The effect of some insecticides, natural compounds and tomato cv. Venezia with Mi gene on the nematode *Meloidogyne ethiopica* (Nematoda) reproduction

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**ABSTRACT**

Root-knot nematode (RKN) *Meloidogyne ethiopica* is quite unknown tropical plant parasitic species found in Slovenia and Europe in 2003 for the first time. The species is able to parasitize several economically important agricultural crops and therefore is very difficult to control. In our research, several approaches which can be used for *M. ethiopica* control were tested in pot experiment. The effects of some insecticides which are often used in the production in greenhouses, natural compounds as aqueous extract of *Tagetes erecta*, and the commercial natural product Azadirachtin (NeemAzal-T/S) extracted from the seeds of Indian Neem tree (*Azadirachta indica*) on the *M. ethiopica* reproduction were assessed. Test plants treatments with natural compounds reduced nematode multiplication by nearly 3 – 6 times compared to control while foliar application of Thiacloprid as well as Imidacloprid had no effect on nematode reproduction. The treatment with Volaton G granulates (Phoxim) for ground application resulted in no nematode multiplication. Additionally, *M. ethiopica* reproduction ability on the tomato cultivar “Venezia” which have a Mi gene for resistance to *M. incognita* was tested.

Key words: root knot nematode, *M. ethiopica*, control, pesticides, plant extracts, resistant tomato

**IZVLEČEK**


Ključne besede: ogorčice koreninskih šišk, *M. ethiopica*, zatiranje, pesticidi, rastlinski izvlečki, odporen paradižnik

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1 INTRODUCTION

Root-knot nematodes (RKNs) of the genus *Meloidogyne* belong to the economically most important group of plant parasitic nematodes. They are obligate parasites that attack numerous higher plant species including monocotyledons, dicotyledons, herbaceous and woody plants (Eisenback and Hirschmann, 1991). RKN causes development of galls on the plant host roots and therefore water and nutrient intake are limited. Plants infested with RKNs do not show specific above ground symptoms but different symptoms like stunting, wilting and sometimes early flower and fruit drop can be observed.

Two RKN species, *M. chitwoodi* and *M. fallax*, are quarantine pests for EU and EPPO region but several RKN species are also important for vegetable production causing substantial yield losses worldwide. Four species of RKN have been detected in Slovenia so far: *M. arenaria*, *M. incognita*, *M. hapla* and *M. ethiopica* (Širca and Urek, 2004; Širca et al., 2004a; Širca et al., 2004b). *M. ethiopica* was established in 2003 in the greenhouse situated in the Primorska region as the first finding of this species in Slovenia and Europe (Širca et al., 2004b). RKNs are very difficult to control because of their polyphagous nature of parasitism and more limited approaches to their chemical control recently available. Several RKN species are able to parasitize and successfully multiply on monocotyledonous plants as well as on woody and herbaceous dicotyledonous plants. *Meloidogyne ethiopica* was shown to be such a case with the wide host range on the following plants: broad bean (*Vicia faba*), black wattle, cabbage, cowpea, pepper, potato, pumpkin, tobacco, tomato (Whitehead, 1969). Carneiro et al. (2004, 2007) found *M. ethiopica* on grapevine, kiwi fruit and watermelon. Besides, barley, bean, beet, broccoli, carrot, cauliflower, celery, chicory, cucumber, curled dock, eggplant, endive, florence fennel, kale, kohlrabi, lettuce, melon, onion, pea, radish, spinach, strawberry, sunflower and sweet corn were also shown to be the host plant species (Strajnar et al., 2009).

*Meloidogyne ethiopica* is quite unknown tropical RKN species found in Europe for the first time and, because of that, we decided to test several approaches which can be used for their control. In this research we studied the effects of some insecticides which are frequently used in greenhouse production, natural compounds as aqueous extract of *Tagetes erecta*, and the commercial natural product of 1% Azadirachtin (NeemAzal-T/S) extracted from the seeds of Indian Neem tree (*Azadirachta indica*) on the *M. ethiopica* reproduction. Additionally, *M. ethiopica* reproduction ability on the tomato cultivar “Venezia” which have a Mi gene for resistance to *M. incognita* was tested.

2 MATERIALS AND METHODS

The experiment was carried out in the glasshouse on the tomato plants *Lycopersicon lycopersicum* cv. Volovško srce which is susceptible to RKN *M. ethiopica* (Strajnar et al., 2009). Tomato variety “Venezia” was obtained from Seminis. The experiment was randomized in blocks with five replications of each treatment including untreated pots as a control. Three chemical treatments and two treatments with natural compounds were tested (Table 1).

The tomato plants were planted individually in 16 cm diameter pots filled with 1960 g of fine sterilized sand (particle size: 0.25 – 1.0 mm). The daily growing temperatures ranged from 20 to 30°C. Watering was obtained for manual keeping 15% of moisture of the dry sand weight (Kutywa and Been, 2006). Nutrients for hydroponic grow (Flora series) were used. Nutrient concentration depended on the stage of plant development.

**Nematode inoculum**

The cultures of *M. ethiopica* were maintained on the bean *Phaseolus vulgaris* cv. Meraviglia di Venezia nano planted in sterilized sand (particle size: 0.25 – 1.0 mm) and kept in a glasshouse at 20 to 27°C. After forty-five days, the cultures were used for inoculum preparation. Inoculum of nematode eggs was prepared by shaking chopped galled bean roots in 1% sodium hypochlorite (NaClO) for 4 min to dissolve the gelatinous matrix surrounding root-knot nematode eggs (Hussey and Barker, 1973). The suspension of eggs was washed through 850, 250 and 32 µm banked sieves. The eggs on the lower sieve were washed with tap water to remove NaClO (Ehwaeti et al., 1998) and rinsed from the sieve in 40 ml of water into 50 ml polycarbonate centrifuge tubes. The tubes were centrifuged at 1500 rpm for 5 min. The pellets which contained the eggs and plant tissues were re-suspended in 40 ml of sucrose solution (454 g sucrose per 1 liter of tap water) and centrifuged at 1000 rpm for 1 min (McClure et al., 1973). The supernatants were poured through 32 µm banked sieves. The eggs were rinsed from the sieve and counted. Forty-five days old tomato plants were inoculated with the aqueous solution of 5000 eggs of *M. ethiopica* per plant.
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Chemical and natural compound treatments

Table 1: Application times and applied doses used for chemical and natural compound treatment.

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Application times</th>
<th>Dose</th>
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<tbody>
<tr>
<td>Calypso SC 480</td>
<td>Twice: two WAI and four WAI</td>
<td>3 ml/10 l H$_2$O (0.03%)</td>
</tr>
<tr>
<td>Confidor SL 200</td>
<td>Twice: two WAI and four WAI</td>
<td>6 ml/10 l H$_2$O (0.06%)</td>
</tr>
<tr>
<td>Aqueous extract of</td>
<td>Twice: one DBI and two WAI</td>
<td>100 ml/1 l H$_2$O (10%)</td>
</tr>
<tr>
<td>Tagetes erecta</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NeemAzal-T/S a</td>
<td>Twice: one DBI and two WAI</td>
<td>30 ml/10 l H$_2$O (0.3%)</td>
</tr>
<tr>
<td>NeemAzal-T/S b</td>
<td>Twice: one DBI and two WAI</td>
<td>100 ml/10 l H$_2$O (1%)</td>
</tr>
<tr>
<td>Volaton G 5%</td>
<td>Once: one DBI</td>
<td>1 kg/100 m$^2$</td>
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WA1 Week after inoculation  
DB1 Day before inoculation  
a NeemAzal-T/S was applied at the dose of 0.3%.  
b NeemAzal-T/S was applied at the dose of 1%.

Thiacloprid – Calypso SC 480 (Bayer CropScience) and Imidacloprid – Confidor SL 200 (Bayer CropScience) were used in the dosage according to manufacturer instructions for insect control applications. The applications of systemic insecticides on the leaves were performed twice. The first application was carried out two weeks after inoculation and the second four weeks after inoculation. Phoxim – Volaton G 5% (Pinus TKI d.d., Slovenia) granules were applied under the surface of the sand according to manufacturer’s instruction one day before inoculation.

Aqueous extract of *Tagetes erecta* was made from the three month old plants. 200 g of the whole plant material (roots, steam, leaves and blossoms) was mashed by a blender and soaked in 100 ml of tap water across the night. The aqueous extract was filtered through the gauze and stored at –20°C before use. 50 ml of 10% *T. erecta* aqueous extract was applied per plant twice, one day before inoculation and 14 days after inoculation. Azadirachtin – NeemAzal-T/S (Trifolio-M GmbH, Lachnau, Germany) was applied in 0.3% and 1% concentrations to the sand around the plant. The first application was done one day before inoculation and the second two weeks after inoculation.

**Analysis of nematode reproduction extent**

The experiment was terminated forty-five days after inoculation when tomato roots were removed and nematode reproduction was assessed. Males and second stage juveniles were isolated from the sand by a decanting method (Hržič, 1973) followed by Berman’s funnel extraction. Nematode eggs were isolated as previously described. Reproduction factor (R) was calculated as final nematode population divided by initial nematode population (R= Rf/Ri). The data were statistically analyzed by analysis of variance (ANOVA) and significant differences in means ranked by least significant difference multiple range test (LSD) (Statgraphics versions XV).

**3 RESULTS**

Application of systemic insecticides Calypso SC 480 and Confidor SL 200 did not significantly differ from the control on nematode multiplication (Fig. 1). The reproduction factor ranged from 25.1 to 28.2 (Tab. 2). The treatment with aqueous extract of *Tagetes erecta* was significantly different from all chemical treatments, NeemAzal-T/S treatments and from tomato cv. Venezia with Mi gene (Fig.1). The *Tagetes* aqueous extract reduced the nematode population by nearly 3 times (2.6) compared to untreated control and the reproduction factor was 10.4. Treatments with Azadirachtin had a significant effect on the reproduction of nematodes compared to untreated plants (Fig. 1, Fig. 2). The reproduction factors were 4.3 and 6.2 for higher (1%) and lower (0.3%) concentration treatments, respectively. The analyses of Azadirachtin treatments also revealed deformed eggs of *M. ethiopica* (Fig. 3A) which were not viable and therefore not included in the final population sum.
Figure 1: The effect of different chemical and natural compound treatments and tomato cv. Venezia on the final population of *M. ethiopica*. The data are presented as means and standard deviation of five replicates. a, b, c and d represent homogenous groups (not significantly different) at $P \leq 0.05$ by LSD multiple range test. NeemAzal-T/S a: application at 0.3% dose. NeemAzal-T/S b: application at 1% dose.

Figure 2: The reduction of nematode infection and reproduction on tomato treated with 1% NeemAzal-T/S (A) compared to an untreated control plant (B).
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Figure 3: Deformed eggs of *M. ethiopica* were observed after Azadirachtin treatments (A); Healthy egg of *M. ethiopica* from an untreated control tomato plant (B).

No nematode reproduction was observed on the roots of plants where Volaton G 5% was used and on the roots of tomato cv. Venezia. The reproduction factors of those two variants significantly differed from all the other variants including control.

Table 2: Reproduction factors of *M. ethiopica* for different chemical and natural compound treatments and tomato cv. Venezia.

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Reproduction factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>27.3</td>
</tr>
<tr>
<td>Calypso SC 480</td>
<td>28.2</td>
</tr>
<tr>
<td>Confidor SL 200</td>
<td>25.1</td>
</tr>
<tr>
<td>Aqueous extract of <em>Tagetes erecta</em></td>
<td>10.4</td>
</tr>
<tr>
<td>NeemAzal-T/S 0.3%</td>
<td>6.2</td>
</tr>
<tr>
<td>NeemAzal-T/S 1%</td>
<td>4.3</td>
</tr>
<tr>
<td>Volaton G 5%</td>
<td>0.0</td>
</tr>
<tr>
<td>Tomato cv. Venezia (Mi gene)</td>
<td>0.0</td>
</tr>
</tbody>
</table>

4 DISCUSSION

Root knot nematodes are important plant pests with a worldwide distribution. In the agricultural production they cause great yield losses when they over reproduce therefore it is important to develop useful control methods. RKN *M. ethiopica* is difficult to control because of their wide host range attacking most of the vegetable grown in the greenhouses and high rates of reproduction. The nematode can parasitize monocotyledons as well as dicotyledons which limits the use of crop rotation as a strategy to control the nematode reproduction. Besides, chemical control approaches are becoming less available recently. On the contrary, natural compounds are more acceptable for the environment and they are becoming more and more popular. However, their usefulness and efficiency for pest control needs to be assessed.

The aim of our experiment was to establish if any insecticides or natural compound influence the *M. ethiopica* reproduction, which is new species for Slovenian environment. The effects of some insecticides which are frequently used in greenhouse production, natural compounds as extract of *Tagetes erecta* and natural product of 1% Azadirachtin (NeemAzal-T/S) extracted from the seeds of Indian Neem tree (*Azadirachta indica*) as well as tomato cv. Venezia with Mi resistance gene on *M. ethiopica* reproduction were tested. The data showed that the insecticide Volaton G 5% and tomato with Mi gene prevented the reproduction of the nematode. Our results showed that Mi gene has significant effect on *M. ethiopica* reproduction suggesting that production of resistant cultivars with Mi gene could be successful non-chemical measure for controlling the nematode. Foliar application of...
Thiacloprid or Imidacloprid had no effect on nematode reproduction as both treatments showed no significant difference from the control plants.

Natural compounds treatments reduced nematode reproduction by 2.6, 4.4 and 6.3 times for Tagetes extract, 0.3% NeemAzal-T/S and 1% NeemAzal-T/S, respectively, compared to untreated controls. The analyses of final nematode populations after Azadirachtin treatments revealed deformed non-viable eggs of *M. ethiopica*. Such observations are reported for the first time on nematodes. However, similar observations were reported for common cockchafer *Melolontha melolontha* L. where 80-90% of females which were fed with Azadirachtin treated oak leaves were not able to produce viable eggs (Malinowski et al., 2003). The treatment with the *Tagetes* aqueous extract was not so efficient as Azadirachtin treatments but more concentrated extract could give better results which we plan to examine in the future.

5 ACKNOWLEDGEMENTS

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6 REFERENCES


