Integration of Soil Solarization, Arbuscular Mycorrhizal Fungi, Trichoderma viride, Azotobacter chroococcum and Soil Amendments for the Management of Carnation (Dianthus caryophyllus L.) Wilt (Fusarium oxysporum f.sp. dianthi (Prill and Del.) Snyd. and Hans.)

Savita Sharma*, Harender Raj and Neeraj Sharma

Dr. Y S Parmar University of Horticulture and Forestry, Nauni, Himachal Pradesh 173032, India

*Corresponding author

Studies were conducted on Carnation (Dianthus caryophyllus L.) to find out the effect of integrated inoculation of potent native isolates of Arbuscular mycorrhizal fungi (AM fungi) and Azotobacter chroococcum with other approaches of management like cultural and biological methods in solarized soil on the incidence of wilt caused by Fusarium oxysporum f.sp. dianthi (Prill. and Del.). Initially, organic amendments, botanicals and bio-control agents were evaluated against wilt to find out the best treatments. Among amendments, neem cake was found most effective with 75.0 per cent reduction in the wilt incidence. Different fungicides, botanicals, bio-pesticides and bio-control agents were also evaluated against wilt by dip treatment of unrooted carnation cuttings. Bavistin among fungicides, Neemazal among botanicals and Trichoderma viride among bio-control agents were found effective with 100.0, 71.1 and 93.6 per cent reduction in the wilt incidence. Based on the best individual treatments, fourteen treatment combinations were evaluated in solarized and unsolarized plots for their efficacy against the disease. Among different treatments, root dip of cuttings in Bavistin (0.1%), soil amendment with Neemcake (1kg/m²), root inoculation with culture of AM fungi and A. chroococcum (5g culture/plant) and soil application of T. viride formulation (10g/m²) in solarized soil was found most effective with 97.1 per cent reduction in the wilt incidence. This treatment combination also resulted in maximum increase of 50.97, 100.4, 39.2, 57.3, per cent in plant height, number of flowers per plant, flower size and length of flowering stem, respectively in comparison to control.

Keywords: Fusarium wilt, Carnation, Soil solarization, Azotobacter chroococcum, Arbuscular mycorrhizal fungi, Integrated disease management

Abstract

Introduction

Vascular wilt caused by Fusarium oxysporum f.sp. dianthi is most prevalent disease in carnation and upto 79 per cent incidence has been recorded in different parts of the Himachal Pradesh (Chandel and Katoch, 2001). Soil-borne pathogens are difficult to control due to repeated cultivation of the crop in the same piece of the land. Use of chemicals in the management of soil-borne disease results in high cost of production and also has drastic adverse effect on soil microflora (Aktar et al., 2009). Thus, there is
an urgent need for development of integrated disease management strategy by evaluation of other physical, biological and cultural methods effective against the wilt pathogen. Soil solarization (SS) is one of the important and cost-effective methods for the management of soil-borne pathogens in different crops in different regions (Katan 1981). Bio-control agents and soil amendments have also been reported effective against the soil-borne diseases (Lodha and Israel 2005, Karimi et al., 2007). Soil is rich in many beneficial microorganisms like VA mycorrhizal fungi and A. chroococcum which are beneficial to the plants in enhancing plant growth and productivity, and these organisms also help in reducing incidence of different soil-borne pathogens (Smith, 2002, Dehne, 1982 and Brown, 1974). Soil solarization has been found more effective against soil-borne pathogens when integrated with biological control agents, soil amendments and chemical treatment (Gamliel and Stapleton 1993, Raj and Sharma 2009). Hence, the present investigation was undertaken to evaluate the integrated efficacy of SS, botanicals, bio-control agents, soil amendments and chemicals for management of the disease.

Materials and Methods

Soil amendments

Soil amendments were evaluated to find their effect on the incidence of Fusarium wilt of carnation and to know their effect on important plant growth characteristics and quality parameters of the flowers. In this trial, seven soil amendments viz., neem (Azadirachta indica L.) cake, vermicompost, darek (Melia azedarach L.) seed meal, karu (Roylea elegans Wall.) leaves, cauliflower (Brassica oleracea L. var. botrytis L.) leaves and banna (Vitex negundo) leaves were used at the rate of 100 g/pot which contained 5 kg of soil. In addition, neem granules (Azadirachtin 0.15 % (E.I.D. Parry (India) Ltd.) were also used and were applied at the rate of 10 g/pot. These amendments were mixed thoroughly in the upper 15cm soil layer. Soil was then irrigated to saturation level and left for the decomposition for two weeks before planting the carnation cuttings.

Root dip treatments

Unrooted carnation cuttings of variety ‘Sunrise’ were dipped in different treatments of fungicides, botanicals and bio-pesticides for 30 minutes followed by a quick dip with NAA at 500 ppm before planting them into rooting media containing sand and soil in the ratio of 1: 1. Fungicides viz., carbendazim (Bavistin 50% WP) (0.1%), hexaconazole (Contaf 5% EC) (0.05%), difenoconazole (Score 25% EC) (0.025%), mancozeb (Dithane M-45 75% WP) (0.25%), iprodione 25% + carbendazim 25% WP (Quintal) (0.2%), carbendazim 12% + mancozeb 63% WP (Saaf) (0.2%), captaf (captan 50% WP) (0.2%), pyraclostrobin 5% + metiram 55% WG (Cabrio Top) (0.2%) and myclobutanil (Sythane 10% WP) (0.05%) were taken. In botanicals and bio-pesticides plants like darek (Melia azedarach L.) (1%), karu (Roylea elegans Wall.) (1%), dudhli (Cryptolepsis buchanani Roem. & Schult.) (1%), tulsi (Ocimum sanctum L.) (1%) shambri (Artemisia roxburghiana) (1%), safeda (Eucalyptus globulus) (1%), gharit kumari (Aloe vera) (1%), commercial formulation of neem (Neemazal 1.0% EC) (1%) and also vermiwash (1%) were taken. In fungal antagonists like Trichoderma viride (1%) and T. harzianum (1%) were taken where one percent formulation was made by taking one gm of commercially available formulation made in talc powder and then dissolving it in 100 ml water. In bacterial antagonists, Bacillus subtilis (1%), Brevibacillus brevis (1%), Azotobacter chroococcum (1%) and Pseudomonas fluorescens (1%) were taken.
where one per cent formulation was made by dissolving 1ml of Nutrient Agar broth culture of bacteria in 100 ml water. Cuttings were inserted in the rooting media up to two nodes and then kept in the mist chamber. Data on disease incidence, root length and plant height were recorded after 30 days.

**Isolation, identification and mass multiplication of native potent isolates of AM Fungi and *A. chroococcum***

Soil samples were collected from different carnation growing areas of the State to isolate potent isolates of AM fungi. Seven potent isolates, viz *Glomus mosseae*, *G. fasciculatum*, *G. macrocarpum*, *G. constrictum*, *Acaulospora bireticulata*, *Gigaspora* sp., *Entrophospora* sp. were selected on the basis of occurrence and frequency of distribution in the carnation growing areas. The consortium of these seven potent isolated isolates of AM fungi was made and named as AMUHF. The AMF spores were isolated by wet sieving and decanting method of Gerdemann and Nicolsan (1963) and identified to the genus level under tri-nocular biological microscope (Leica DMLB) attached with a digital camera. Spores were identified by different synoptic keys (Morton 1988). These isolates were multiplied on green gram (*Vigna radiata* L. Wilczek) in sterilized soil in earthen pots for 3 months. These plants were uprooted after 3 months and their roots were chopped into pieces to develop mass culture of consortium of AM fungi for inoculation into soil. The inoculum of different isolates used in the field experiments contained spores of the isolate, pieces of infected chopped roots and mycelium in the pot culture soil. Isolate of *A. chroococcum* was selected from the rhizosphere soil of carnation by serial dilution technique and it was named as AZUHF. 10g soil from the rhizospheric soil of carnation was drawn and serially diluted aseptically to $10^{-3}, 10^{-4}, 10^{-5}$ and $10^{-6}$ dilutions and out of this 1ml of suspension was spreaded on Jenson’s medium (Subba Rao, 1986). Culture carrier of each isolate was prepared in 10 % jaggery slurry added with gum to stick. This slurry of the culture was prepared to apply the culture to the roots.

**Soil solarization**

Soil solarization was done for 40 days during 1st May to 9th June 2011 with thin transparent polyethylene sheet (25 µm thick). Beds (1 x 1m) were irrigated to saturation level and then covered with thin transparent polyethylene sheet. The sheets were removed after 40 days of solarization. In the second set, beds were not covered with any sheet and served as control for comparison. During the period of solarization, soil temperature was recorded every day for 40 days at 2 pm in both solarized as well as unsolarized beds with dial type digital thermometer at 5 and 15 cm soil depths.

**Integrated disease management**

Treatments that proved effective under *in vitro* and polyhouse experiments were then integrated with soil solarization to know and compare their individual and combined effect on the incidence of carnation wilt. The experiment was laid out in the polyhouse during the year 2011, which comprised of effective treatments of soil amendments, root dip/treatment of cuttings with fungicides /botanicals/bio-pesticides/bio-control agents and effective combination of AM fungi, *Azotobacter chroococcum* and *T. viride* in different combinations in solarized and unsolarized soil. Talc based formulation of *T. viride* was applied before planting @ 1% i.e. by mixing 10 g of the talc powder formulation (6×106 cfu/g) in 1kg well rotten farm yard manure per bed. Neem cake was applied at the rate of one kg/m² both in solarized and
unsolarized beds (lm x lm) and were mixed thoroughly in the upper 15 cm soil layer irrigated to saturation level and left for the decomposition for two weeks. The roots of the carnation cuttings were dipped for 15 minutes in culture slurry of the A. chroococcum so that bacteria could adhere on the root surface. Among fungicides, bavistin (0.1%) was used as root dip treatment of the cuttings for 30 minutes and among botanicals/bio-pesticides, Neemazal was used at 20 per cent concentration as root dip treatment of cuttings for 30 minutes before planting. The carnation cuttings were planted in solarized and unsolarized beds in planting holes which were added with 5g inoculum of AMUHF before planting. Recommended dosages of chemicals fertilizers used in the polyhouse experiments were urea (46% N), single super phosphate (16% P2O5, 19% Ca, 12% Sulphur) and muriate of potash (60% K2O).

IDM treatments

Fourteen treatments, viz T1, root dip of cuttings in Bavistin @ 0.1 %; T2, soil application of T. viride @ 10g/1 kg of FYM; T3, root dip of cuttings in Neemajal @ 20 %; T4, soil amendment with Neem cake @ 1 kg/m²; T5, (T4 + T2); T6, (T4 + T1); T7, (T4 + T3); T8, root inoculation of cuttings with AMUHF @ 5g/plant + AZUHF @ 5g/plant + Soil application of T. viride @10g/1kg of FYM/m³; T9, (T4 + T8); T10, (T1 + T8); T11, (T3 + T8); T12, (T1 + T4 + T8); T13, (T3 + T4 + T8) and T14 Control (Unamended and unsolarized) were applied in the field in the poly-house each comprising of three replications in Randomized Block Design. Carnation cuttings of variety ‘Master’ were planted at a distance of 20 x 20 cm in 1m x 1m bed with 25 cuttings per bed. Per cent disease incidence was calculated during the growing period in each bed. Data pertaining to plant growth and quality parameters viz., plant height (cm), number of days taken for first flowering (days), number of flowers per plant, length of flowering stem (cm) and flower size (cm) were recorded by selecting 5 plants per replication in each treatment.

Statistical analysis

The data recorded from pots and mist chamber experiments were analyzed as per the procedure of Completely Randomized Design (CRD) and data of field experiments were statistically analyzed using Randomized Block Design (RBD) as described by Gomez and Gomez (1984). Least significance difference at 5% level was used for testing significant differences. The data on per cent disease incidence were arc sine transformed (in parentheses) then subjected to statistical analysis.

Results and Discussion

Effect of soil amendments

Soil amendments were found effective in reducing the incidence of the wilt. However, neem cake was found most effective among all the treatments which resulted in 16.67 per cent reduction in the incidence of wilt in comparison to 66.67 per cent in control. This treatment also resulted in maximum increase (44.7 %) in plant height and took 8.9 per cent less days to 1st flowering. Soil amendments have been reported to enhance the activity of the soil microflora which are potentially competitive or antagonistic against several soil-borne pathogens by different modes of actions including production of various biochemical substances during decomposition (Hortink and Fahy 1986). Negi (2009) reported that soil amendment with neem cake was found effective with 35.4 per cent reduction in incidence of wilt of carnation. Soil amendment with neem cake has also been reported most effective with 71.0 per
cent reduction in incidence of *Fusarium* wilt (*F. oxysporum* f.sp. *dianthi*) of carnation (Chandel, 2011). The mechanism of disease control for high nitrogen containing amendments like oil cakes is the generation of ammonia and nitrous acid following degradation of the amendments by microorganisms which is lethal to pathogens (Lazarovits et al., 2001). Application of nitrogen rich soil amendments (oil cakes) reduced soil-borne diseases by releasing allelochemicals (Bailey and Lazarovits, 2003) (Table 1).

**Effect of root dip treatments**

Among fungicides, dip of carnation cuttings in Bavistin and Quintal were found most effective with complete reduction of incidence of wilt. However, treatment with Bavistin also resulted in maximum increase of 66.0 and 440.5 per cent in average plant height and root length followed by Quintal with 54.1 and 372.7 per cent, respectively in comparison to control (Table 2). Kishore and Kulkarni (2008) also reported effectiveness of carbendazim against *Fusarium* wilt of carnation. Drenching of rooting media of carnation with carbendazim @ 0.2% has also been reported effective in reducing the incidence of *Fusarium* wilt of carnation (Sharma 2000).

Among bio-control agents, *T. viride* has been found most effective with 2.3 per cent incidence of wilt followed by *A. chroocococum* with 6.3 per cent disease incidence. Treatment of the cuttings with *T. viride* resulted in maximum increase of 73.1 and 586.4 per cent in average plant height and root length followed by *A. chroocococum* with maximum average increase of 63.2 and 575.7 per cent, respectively in comparison to control (Table 3). Chandel (2011) reported that root dip of carnation cuttings in *T. viride* is effective with 68.55 per cent reduction in the incidence of wilt in comparison to control. *T. viride* and *T. harzianum* applied during rooting of carnation cuttings strongly promoted growth of plants and gave good control of *F. oxysporum* f.sp. *dianthi* (Manka et al., 1997; Weber et al., 1998). Martinez and Pinzon (1999) also reported that application of *Trichoderma* spp. to unrooted carnation cuttings at the time of application of rooting hormone and one more application to the soil immediately before planting resulted in reduction in incidence of *Fusarium* wilt.

Among different treatments of the botanicals and bio-pesticides, dip of cuttings in Neemajal was found most effective with 11.9 per cent incidence of the wilt. Further, treatment of unrooted carnation cuttings with Neemajal resulted in maximum increase of 597.6 and 51.9 per cent in average root length and plant height in comparison to control. Chandel and Tomar (2008) reported that dip of carnation cuttings in neem formulation (Ahook) is most effective with 89.6 per cent reduction in the incidence of wilt (*F. oxysporum* f.sp. *dianthi*) in carnation followed by Neemajal in comparison to control. The mechanism behind the disease control may be the Azadirachtin from neem which act as a chitin inhibitor and cause lysis of cell walls of resting pathogenic spores present in sick soil and stimulation of fungal antagonist in soil may have an indirect effect (Bhattacharya and Pramanik, 1998) (Table 4).

**Effect of soil solarization**

Soil solarization with transparent polyethylene sheet resulted in average increase of 8.3 °C in the soil temperature at 5 cm soil depth with average maximum soil temperature of 41.0°C in the solarized soil in comparison to unsolarized beds (Table 5). However, increase in average maximum soil temperature at 15cm soil depth was 6.0°C. In general, transparent polythenes mulch (25µm
thick) has been reported to be effective in increasing the average maximum soil temperature (Katan, 1981). Melero-Vara et al., (2005) also reported increase of 5-7 °C in average maximum temperature in the poly-house in an experiment on use of soil solarization for the management of *Fusarium* wilt of carnation.

**Effect of IDM on disease incidence and growth characteristics**

Integration of different effective treatments had enhanced efficacy than the individual treatments in the management of the wilt and in the improvement of plant growth and flower quality characteristics in carnation. All the treatment combinations were found effective and these treatments were more effective under solarized conditions (Table 6 and 7). These treatments reduced the wilt incidence ranging from 54.2 to 97.2 per cent under solarized plots in comparison to 51.5 to 80.0 per cent under unsolarized plots. Treatment combination T12 (root dip of carnation cuttings in Bavistin @ 0.1% + Neemcake @1kg/m² as soil amendment + AMUHF @ 5g/plant as soil application + AZUHF @ 5g/plant as root inoculation of cuttings and *T. viride* @ 10g/1kg of FYM/m² as soil application) in solarized plots was found most effective with wilt incidence of 1.3 per cent in comparison to 46.6 per cent in unsolarized control (Table 6). Different components of the Treatment T12 have a distinctive effect in enhancing the efficacy of the treatment. All the treatment combinations were found statistically superior in solarized soil with 2.7 to 17.2 per cent more control in the incidence of the wilt. Soil solarization has been reported effective for the management of wilt of carnation (Melero-Vara et al., 2005). Reduction in disease incidence due to the application of organic amendments with solarization has been reported in *Fusarium* and *Phytophthora capsici* infestation in pepper (Martínez et al., 2011; Núñez-Zofio et al., 2011) and was at least partially attributed to the production of NH₃ and an increase in soil microbial activity, which can help control soil-borne pathogens through competition, antibiosis, parasitism/predation, etc. (Núñez-Zofio et al., 2011). Soil solarization in combination with soil amendments, crucifer residues and microbial pesticides like *Trichoderma* spp., *Gliocladium* sp., *Pseudomonas* sp. has also been reported to be effective in strawberry, gladiolus, vegetables and other crops against different soil-borne diseases (Porras et al., 2009, Raj and Upmanyu, 2013).

In Treatment combination T12, root inoculation with culture of AM fungi and *A. chroococcum* have a significant effect in the management of the wilt. If we compare treatments T6 and T12, it is evident that addition of T8 with T6 resulted in 11.4 per cent more reduction in the incidence of the wilt. There are number of reports in the literature which explain the role of AM fungi, *A. chroococcum* and *Trichoderma* spp. in the management of different soil-borne diseases. Inoculation of carnation cuttings with *Glomus intraradices* has been reported to reduce *Fusarium* wilt (*F. oxysporum* f.sp. *dianthi*) and the reduction in disease incidence was associated with reduction of number of propagules of the wilt pathogen. Reduction in the wilt incidence has been attributed either to the induction of disease resistance mechanism by the mycorrhizal fungus, or by direct/indirect interaction between VAM fungus and *F. oxysporum* f.sp. *dianthi* inoculum in the soil (St-Arnuad et al., 1997). Application of VA-mycorrhizal fungi *Gigaspora margarita* in pea against *Fusarium* wilt (*Fusarium oxysporum* f.sp. *pisi*) resulted in minimum incidence (10.8%) of wilt in comparison to 45.8 per cent in control (Verma and Dohroo 2005). VA-mycorrhizal fungi exert number of factors, like lignifications of
mycorrhizal roots, increased respiration, increased production of arginine and isoflavonoids, better Phosphorus nutrition, changes in the antioxidant enzymes, phenolic compounds and pathogen related-proteins which are reported to contribute in imparting resistance against soil-borne pathogens (Dehne et al., 1978; Dehne, 1982; St-Arnaud et al., 1994; Morandi, 1996; Khalil, 2007).

Efficacy of *Trichoderma* spp. in different treatment combinations has been reported against different soil-borne diseases. Dipping of corms of gladiolus in carbenazim (0.05 %) for 30 mintues along with soil application of neem cake (100 g per row) and *T. viride* (2.5 % w/w) has been resulted in 74.51 per cent reduction in disease incidence of *Fusarium* yellows (*Fusarium oxysporum* f.sp. *gladioli*) in gladiolus in comparison to control (Sharma et al., 2005). Inoculation of four AMF (*Glomus intraradices*, *Glomus mosseae*, *Glomus claroideum* and *Glomus constrictum*) and *Trichoderma* sp. in the seedlings nurseries has been reported to reduced the incidence of *Fusarium* wilt (*Fusarium oxysporum* f.sp. *melonis*) in melon seedlings (Martinez-Medina et al., 2009). Integration of SS, *Glomus fasciculatum* isolate of Vamcorrhiza and native isolate of *A. chroococcum* was found most effective with no incidence of white root rot of apple caused by *Dematophora necatrix* in comparison to 33.6-35.4 per cent in control (Raj and Sharma, 2009). Tomato seedlings inoculated with *T. harzianum* and arbuscular mycorrhizal fungi (AMF) has been reported to have reduced disease severity of wilt caused by *Fusarium oxysporum* f.sp. *lycopersici* (Mwangi et al., 2011). Integration of SS along with application of neem cake (30g/ha) and *Azotobacter* (40kg/ha) has been resulted in 44.3 per cent reduction in the disease incidence of wilt (*Fusarium oxysporum* f. sp. *cuminii*) of cumin in comparison to control (Bijarniya and Lal, 2009). *A. chroococcum* has been reported to have antagonistic effect against *Fusarium oxysporum* by degradation and digestion of cell wall components, empty cell (halo) formation, shrinking and lysis of fungal mycelia along with significant degeneration of conidia (Maheshwari et al., 2012). Similarly, many researchers have reported inhibitory effects of *A. chroococcum* on different soil-borne diseases (Ebtehag et al., 2009; Umesh and Mane, 2010).

**Effect of IDM on quality parameters**

Different treatment combinations also resulted in improvement of important plant growth and quality characteristics of carnation both in solarized and unsolarized plots. Treatment combination T12 in solarized plots was found most effective with an increase of 50.97, 100.4, 39.2 and 57.3 per cent in plant height, number of flowers per plant, flower size and length of flowering stem, respectively and also recorded 15.22 days to 1st flowering, respectively in comparison to unsolarized and unamended control (Table 2 and 3). Different components of Treatment combination T12 have been reported to have positive effect on different growth and quality characteristics of different plants raised in soil infected by different soil-borne pathogens. SS has been reported to support higher growth and yield in different crops including nursery of fruits and vegetables (Patel, 2001; Raj, 2004). The mechanism for explaining increased growth responses and yield in plants has been attributed to chemical factors (like release of nutrients and other growth factors, nullification of toxins) and biological factors (elimination of minor or unknown pathogens) and stimulation of beneficial micro-organisms (Stevens et al., 2003). Gawande et al., (2001) reported that SS and *Trichoderma* sp. resulted in recording the least number of days required for flower bud initiation and first flower bud opening per plant in chrysanthemum. Inoculation of
mango seedlings with *Glomus fasciculatum* and *A. chroococcum* in solarized soil has been reported to increase seedlings height, diameter, leaf area, total root length, leaf N, P, K and Zn content in comparison to control (Sharma *et al.*, 2011). The direct mechanisms of increase in root development and plant growth by *Azotobacter* has been attributed to the secretion of vitamins and amino acids; production of siderophores and auxins (Akbari *et al.*, 2007). Similarly, conjoint inoculation of plants with *Azotobacter* and Va-mycorrhizae has been reported to increase in the rhizosphere populations of bacteria and actinomycetes and resulted in synergistic growth enhancement of the host plant (Bagyaraj and Menge, 1978). Thus, root dip of cuttings in Bavistin (0.1%), soil amendment with Neemcake (1kg/m²), root inoculation with culture of AM fungi and *A. chroococcum* (5g culture/ plant) and soil application of *T. viride* formulation (10g/m²) in solarized soil is effective with 97.1 per cent reduction in the wilt incidence.

**Table.1** Effect of different organic amendments on the incidence of *Fusarium* wilt and important plant growth characters

| Treatments                  | (Rate of application in g/ 5kg of pot soil) | Disease incidence (%) | Plant height (cm) | Number of days taken for 1st flowering |
|-----------------------------|---------------------------------------------|-----------------------|-------------------|---------------------------------------|
| Neemcake                    | 100g                                        | 16.67 (19.99)         | 75.21             | 128.67                                |
| Cauliflower leaves          | 100g                                        | 33.33 (34.99)         | 70.97             | 131.00                                |
| Neem granules               | 5g                                          | 25.00 (24.99)         | 73.15             | 129.33                                |
| *Melia azedarach* (S)       | 100g                                        | 33.33 (34.99)         | 68.44             | 133.67                                |
| *Roylea elegans* (L)        | 100g                                        | 41.67 (39.98)         | 66.20             | 133.00                                |
| *Vitex negundo* (L)         | 100g                                        | 58.33 (49.98)         | 62.34             | 134.67                                |
| Vermicompost                | 100g                                        | 50.00 (44.98)         | 64.57             | 135.33                                |
| Control                     | _                                           | 66.67 (59.97)         | 51.95             | 141.33                                |
| CD<sub>0.05</sub>           | NS                                          | 5.48                  | 6.7               |

S, Seed meal; L, Leaves  
* Figures in parentheses are arc sine transformed values
Table 2 Effect of dip treatment of unrooted carnation cuttings in fungicides on the incidence of *Fusarium* wilt and important plant growth parameters

| Fungicides | Conc. (%) | Diseases incidence (%) | Mean Plant height (cm) | Mean Root length (cm) | Mean |
|------------|-----------|-------------------------|------------------------|-----------------------|------|
|            | 2010      | 2011                    | 2010                  | 2011                  |      |
| Bavistin   | 0.1       | 0 (0)                   | 0 (0)                 | 0 (0)                 |      |
| Dithane M-45 | 0.25   | 17.45 (4.17)            | 15.87 (3.97)          | 16.66 (4.07)          | 13.87 |
| Score      | 0.025     | 11.11 (3.32)            | 11.11 (3.32)          | 11.11 (3.32)          | 13.06 |
| Contaf     | 0.05      | 15.87 (3.97)            | 20.63 (4.54)          | 18.25 (4.25)          | 12.98 |
| Quintal    | 0.2       | 0 (0)                   | 0 (0)                 | 0 (0)                 | 15.97 |
| Saaf       | 0.2       | 6.35 (2.48)             | 1.59 (0.73)           | 3.97 (1.61)           | 15.72 |
| Captaf     | 0.2       | 20.63 (4.54)            | 19.04 (4.34)          | 19.83 (4.44)          | 12.22 |
| Systhane   | 0.05      | 25.39 (5.03)            | 26.98 (5.19)          | 26.19 (5.11)          | 11.61 |
| Cabrio Top | 0.2       | 9.52 (3.09)             | 11.11 (3.32)          | 10.31 (3.2)           | 13.01 |
| Control    | –         | 30.15 (5.47)            | 25.39 (5.01)          | 27.77 (5.24)          | 10.28 |
| Mean       |           | 13.65 (3.21)            | 13.17 (3.04)          | 13.54 (3.18)          | 3.53  |
| **CD**     |           | **0.53**                |                       |                       | **0.77** |

*Figures in parentheses are arc sine transformed values*
**Table 4** Effect of dip treatment of unrooted carnation cuttings in botanicals and bio-pesticides on the incidence of *Fusarium* wilt and important plant growth parameters

| Treatment                  | Conc. (%) | Diseases incidence (%) | Mean | Plant height (cm) | Mean | Root length (cm) | Mean |
|---------------------------|-----------|------------------------|------|------------------|------|------------------|------|
|                           |           | 2010                   | 2011 | Mean             | 2010 | 2011             | Mean |
| Neemajal (Azadirachta indica) | 1.0       | 11.11 (19.37)          | 12.69 (20.78) | 11.90 (20.08) | 15.06 | 15.17 | 15.12 | 6.07 | 5.93 | 6.0 |
| Melia azedarach (S) (Darek) | 1.0       | 12.69 (20.78)          | 12.69 (20.78) | 12.69 (20.78) | 13.36 | 14.60 | 13.98 | 4.88 | 4.89 | 4.89 |
| Roylea elegans (L) (Karu)   | 1.0       | 15.87 (23.41)          | 14.28 (22.19) | 15.07 (22.8)  | 15.55 | 14.32 | 13.94 | 4.75 | 4.74 | 4.75 |
| Artemisia roxburghiana (L) (Shambri) | 1.0       | 19.04 (26.97)          | 19.04 (25.86) | 19.04 (25.86) | 13.16 | 13.10 | 13.13 | 3.32 | 2.75 | 3.04 |
| Cryptolepis buchanani (L) (Dudhli) | 1.0       | 19.04 (26.97)          | 19.04 (25.86) | 19.04 (25.86) | 13.15 | 12.99 | 13.07 | 2.92 | 2.34 | 2.63 |
| Ocimum sanctum (L) (Tulsi)  | 1.0       | 20.63 (24.64)          | 20.63 (26.97) | 20.63 (26.97) | 12.63 | 12.95 | 12.79 | 2.89 | 2.87 | 2.88 |
| Eucalyptus globulus (L) (Safeda) | 1.0       | 17.45 (26.97)          | 19.04 (25.86) | 18.25 (25.25) | 13.26 | 13.37 | 13.32 | 3.3  | 3.31 | 3.3 |
| Aloe vera (L) (Gharit kumari) | 1.0       | 20.63 (30.2)           | 22.21 (28.08) | 21.42 (27.53) | 12.97 | 13.01 | 12.99 | 2.85 | 2.6  | 2.73 |
| Vermiwash                  | 1.0       | 25.47 (38.98)          | 26.98 (31.26) | 26.23 (30.73) | 12.48 | 12.85 | 12.67 | 2.33 | 2.47 | 2.4 |
| Control                    | -         | 39.68 (38.98)          | 44.44 (41.77) | 42.06 (40.38) | 9.94  | 9.95  | 9.95  | 1.12 | 0.60 | 0.86 |
| Mean                       |           | 20.16 (26.3)           | 21.10 (26.94) | 12.96 | 13.23 | 3.44 | 3.25 |

CD<sub>0.05</sub> 2.52 0.49 0.44

S: Seed meal; L: Leaves
* Figures in parentheses are arc sine transformed values

**Table 5** Effect of soil solarization with transparent polyethylene sheet (25µm thick) on soil temperature in the polyhouse

| Treatment | Soil depth (cm) | Maximum soil temperature (ºC) during 1 May-9 June (2011) |
|-----------|----------------|----------------------------------------------------------|
|           |                | Average | Range          |
| Solarized with transparent polyethylene mulch (25 µm thick) | 5 | 41.0 | 35.00 - 45.00 |
|           | 15 | 35.8 | 29.00 - 39.00 |
| Unsolated | 5  | 32.7 | 27.00 - 37.00 |
|           | 15 | 29.8 | 26.00 - 34.00 |
Table 6 Effect of integration of soil solarization with effective root dip treatment of cuttings, soil amendment and combination of AM fungi, *Azotobacter chroococcum* and *Trichoderma viride* on the incidence of *Fusarium* wilt and important plant growth characters

| Treatment | Disease incidence (%) | Plant height (cm) | Number of days taken for 1st flowering |
|-----------|------------------------|-------------------|--------------------------------------|
|           | S          | US          | Mean      | S          | US          | Mean      | S          | US          | Mean      |
| **T1**    | Root dip of cuttings in Bavistin @ 0.1% | 6.67 (14.79) | 17.33 (24.56) | 12.00 (19.68) | 79.00 | 72.44 | 75.72 | 138.33 | 140.67 | 139.5 |
| **T2**    | Soil application of *T. viride* @ 10g/1 kg of FYM/m² | 9.33 (18.98) | 17.33 (24.56) | 13.33 (21.13) | 78.67 | 70.33 | 74.5 | 138.00 | 141.67 | 139.83 |
| **T3**    | Root dip of cuttings in Neemajal @ 20% | 10.67 (18.98) | 22.67 (28.4) | 16.67 (23.69) | 75.76 | 67.89 | 71.78 | 140.33 | 144.00 | 142.17 |
| **T4**    | Soil amendment with Neemcake @1 Kg/m² | 12.00 (20.26) | 21.33 (27.47) | 16.67 (23.87) | 77.78 | 69.56 | 73.67 | 138.67 | 143.00 | 140.83 |
| **T5**    | (T4 + T2) | 8.00 (16.42) | 14.67 (22.47) | 11.33 (19.44) | 82.66 | 76.78 | 79.72 | 136.67 | 137.00 | 136.83 |
| **T6**    | (T4 + T1) | 6.67 (14.79) | 16.00 (23.57) | 11.33 (19.18) | 81.55 | 74.00 | 77.78 | 135.00 | 137.33 | 136.17 |
| **T7**    | (T4 + T3) | 8.00 (16.42) | 16.00 (23.57) | 12.00 (20.00) | 80.83 | 73.67 | 77.25 | 137.33 | 139.00 | 138.17 |
| **T8**    | Root inoculation of cuttings with AMUHF @ 5g/plant + AZUHF @ 5g/plant + Soil application of *T. viride* @10g/1kg of FYM/m² | 5.33 (13.16) | 13.33 (21.36) | 9.33 (17.26) | 83.78 | 74.89 | 79.34 | 133.67 | 136.00 | 134.83 |
| **T9**    | (T4 + T8) | 2.67 (7.69) | 13.33 (21.36) | 8.00 (14.53) | 89.11 | 80.78 | 84.94 | 132.00 | 134.67 | 133.33 |
| **T10**   | (T1 + T8) | 2.67 (7.69) | 10.67 (18.98) | 6.67 (13.33) | 89 | 82.22 | 85.61 | 132.00 | 134.00 | 133.00 |
| **T11**   | (T3 + T8) | 4.00 (11.53) | 12.00 (20.26) | 8.00 (15.9) | 86.55 | 81.55 | 84.05 | 132.67 | 135.33 | 134.00 |
| **T12**   | (T1 + T4 + T8) | 1.33 (3.84) | 9.33 (17.7) | 5.33 (10.77) | 94.78 | 86.44 | 90.61 | 130.00 | 131.6 | 130.83 |
| **T13**   | (T3 + T4 + T8) | 2.67 (7.69) | 9.33 (17.7) | 6.00 (12.69) | 91.67 | 82.78 | 87.22 | 132.00 | 133.67 | 132.83 |
| **T14**   | (Control) | 21.33 (27.35) | 46.67 (43.06) | 34.00 (35.21) | 71.43 | 62.78 | 67.11 | 146.33 | 153.33 | 149.83 |

**CD<sub>0.05</sub>**

| Treatment | 3.43 |
| Treatment x Solarization | 4.85 | 4.12 | 6.67 |

*S, Solarized; US, Unsoarized

* Figures in parentheses are arc sine transformed

AMUHF, consortium made from different isolates of AM fungi; AZUHF, isolate of *Azotobacter chroococcum*
Table 7 Effect of integration of soil solarization with effective root dip treatment of cuttings, soil amendment and combination of AM fungi, *Azotobacter chroococcum* and *Trichoderma viride* on the important quality parameter of the flowers

| Treatment                                                                 | Number of flowers per plant | Flower size (cm) | Length of flowering stem (cm) |
|--------------------------------------------------------------------------|----------------------------|-----------------|-------------------------------|
|                                                                          | S  | US  | Mean | S  | US  | Mean | S  | US  | Mean |
| **T_1** (Root dip of cuttings in Bavistin @ 0.1%)                          | 3.44 | 3.11 | 3.28 | 6.19 | 6.08 | 6.14 | 71.89 | 65.44 | 68.67 |
| **T_2** (Soil application of T. viride @ 10g/1 kg of FYM/m²)              | 3.56 | 3.22 | 3.39 | 6.11 | 6.00 | 6.06 | 71.44 | 64.89 | 68.17 |
| **T_3** (Root dip of cuttings in Neemajal @ 20%)                          | 3.22 | 2.78 | 3.00 | 6.00 | 5.57 | 5.78 | 69.67 | 60.67 | 65.17 |
| **T_4** (Soil amendment with Neemcake @ 1 kg/m²)                         | 3.33 | 3.00 | 3.17 | 6.00 | 5.73 | 5.87 | 71   | 63.44 | 67.22 |
| **T_5** (T_4 + T_2)                                                      | 3.89 | 3.56 | 3.72 | 6.3  | 6.2  | 6.25 | 75.56 | 69.56 | 72.56 |
| **T_6** (T_4 + T_1)                                                      | 3.78 | 3.33 | 3.56 | 6.22 | 6.17 | 6.2  | 75.22 | 67.44 | 71.33 |
| **T_7** (T_4 + T_3)                                                      | 3.68 | 3.33 | 3.51 | 6.36 | 6.25 | 6.31 | 73.89 | 67.33 | 70.61 |
| **T_8** (Root inoculation of cuttings with AMUHF @ 5g/plant + AZUHF @ 5g/plant + Soil application of T. viride @ 10g/1 kg of FYM/m²) | 4.00 | 3.67 | 3.83 | 6.36 | 6.25 | 6.31 | 77.67 | 68.22 | 72.94 |
| **T_9** (T_4 + T_8)                                                      | 4.33 | 4.00 | 4.17 | 6.7  | 6.37 | 6.53 | 83   | 75.67 | 79.33 |
| **T_10** (T_1 + T_8)                                                    | 4.11 | 3.89 | 4.00 | 7.07 | 6.45 | 6.76 | 82.67 | 75.56 | 79.11 |
| **T_11** (T_1 + T_4)                                                    | 4.00 | 3.78 | 3.89 | 6.55 | 6.37 | 6.46 | 79.89 | 74.78 | 77.33 |
| **T_12** (T_1 + T_4 + T_8)                                              | 4.67 | 4.11 | 4.39 | 7.42 | 6.95 | 7.18 | 88.29 | 80.11 | 84.2  |
| **T_13** (T_3 + T_4 + T_8)                                              | 4.44 | 4.00 | 4.22 | 7.36 | 6.78 | 7.07 | 84.89 | 76.78 | 80.83 |
| **T_14** (Control)                                                      | 3.00 | 2.33 | 2.67 | 5.83 | 5.33 | 5.58 | 64.89 | 56.11 | 60.50 |

**CD**<sub>0.05</sub>

| Treatment | Number |       |
|-----------|--------|-------|
|           | 0.28   | 0.23  |

| Treatment x Solarization |       |
|--------------------------|-------|
|                          | 0.40  | 0.33  |

*S*, Solarized; *US*, Unsolarized
* Figures in parentheses are arc sine transformed values
AMUHF, consortium made from different isolates of AM fungi; AZUHF, isolate of *Azotobacter chroococcum*
This treatment combination is also effective in improving important plant growth and quality parameters with increase of 50.97, 100.4, 39.2, 57.3, per cent in plant height, number of flowers per plant, flower size and length of flowering stem, respectively in comparison to unsolarized and unamended control. This treatment combination also resulted in early flowering by 23.3 days in comparison to unsolarized control.

References

Akbari G A, Arab S M, Alikhani H A, Allahabadi I and Arzanesh M H. 2007. Isolation and selection of indigenous Azospirillum spp. and the IAA of superior strains effects on wheat roots. World Journal of Agriculture Science 3: 523-529.

Aktar M W, Sengupta D and Chowdhuri A. 2009. Impact of pesticides use in agriculture: their benefits and hazards. Interdisciplinary Toxicology 2: 1-12.

Bagyaraj D J and Menge J A. 1978. Interaction between a Va-mycorrhiza and Azotobacter and their effects on rhizosphere microflora and plant growth. New Phytopathologist 80: 567-573.

Bailey K L and Lazarovits G. 2003. Suppressing soil-borne diseases with residue management and organic amendments. Soil and Tillage Research 72: 169-180.

Bhattacharya I and Pramanik M.1998. Effect of different antagonist rhizobacteria and neem products on clubroot of crucifers. Indian Phytopathology 51:87-90.

Bijarniya D and Lal G. 2009. Integrated strategy to control wilt disease of cumin (Cuminum cyminum L.) caused by Fusarium oxysporum f. sp. cumini (Schlecht). Journal of Spices and Aromatic Crops 18: 13-18.

Brown M E. 1974. Seed and root bacterization. Annual Review of Phytopathology 12: 181-197.

Chandel S and Katoh R. 2001. Chemical control of Fusarium oxysporum f.sp. dianthi- an incitant of carnation wilt. Indian Journal of Microbiology 41: 135-137.

Chandel S and Tomar M. 2008. Effectiveness of bioagents and neem formulations against Fusarium wilt of carnation. Indian Phytopathology 61: 152-154.

Chandel S. 2011. Management of carnation wilt using bioresources and soil solarisation. Plant Disease Research 26: 168.

Dehne H W, Schonbeck F and Bltruschat H.1978. The influence of endotrophic mycorrhiza on plant diseases. 3, chitinase activity and the ornithine cycle. Zeitschrift fur Pflanzenkrankheiten und Pflanzenschutz 85: 666-678.

Dehne H W. 1982. Interaction between vesicular arbuscular mycorrhizal fungi and plant pathogens. Phytopathology 72: 1115-1119.

Ebtehag E B, Nemat M A, Azza S T and Hoda A H. 2009. Antagonistic activity of selected strains of Rhizobacteria against Macrophomina phaseolina of soyabean plants. American-Eurasian Journal of Agriculture and Environment Science 5: 337-347.

Gamliel A and Stapleton J J. 1993. Effect of chicken compost or ammonium phosphate and solarization on pathogen control, rhizosphere organisms and lettuce growth. Plant Disease 77: 886-889.

Gawande A A, Mahorkar V K, Joshi P S, Deo D D and Jadhao B J. 2001. Effect of soil solarization and bioagents on yield and quality of annual chrysanthemum (Chrysanthemum coronarium) cv. local bijali. Agricultural Science Digest 21: 52-54.
Gerdemann J W and Nicolson T H. 1963. Spores of mycorrhizal Endogone species extracted from soil by wet sieving and decanting. *Transaction of the British Mycological Society* 46: 235-244.

Hortink H A J and Fahy P C. 1986. Basis for the control of soil-borne plant pathogens with composts. *Annual Review of Phytopathology* 24: 93-114.

Karimi E, Rouhani H, Zafari D, Khodakaramian G and Taghinasab M. 2007. Biological control of vascular wilt disease of carnation caused by *Fusarium oxysporum* f.sp. *dianthi* by *Bacillus* and *Pseudomonas* strains isolated from rhizosphere of carnation. *Journal of Science and Technology of Agriculture and Natural Resources* 11: 309-20.

Katan J. 1981. Solar heating (Solarization) of soil for the control of soil-borne pests. *Annual Review of Phytopathology* 19: 211-236.

Khallal S M. 2007. Induction and modulation of resistance in tomato plants against *Fusarium* wilt disease by bioagent fungi (arbuscular mycorrhiza) and/or hormonal elicitors (jasmonic acid & salicylic acid): 2 - changes in the antioxidant enzymes, phenolic compounds and pathogen related-proteins. *Australian Journal of Basic and Applied Sciences* 1: 717-732.

Kishore C and Kulkarni S. 2008. Management of carnation wilt caused by *Fusarium oxysporum* f.sp. *dianthi*. *Journal of Plant Disease Sciences* 3: 17-20.

Lazarovits G, Tenuta M and Kenneth L C. 2001. Organic amendments as a disease control strategy for soil-borne diseases of high-value agricultural crops. *Australasian Plant Pathology* 30: 111-117.

Lodha S and Israel S. 2005. Combining solar heat, on-farm wastes and summer irrigation for improved control of *Fusarium oxysporum* f.sp. *cumini*. *Acta Horticulturae* 698: 181-6.

Maheshwari D K, Dubey R C, Aeron A, Kumar B, Kumar S, Tewari S and Arora N K. 2012. Integrated approach for disease management and growth enhancement of *Sesamum indicum* L. utilizing *Azotobacter chroococcum* TRA2 and chemical fertilizer. *World Journal of Microbiology Biotechnology* 28: 3015-3024.

Manka M, Fruzynska-Jozwiak D, Pokojska-Burdziej A, Dahm H. 1997. Promoting effect of *Trichoderma* on cutting growth in biocontrol of *Fusarium* carnation wilt. *Folia Horticulturae* 9: 3–13.

Martinez G and Pinzon L. 1999. New strategies in the integrated disease management of the vascular wilt of carnation (*Dianthus caryophyllus* L.) caused by *Fusarium oxysporum* f.sp. *dianthi* (Prill. & Del.) Snyd. & Hans. *Acta Horticulturae* 482:139-144.

Martínez MA, Martínez MC, Bielza P, Tello J, Lacasa A. 2011. Effect of biofumigation with manure amendments and repeated biosolarization on *Fusarium* densities in pepper crops. *Journal of Industrial Microbiology and Biotechnology* 38:3–11.

Martinez-Medina A, Pascual J A, Lloret E and Roldan A. 2009. Interactions between arbuscular mycorrhizal fungi and *Trichoderma harzianum* and their effects on *Fusarium* wilt in melon plants grown in seedling nurseries. *Journal of Science Food Agriculture* 89: 1843-1850.

Melero-Vara J M, Lopez-Herrera C J, Basallote-Ureba M J, Navas J A, Lpez M, Vela M D, Gonzalez L. 2005.
Physical and Chemical Methods of Controlling Fusarium Wilt of Carnation as Alternatives to Methyl Bromide Treatments. Acta Horticulturae 698: 175-179.

Morandi D. 1996. Occurrence of phytoalexins and phenolics compounds in endomycorrhizal interaction and their potential role in biological control. Plant and Soil 185: 241-245.

Mortan J B. 1988. Taxonomy of Vamycorrhizal fungi: classification nomenclature and identification. Mycotaxon 32: 267-324.

Mwangi M W, Monda E O, Okoth S A and Jefwa J M. 2011. Inoculation of tomato seedlings with Trichoderma harzianum and arbuscular mycorrhizal fungi and their effect on growth and control of wilt in tomato seedlings. Brazilian Journal of Microbiology 42: 508-513.

Negi H S. 2009. Integrated management of wilt of carnation caused by Fusarium oxysporum f.sp. dianthi (Prill. and Del.) Snyd. and Hans. M Sc. Thesis, Dr. Y S Parmar UHF, Nauni, Solan, pp. 39-58.

Núñez-Zofío M, Larregla S, Garbisu C. 2011. Application of organic amendments followed by soil plastic mulching reduces the incidence of Phytophthora capsici in pepper crops under temperate climate. Crop Protection 30:1563–1572.

Patel D J. 2001. Soil solarization for management of soil-borne plant diseases. Journal of Mycology and Plant Pathology 31: 1-8.

Porras M, Barrau C. and Romero F. 2009. Effects of Trichoderma and soil solarisation on Phytophthora cactorum in strawberry fields. Acta Horticulturae 842: 649-52

Raj H and Sharma S D. 2009. Integration of soil solarization and chemical sterilization with beneficial microorganisms for the control of white root rot and growth of nursery apple. Scientia Horticulture 119: 126-131.

Raj H. 2004. Effect of solarization of farmyard manure amended soil for management of damping-off caused by Pythium ultimum, Rhizoctonia solani, Sclerotium rolfsii in vegetable crops in nurseries. Indian Journal of Agricultural Sciences 74: 425-429.

Raj H. and Upmanyu S. 2013. Integration of soil solarization, corn treatment and soil amendment for the management of Fusarium wilt of gladiolus. Indian Phytopathology 66: 28-33.

Sharma P. 2000. An integrated approach for the management of carnation wilt caused by Fusarium oxysporum f.sp. dianthi (Prill. and Del.) Snyd. and Hans. New Botanist 27: 143-50.

Sharma S D, Kumar P and Bhardwaj S K. 2011. Screening and selecting novel AM fungi and Azotobacter strain for inoculating apple under soil solarization and chemical disinfestation with mulch practices for sustainable nursery management. Scientia Horticulturae 130: 164-174.

Sharma S N, Chandel S and Tomar M. 2005. Integrated management of Fusarium yellows of gladiolus caused by Fusarium oxysporum f.sp. gladioli Snyder and Hans. under polyhouse conditions. In: Integrated Plant Disease Management, Sharma R C and Sharma J N (eds.), Scientific Publisher, Jodhpur, pp. 221-229.

Smith F W. 2002. The phosphate uptake mechanism. Plant and Soil 245: 105-114.

St-Arnaud M, Hamel C, Caron M and Fortin J A. 1994. Inhibition of Pythium ultimum in roots and growth substrate of mycorrhizal Tagetes patula
colonized with *Glomus intraradices*. 
*Canadian Journal of Plant Pathology* 16:187-194.

St-Arnaud M, Hamel C, Vimard B, Caron M and Fortin J A. 1997. Inhibition of *Fusarium oxysporum* f.sp. *dianthi* in the non-VAM species *Dianthus caryophyllus* by co-culture with *Tagetes patula* companion plants colonized by *Glomus intraradices*. *Canadian Journal of Botany* 75: 998-1005.

Stevens C, Khan V A, Rodriguez-Kabana R, Ploper C D, Backman P A, Collins D J, Brown J E, Wilson M A and Igwegde E C K. 2003. Integration of soil solarization with chemical, biological and cultural control for the management of soil-borne diseases of vegetables. *Plant and Soil* 253: 493-6.

Subba Rao N S. 1986. *Soil Micro-organisms and Plant Growth*. Oxford and IBH Co. Pvt. Ltd, New Delhi. 314p.

Umesh M P and Mane R Y. 2010. Antagonistic effect of Bio-fertilizers against seed borne mycoflora of tomato (*Lycopersicum esculentum*). *Journal of Agricultural Sciences* 1: 255-258.

Verma S and Dohroo N P. 2003. Population dynamics of VAM fungi and its effect on wilt of pea caused by *Fusarium oxysporum* f.sp. *pisi*. In: *Integrated plant disease management*, Sharma R C and Sharma J N (eds.), Scientific Publishers, Jodhpur, pp. 85-92.

Weber Z, Werner M, Frużyńska-Jóźwiak D. 1998. Biological protection of carnation, asparagus and babies'-breath against different *formae speciales* of *Fusarium oxysporum* Schlecht. *Phytopathologia Polonica* 16: 37-43.

How to cite this article:

Savita Sharma, Harender Raj and Neeraj Sharma. 2019. Integration of Soil Solarization, Arbuscular Mycorrhizal Fungi, *Trichoderma viride, Azotobacter chroococcum* and Soil Amendments for the Management of Carnation (*Dianthus caryophyllus* L.) Wilt (*Fusarium oxysporum* f.sp. *dianthi* (Prill and Del.) Snyd. and Hans.). *Int.J.Curr.Microbiol.App.Sci.* 8(01): 2484-2499. doi: https://doi.org/10.20546/ijcmas.2019.801.263