylenchorhynchus vulgaris*, *Pratylenchus thornei*, *Hoplolaimus indicus*, *Longidorus pisi* and *Meloidogyne* sp. have been reported to be associated with different species of henbane but the nematode which has been reported to cause serious damage to the crop is root-knot nematode (*Meloidogyne* spp.). Root-knot nematode causes poor growth of the seedling establishment and leads to significant decrease of the herbage yield of the plant (Figure 24).

Root-knot nematodes of henbane: The different species of *Hyoscyamus* are found to be heavily infested with root-knot nematode, *M. incognita* & *M. javanica*, which cause significant damage to the crop (Pandey, 1990).

Symptoms of damage: In the field the root-knot infested plants of *Hyoscyamus muticus*, *H. niger* & *H. albus* show chlorosis, stunted plant growth, a patchy appearance fewer smaller leaves and flowers. The roots of infested plants depict severe galling of various degrees. Experiments carried out in CIMAP indicate that even 3-4 larvae/g soil cause significant damage to the crop (Haseeb and Pandey 1989).

Control measures:

In the control programme of root-knot nematode the major attention has been paid to protect the young seedlings. Reduction of nematode population in soil prior to sowing is very important to avoid the poor plant growth. Pre-sowing measures to suppress root-knot nematode population include rotation with non host crops, fallowing, green manuring, organic amendments and application of some nematode antagonistic organisms.

Growing of non host crop such as *Cymbopogon flexuosus*, *C. winterianus*, and *C.*

martinii can reduce the incidence of damage caused by root-knot nematode. Marigold (*Tagetes spp*) are one of the most highly studied crops for its ability to suppress nematode population with antagonistic phytochemical in root exudates i.e. polythienyls. Therefore prior to sowing henbanes, it is very important to grow *Tagetes minuta* at the distance of 30 X 30 cm which drastically reduced the phytonematode population (Pandey 2011 Unpublished). Removal of weed host of root-knot nematode from the field is an integral part to avoid the infestation of root-knot nematode as the susceptible weed hosts may serve as a reservoir for root-knot nematode inoculum. Avoiding the general practice of irrigation of noninfested field with water from infested field, will help to restrict *Meloidogyne* sp. population spreading from one field to another.

Certain crops resistant to root-knot nematodes could be used in rotation with henbanes to reduce the root-knot nematode population. However population explosion renders the practice of fallowing not possible due to increased economic pressure on agricultural land. Experiment has been carried out at Central Institute of Medicinal and Aromatic Plants, Lucknow and its resource canter to screen large number of tetraploid henbane species but none was found to be resistant to root-knot nematodes. Chemicals such as carbofuran (@2kg a.i./ha soil) and

monocrotophos (@0.1% SS) have been used to prevent root-knot nematode infestation to henbane. Monocrotophos can be used for seed soaking whereas carbofuran is preferred prior to sowing the seeds of the crop. It helps up to some extent to eliminate the root knot nematode infestation. The harvesting period of this winter crop is only four months, sowing daicha as green manuring crop can be used which decreases nematode population below the economic damage levels.

Biological control: Some ecofriendly approaches have also been made to manage this important pest on these plants. In one of the experiment Pandey *et al.* (1999) observed that the plants with inoculum of different mycorrhizal fungi showed better growth in comparison to untreated - *M. incognita* inoculated and untreated-uninoculated one.

The best results were obtained with *Glomus aggregatum* for increasing growth/ biomass of plants and reduction in root-knot nematode population. For example in *Hyoscyamus niger* the nematode population was observed maximum in nematode alone inoculated plants than plants inoculated in combination of nematode and bio-inoculants resulting significant reduction of *M. incognita* (Mi) population especially in combined treatment followed by *Pseudomonas fluorescens* (Pf) or *G. aggregatum* (Ga) inoculated treatments. Thus, the degree of losses in biomass



Figure 24. Healthy & root-knot infested plant and roots of Egyptian henbane (*H. muticus*)

production caused by Mi in combination with the bio inoculants were significantly reduced over the losses observed with Mi alone. Among the four combined inoculations, a combination of all the four with Mi appeared to be the best followed by GA or Pf. Root colonization and spore population of VAM-fungi was observed 16 weeks after inoculation. Maximum spore population and percent root colonization was observed in combined treatment, which was followed by GA, Gm and Gf. It is worth mentioning here that root colonization was found directly proportional to spore population and biomass yield from the crop. Experimental findings indicated that application of bio-inoculants have not only enhanced the total biomass yield of *H. niger* but it have also significantly decreased the multiplication of nematode, however, a significantly higher reduction was recorded in the treatment where all bio-inoculants are combined . This may be attributed to the fact that these bio-agents may be secreting potent chemicals which are either non favorable for multiplication of Mi or inducing tolerance in the plant against the attack of root knot nematodes. Nematode reproduction was higher in plants inoculated with nematode alone than in plants with combined inoculation of nematodes with bio-inoculants. Thus, experimental evidences indicated that mixed inoculation of rhizobacterium with VAM fungi could be considered as biological management instead of nematicides for reducing the deleterious effect of root knot disease in black henbane. Similar results were also obtained with different bio-agents in *H. muticus* plants (Pandey 1997c, Pandey *et al.* 2000c).

Essential oils were also used in another experiment to manage root-knot nematode population in *H. niger* (Pandey *et al.* 2000a). It was observed that oils of *Cymbopogon martinii*, *C. wintrianus*, *Ocimum basilicum* and *Mentha arvensis* were quite effective in reducing *M. incognita* population and improving the growth of plant; the oil of *C. martinii* (@2ml/pot) being most effective.

Summary

The cultivation of medicinal and aromatic plants has considerably increased in tropical and sub tropical countries of the world to meet the requirement of pharmaceutical and perfumery industries. The disease caused by variety of pest and pathogen is a major limiting factor for its production. Although chemical control of pest and pathogen has proved effective, but it can not be recommended to the farmers because of cost, harmful effect on human and animal health and its percolation in ground water resources. Other alternatives are available in medicinal plants like resistant germplasms, useful organic materials, effective biocontrol agents and other cultural/physical methods of disease management. Such resistant and tolerant germplasms could be exploited in future plant breeding programmes for developing resistant/ tolerant genotypes against major pest and pathogens. Various organic materials are available which has been proven useful to decrease disease incidence and enhanced medicinal and aromatic plant yield could be used in ongoing programmes for better healthy plants. Large number of bio agents is available which could be used on large scale to save medicinal and aromatic plants against various pest and pathogens. Based on the contribution added to the science of mycology, virology and nematology in relation to medicinal and aromatic plants, CIMAP plant pathology, virology and nematology labs have been placed in national and international records. However much attention also is needed to study and develop some new strategy to manage major diseases in ecofriendly way, which should be cost effective and environmentally friendly

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