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Alleviation of Nematode-Mediated Apple Replant Disease by Pre-Cultivation of *Tagetes*

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Abstract: Apple replant disease (ARD) is a severe problem in orchards and tree nurseries caused by yet unknown soil biota that accumulate over replanting cycles. This study tested the contribution of nematodes to ARD, and cultivation of *Tagetes* as a control option. In a pot experiment, *Tagetes patula* or *Tagetes tenuifolia* were grown in ARD soil, incorporated or removed. Nematodes extracted from untreated ARD soil and washed on 20 µm-sieves induced ARD symptoms when inoculated to apple saplings growing in a sterile substrate. In contrast, nematodes from *Tagetes* treated ARD soil did not reduce root growth compared to uninoculated plants, irrespective of *Tagetes* species and incorporation. In plots of five apple tree nurseries or orchards, either *Tagetes* or grass was grown on ARD soil. Nematodes extracted from the grass plots and inoculated to apple saplings significantly reduced plant growth compared to nematodes from *Tagetes* plots for all five farms. Apple rootstocks showed overall a significantly higher increase in shoot base diameter when grown on *Tagetes*-treated plots compared to grass plots, while this effect differed among farms. Plant-parasitic nematodes were too low in abundance to explain plant damage. In conclusion, the free-living nematodes involved in ARD can be controlled by *Tagetes*.

Keywords: *Tagetes*; marigold; apple replant disease; nematodes; pest control; soil biome management; *Malus*

1. Introduction

Apple replant disease (ARD) has been recognized throughout pome fruit production regions of the world and has been studied extensively [1–4]. When establishing a new orchard on replant sites, trees commonly exhibit poor growth that shows in the significantly reduced shoot growth, necrosis and patchy blackening of root cells, impaired root hair development, and low cell vitality, which may lead to root death [5]. As a result, fruit yield and quality are significantly reduced [6]. Although consensus regarding the causality of replant disease has not been fully realized, mitigating measures such as soil pasteurization or fumigation significantly improved the growth of apple plants which gives evidence that the disease is caused by biotic factors [7]. Originally, replant disease, also known as soil fatigue, was described as the phenomenon that soil gradually loses its

capacity to support growth of a specific plant after replanting without any obvious reasons, especially if plant damage can not be attributed to known pathogens or plant-parasitic nematodes [8,9]. However, many different pathogens of apple plants have been occasionally found in ARD soil or in apple roots growing in ARD soil, including fungi, bacteria, oomycetes, and nematodes [10–17], while none of them was consistently associated with disease severity. Apple roots show a non-systemic, localized reaction where they are in direct contact with ARD soil, resulting in local brownish discoloration and decreased ^{15}N uptake [18]. Accumulation of phenolic compounds or phytotoxins in disease-affected parts of the roots has been discussed to play a role in the disease [16,19]. Recently, a transcriptomic analysis of the molecular responses of apple plants to ARD soils showed peculiar defense reactions to biotic stress, especially up-regulation of genes for phytoalexin synthesis [20,21].

Nematodes are abundant in soils and play an essential role in ecosystem functions and services [22]. Some of the nematodes, mainly the endoparasitic phytonematodes [23,24], are a major limiting biotic factor of productivity [25,26]. These nematodes migrate through the soil in search of a host plant, invade roots and feed on the cytoplasm of cells. Etiological relations between some nematodes and microbes in soil-borne disease complexes are known [27,28]. *Pratylenchus penetrans* was reported to exacerbate apple replant disease (ARD), in addition to unknown biotic factors [29,30]. The peculiar role of free-living, putatively non-parasitic nematodes in the development of ARD was demonstrated in pot experiments with apple saplings growing in sterile substrate [31]. The microbial fraction extracted from ARD soil hardly induced phytoalexins in the root, while in roots exposed to the washed nematode fraction (including body-associated microbes), phytoalexins increased by 1.7 log units. The combination of nematodes and microbes further increased phytoalexins by 3.7 log units [31]. This gave evidence that a nematode-microbe complex induced ARD, while *P. penetrans* and other plant-parasitic nematodes were hardly detected.

Effective management of nematode-mediated diseases is still challenging. Apple growers consistently rely upon the use of pre-plant soil fumigation [32], which has revealed a large site-to-site variation in efficiency and often only a short-term growth response [33]. The application of synthetic nematicides targeting nematodes poses health and environmental risks because of animal and human toxicity [34,35]. There is emerging social pressure to develop non-fumigant strategies. Crop rotation is often not applicable in fruit and nursery production systems. Anaerobic soil disinfection or biofumigation with plants or seed meal of Brassicaceae could mitigate ARD [36,37]. These methods are limited by high time requirements, costs, logistical challenges, and often by the environmentally undesirable use of plastic. Biological control options may be costly and of unpredictable efficacy [38,39]. A few *Malus* genotypes with tolerance to ARD have been described but further breeding is necessary to get tolerant rootstocks with growth advantage in different ARD soils [40].

A potential alternative management option of nematodes in ARD is the pre-plant application of *Tagetes*, which has demonstrated the capacity to provide effective and long-term suppression of nematode pests in various other agricultural systems. *Tagetes* species (often referred to as French marigold) are cultivated all over the world for ornamental purposes and also for industrial use. They produce several potentially bioactive compounds among which alpha-terthienyl is acknowledged as one of the most toxic chemicals present in the marigold tissues and roots [41]. They exhibit insecticidal, nematocidal, fungicidal, antiviral, and cytotoxic activities [41,42], and were reported to be suppressive to several soil-borne plant pathogenic fungi such as *Rhizoctonia solani* and *Fusarium solani* [43,44]. Recently, precropping with *Tagetes* revealed increased growth of apple in two ARD soils by 175% or 52%, respectively [37,45]. The effect of *Tagetes* on the nematodes related to ARD and its consequences on apple plant growth has not been explored.

The aim of this study was to elucidate the role of nematodes in ARD, and to investigate whether *Tagetes* preculture alleviates ARD by changing the nematode community in soil. The efficacy of *Tagetes* preculture to improve apple plant growth in ARD soils was

evaluated in field trials at a diverse selection of tree nurseries and apple growing farms, and the effect on plant-parasitic and non-parasitic nematodes was investigated. In biotests, the effect of nematodes extracted from *Tagetes* treated and untreated ARD soils on growth and ARD symptoms of susceptible apple plantlets was investigated. Furthermore, we tested the efficiency of different *Tagetes* species and soil incorporation on ARD in pot tests.

2. Materials and Methods

2.1 Pot experiment on the effect of *Tagetes* with or without incorporation into soil on nematode-mediated ARD

In a pot experiment, we investigated whether nematode communities recovered from ARD soil affect apple plant growth and whether preculture of *Tagetes* in ARD soil reduces the negative effect of the nematode community on apple plant growth. Since different *Tagetes* species produce different biocidal compounds, which may be exuded or not from the root, we tested whether *T. patula* and *T. tenuifolia* differ in their effect on nematode-mediated ARD, and whether incorporation of *Tagetes* into the soil is necessary for mitigation of nematode-mediated ARD. The soil was obtained from a field in the Pinneberg area near Heidgraben, Germany (53°41'57.1''N 9°40'59.4''E). Since 2009, the rootstock of the cultivar 'Bittenfelder Sämling' was planted repeatedly in two-years cycles [46]. The soil was sampled around the roots of plants at a depth of 0–30 cm and sieved through a 5-mm mesh. Samples were stored at 4°C for 1 week before the pot experiments. Seeds of *T. patula* 'Single Gold' or *T. tenuifolia* (Saatzucht Bardowick, Bardowick, Germany) were nursed separately in sterile sand. Six two-week seedlings were transferred into each pot filled with 1 L of the ARD soil. Plants were grown for 8 weeks until flowering. To investigate the effect of plant incorporation into soil, *Tagetes* plants were removed from 20 pots, chopped into 0.5-cm pieces, and 50 g per pot was mixed with the soil. *Tagetes* in the rest of the 20 pots were removed and not incorporated into the soil. Control pots were not treated with *Tagetes*. After four weeks, nematode communities (plant-parasitic and free-living) were extracted from each pot by centrifugal floatation using MgSO₄ at 1.18 specific density [47]. Nematodes were collected on 20-µm sieves and washed with sterile water. Nematodes were inoculated to in vitro propagated and acclimatized, 5-weeks old M26 apple plantlets growing in 500 mL sterile sand, by equally distributing the suspension into four 5-cm deep 1-cm wide holes in 2 cm distance around the shoot. Pots were placed in a randomized complete block design in the greenhouse. The fertilizers Hakaphos NPK (+Mg) [15:10:15(+2)] (Compo, Münster, Germany) (0.5 g per pot) and 36% calciumcarbonate (Vereinigte Kreidewerke, Söhlde, Germany) (2 g per pot) were applied weekly. Plants were watered every 2-3 days as required. The greenhouse conditions were 22 ± 2.5°C, 60 ± 8.7% relative humidity, and a 16 h photoperiod. Pots were sampled eight weeks after inoculation to determine shoot length, shoot fresh mass, leaf fresh mass, and root fresh mass. Overall, the plant growth assay comprised 20 replicates of each treatment with nematodes from ARD soil treated with *T. patula*, *T. tenuifolia*, *T. patula* with incorporation, *T. tenuifolia* with incorporation, or from untreated ARD soil. Additional controls were plants grown in uninoculated sterile substrate, and plants grown in ARD soil.

2.2 Effect of *Tagetes* in mitigating ARD in apple growing farms and tree nurseries

Two apple-growing farms with either organic (farm M) or conventional (farm J) practice, and three tree nurseries either in Northrhine-Westfalia (farm L) or Schleswig-Holstein (farms S, C) were selected where apple plants were repeatedly replanted and problems with ARD were reported (Table 1). The soils differed in texture, pH as well as N and C contents (Table 2). In in-field trials, plots were either kept under grass to maintain the status of ARD, while avoiding soil erosion and weed growth, or cultured with *T. patula* 'Nemamix' (farms S, C, M, J) or *T. erecta* (farm L) for the vegetation period of 2019. Plants were chopped and incorporated by mulching into the soil before planting of apple (Table

1). The effect of the *Tagetes* pre-treatment on apple plant growth was determined by measuring the increase in shoot diameter during the vegetation period of 2020. In the tree nurseries, the diameter of 40 rootstock plants of the middle rows of each plot was taken at the base (above the soil) shortly after planting and in November before uprooting. In the apple orchards, trees (excluding border plants) were marked at 20 cm above the grafting, and diameter was measured twice (in orthogonal direction) each in May and November 2020.

To determine the contribution of nematodes to ARD, soil samples from all plots were collected in November 2020 at a depth of 0-20 cm with 6-8 sampling points between apple plants. The soil samples were stored at 4°C before nematode communities were extracted. To determine the number of nematodes of plant-parasitic genera and non-parasitic nematodes, 250 mL soil aliquots were extracted using an Oostenbrink elutriator [47]. Nematodes were collected on three mounted 45 µm sieves, washed into a beaker, and transferred onto an Oostenbrink dish to get a clean sample. After 48 h, the nematodes in the Oostenbrink dish were collected on a 20- µm sieve, transferred to 30 mL tap water, and counted on a counting slide under an Olympus SZX12 stereomicroscope at 40×–80× magnification (Olympus, Hamburg, Germany).

Table 1. Experimental details of field trials in companies.

| Farm | <i>Tagetes</i> treatment (2019) | | | Apple cultivation (2020) | | |
|-----------------------|---------------------------------|--------|--------------------|-----------------------------|-------------------------------------|------------------|
| | Species | Sowing | Incor- poration | Apple plants | Measured plants per treatment | No. of blocks |
| L, tree nursery | <i>T. erecta</i> | Jun | Sep | Rootstock A2 | 160 | 4 |
| S, tree nursery | <i>T. patula</i> 'Nemamix' | May | Apr (2020) | Rootstock M9 | 160 | 4 |
| C, tree nursery | <i>T. patula</i> 'Nemamix' | May | Nov | Rootstock 'Bittenfelder' | 235 | 6 |
| M, organic orchard | <i>T. patula</i> 'Nemamix' | May | Nov | 'Red Prince' on M9 | 28 | 1 |
| J, orchard | <i>T. patula</i> 'Nemamix' | May | Nov | 'Sweet Tango' on M9 | 24 | 2 |

2.3 Biotest on ARD-induction by nematodes from apple plots with preceding *Tagetes* or grass cultivation

For the biotest, nematode communities were extracted from 1 L soil samples from apple plots with preceding *Tagetes* cultivation or control plots with preceding grass cover. The extraction was done by centrifugal floatation in MgSO₄ at 1.18 specific density [47]. Nematodes were collected on 20-µm sieves and washed with sterile water. The nematodes were inoculated to apple plants grown in sterile sand and incubated in the greenhouse as described above. Shoot fresh mass, leaf fresh mass, root fresh mass, and shoot length were

determined eight weeks after inoculation. Overall, the biotest comprised 10 replicates for both the *Tagetes* treated ARD soils and the grass treated ARD soils for each farm. Plants growing in uninoculated sterile sand served as control.

Table 2. Location and soil properties of experimental sites.

| Farm | Location | Soil type | Clay [%] | Silt [%] | Sand [%] | pH (CaCl ₂) | C [%] | N [%] |
|------|--|---------------------------|-------------|-------------|-------------|----------------------------|----------|----------|
| L | Northrhine-West- falia 51.83254, 7.42113 | Braunerde- Podsol | 7.4 | 16.0 | 76.6 | 5.9 | 2.95 | 0.18 |
| S | Schleswig Holstein 53.67823, 9.73118 | Podsol-Para- braunerde | 5.9 | 41.9 | 52.2 | 4.7 | 1.59 | 0.11 |
| C | Schleswig Holstein 53.63304, 9.70630 | Podsol | 5.1 | 5.5 | 89.4 | 5.3 | 2.45 | 0.14 |
| M | Lower Saxony 53.50748, 9.68593 | Kleimarsch | 30.5 | 64.0 | 5.5 | 5.9 | 3.37 | 0.32 |
| J | Lower Saxony 53.47639, 9.59167 | Pseudogley- Braunerde | 7.2 | 32.6 | 60.2 | 5.5 | 2.00 | 0.16 |

2.4 Statistical analysis

Statistical analyses were done using the GLIMMIX procedure of the software package SAS 9.4 (SAS Institute Inc., Cary, NC, United States of America). Plant growth parameters were analyzed with the assumption of a normal distribution without data transformation, which was checked by qq- and residual plots. For multiple comparisons, the *P*-value was adjusted by the method of Tukey (ADJUST=TUKEY in the LSMEANS statement). To account for overdispersion, degrees of freedom were approximated by the method of Kenward-Roger (DDFM=KENWARDROGER). To analyze the effect of *Tagetes* (TAGETES = 1 or 0) on shoot base increase (SBI) in plots of five farms, the generalized linear mixed model $SBI = TAGETES \text{ FARM}$ was used, with PLOT as a random effect. The effect for each farm was estimated by using contrasts (LSMESTIMATE statement). The effect of preceding *Tagetes* cultivation on the proportion of plant-parasitic nematodes (PPN) was tested in a generalized linear mixed model $PPN / TOTAL = TAGETES$ with FARM as random effect, binomial distribution and Logit transformation, using GLIMMIX. An effect was regarded as significant at the type III error $P \leq 0.05$.

3. Results

3.1 Effect of *T. patula* or *T. tenuifolia*, with or without incorporation into soil, on nematode-mediated ARD (pot experiment)

Washed nematode fractions from differently treated ARD soil (Heidgraben field), with or without *Tagetes* preculture and with or without soil incorporation of *Tagetes*, were tested for their effect on the growth of roots and shoots of apple plants. The plants that received this nematode inoculum from untreated ARD soil, or that were directly planted in ARD soil discolored roots. Overall, nematodes extracted from the different treatments significantly differed in their effect on the determined parameters of plant growth (MANOVA, $P < 0.0001$). Nematodes extracted from untreated ARD soil significantly reduced root fresh mass compared to nematodes from all *Tagetes* treatments of ARD soil (Table 3). Roots grown in ARD soil or grown in substrate with nematodes from ARD soil

did not significantly differ in fresh mass, but were 44-68% smaller than in the control without nematodes. In contrast, apple roots inoculated with nematodes from the *Tagetes* treatments did not significantly differ from the control without nematodes. Incorporation of *Tagetes* was not a significant factor ($P = 0.84$), while root fresh mass had a trend to be increased in treatments with *T. patula* compared to *T. tenuifolia* ($P = 0.0547$), as revealed by the respective contrasts in ANOVA.

Shoot parameters were significantly reduced by nematodes from ARD soil compared to the control without nematodes (Table 3). Treatments with *T. patula* alleviated this effect, whether incorporated into soil or not. However, nematodes from ARD soil that was treated with *T. tenuifolia* did not significantly differ in their effect on shoot parameters compared to nematodes from untreated ARD soil. Incorporation of *Tagetes* was not a significant factor, as revealed by the respective contrasts in ANOVA ($P > 0.08$). Only *T. tenuifolia*, which produced more shoot biomass than *T. patula*, showed a trend for increased shoot growth in the biotest when treatments with and without incorporation are compared. Shoot parameters were most affected by direct growth in ARD soil compared to growth in substrate that was inoculated with nematodes.

Table 3. Effect of preceding *Tagetes patula* or *T. tenuifolia* cultivation in apple replant disease (ARD) soil, and incorporation into soil after growth, on nematode communities with respect to their inhibitory effect on the growth of apple M26 plantlets.

| Source of nematodes inoculated to apple plantlets | Plant growth parameter ¹ | | | |
|--|-------------------------------------|----------------------|---------------------|---------------------|
| | Shoot length (cm) | Shoot fresh mass (g) | Leaf fresh mass (g) | Root fresh mass (g) |
| Untreated ARD soil | 13.7±3.8c | 3.7±1.4c | 2.3±0.9c | 1.9±0.6b |
| <i>T. patula</i> treated ARD soil | 18.4±1.4ab | 4.9±1.2a | 3.0±0.9ab | 3.7±0.6a |
| <i>T. patula</i> treated ARD soil (with incorporation) | 18.4±3.5ab | 4.9±0.9ab | 3.0±0.6ab | 3.8±0.8a |
| <i>T. tenuifolia</i> treated ARD soil | 11.9±5.0c | 3.5±0.9c | 2.0±0.6c | 3.4±1.1a |
| <i>T. tenuifolia</i> treated ARD soil (with incorporation) | 14.9±3.3bc | 3.9±1.3bc | 2.4±0.8bc | 3.4±1.1a |
| Control without nematodes | 21.2±2.2a | 5.3±0.5a | 3.3±0.5a | 3.4±1.0a |
| M26 directly grown in ARD soil | 5.8±5.3d | 1.6±0.3d | 0.9±0.2d | 1.1±0.6b |

¹ Different letters indicate significant differences (ANOVA, Tukey's adjustment, $n = 20$, $P \leq 0.05$)

3.2 Effect of preceding *Tagetes* cultivation on apple shoot growth in apple orchards and tree nurseries

In three apple orchards and two tree nurseries, the increase in apple trunk diameter over a season was compared among plots that have been precultivated either with *Tagetes* or with grass before planting apple rootstocks. In general, apple plants cultivated in *Tagetes* pre-cultured ARD soils grew significantly better than those in the untreated grass plots (ANOVA, $P = 0.0004$; Fig. 1). The trunk diameter at the base increased in grass plots by 2.3 mm and in *Tagetes* plots by 3.0 mm, on average. However, growth significantly differed among farms ($P = 0.0001$), and the effect of *Tagetes* was dependent on the farm (interaction effect FARM * TAGETES, $P = 0.0001$). The *Tagetes* effect was strongly evident in farms S and C, and less pronounced in farms J and L (Fig. 1). Also blocks of farm C

where hollows cause temporary water logging showed a pronounced positive *Tagetes* effect on apple trunk growth. In farm M, conditions during this season did not allow for increase of the trunk diameter on average, thus no effect of *Tagetes* was realized.

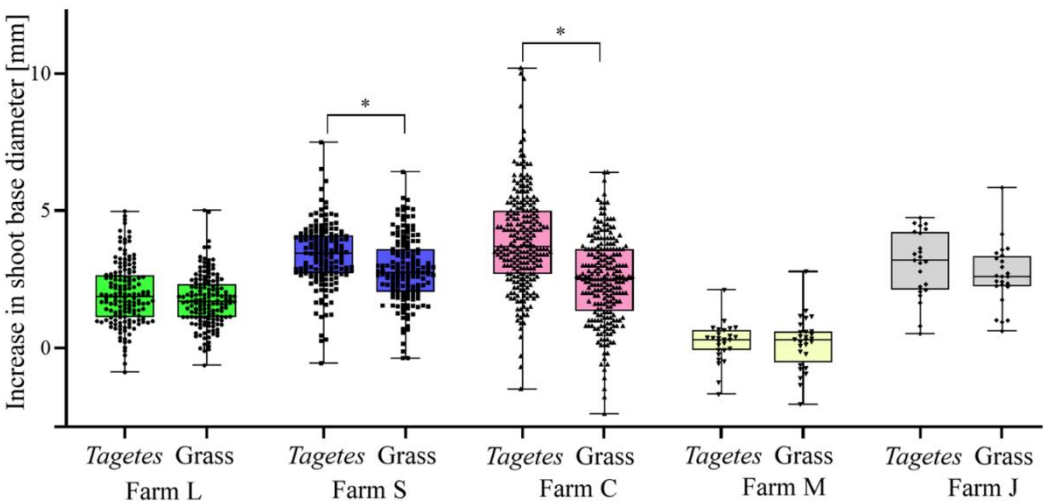


Figure 1. Increase in shoot base diameter of apple plants grown in ARD soils that were either precultured with *Tagetes* or grass. (—) median, whiskers indicate quartiles, stars indicate significant treatment effects as revealed by the respective contrasts in a generalized linear mixed model ($P < 0.05$).

Table 4. Numbers of plant-parasitic and non-parasitic nematodes per 100 ml soil in tree nurseries or apple-growing farms in adjacent plots that were either cultivated with *Tagetes* (*Tag*) or grass (*G*) before replanting of apple rootstocks.

| Genus / type | Tree nursery L | | Tree nursery S | | Tree nursery C | | Apple orchard M | | Apple orchard J | |
|----------------------|----------------|------|----------------|------|----------------|------|-----------------|-----|-----------------|------|
| | G | Tag | G | Tag | G | Tag | G | Tag | G | Tag |
| <i>Rotylenchus</i> | | | 4 | | 112 | 80 | | | | |
| <i>Meloidogyne</i> | | | 8 | | | | 44 | 16 | | |
| <i>Paratylenchus</i> | 4 | | 80 | | | | | | 128 | 200 |
| <i>Pratylenchus</i> | | | 8 | | 20 | | | | | |
| <i>Trichodorus</i> | | | 56 | 4 | 4 | | | | | |
| Non-parasitic | 2744 | 3176 | 1656 | 1472 | 256 | 1104 | 384 | 594 | 1024 | 1464 |

3.3 Effect of preceding *Tagetes* cultivation on plant-parasitic genera and total nematodes in apple orchards and tree nurseries

The numbers of plant-parasitic nematodes in the apple orchards and tree nurseries were generally low, with 77 individuals per 100 ml soil on average (Table 4). The genus *Pratylenchus* that was previously linked to ARD was only detected in low numbers in two tree nurseries, farms S and C. The genera *Rotylenchus*, *Meloidogyne*, or *Trichodorus* were each detected in two of the farms. *Paratylenchus* was the most abundant plant parasite and detected in three farms. None of the genera was detected in all ARD soils. In apple plots with preceding *Tagetes* cultivation, the proportion of plant-parasitic nematodes was sig-

nificantly lower than in the ARD plots ($P = 0.0002$). Farm L was characterized by the highest number of non-parasitic nematodes, while only four *Paratylenchus* were found. A 16 % increment of the non-parasitic nematodes was achieved after pre-treatment with *Tagetes erecta*, while *Paratylenchus* became undetectable. In farm S, the plant-parasitic nematodes were reduced by 97% and non-parasitic nematodes were reduced by 11% after cultivation of *T. patula*. In farm M, the non-parasitic nematodes achieved a 55 % increment after *Tagetes* treatment and *Meloidogyne* was reduced by 64%. In farm J, the *Tagetes* treatment increased the abundance of non-parasitic nematodes by 43% and the genus *Paratylenchus* by 36%. In soil of farm C, the non-parasitic nematodes were increased by 331% after the pre-treatment with *Tagetes* while *Rotylenchus* was reduced by 42%.

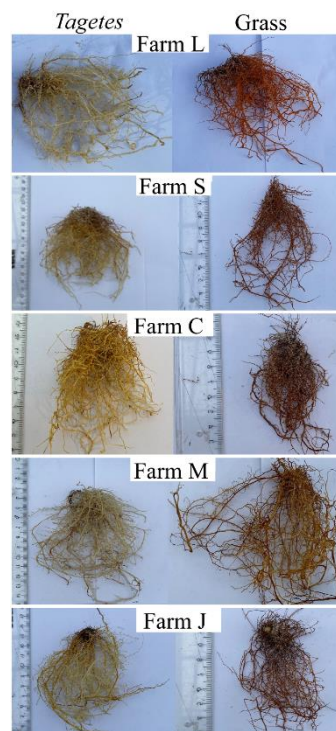


Figure 2. Discoloration and stunting of apple roots in the biotest of nematode fractions extracted from soils of plots in tree nurseries (farms L, S, C) and apple orchards (farms M and J), that were precultured with either *Tagetes* or grass before planting of apple rootstocks.

3.4 Biotest with inoculation of nematodes from apple plots with preceding *Tagetes* or grass cultivation

Nematodes from plots that were not treated by *Tagetes* before replanting of apple rootstocks caused browning and size reduction of roots of inoculated apple plantlets, which is typical for ARD (Fig. 2). A two-factor ANOVA revealed significantly higher root fresh mass of plants that were inoculated with nematodes from *Tagetes*-treated ARD soils compared to grass-treated ARD soils ($P = 0.0001$). The factor FARM also had a significant effect on root fresh mass ($P = 0.0001$). The interaction of FARM and TREATMENT was not significant ($P = 0.14$). Contrasts revealed for each farm a significant *Tagetes* effect on root fresh mass in the biotest ($P < 0.007$), while the difference between treatments was most pronounced for farm L, followed by farms S and M (Fig. 3).

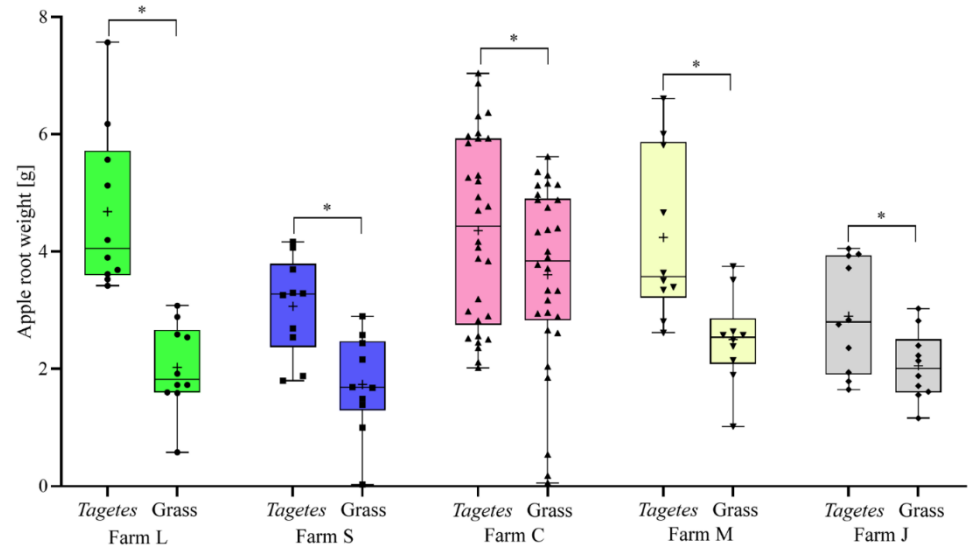


Figure 3. Root fresh mass of apple plantlets in response to inocula of nematodes recovered from ARD soils that were either precultured with *Tagetes* or grass before replanting apple. (—) median; whiskers indicate quartiles; stars indicate significant treatment effects as revealed by the respective contrasts in a generalized linear mixed model.

Table 5. Vegetative growth of apple M26 plants in pots inoculated with nematodes extracted from ARD soils pre-cultured with *Tagetes* or grass.

| Farm | Treatment | Plant growth parameter ¹ | | |
|------|----------------|-------------------------------------|----------------------|---------------------|
| | | Shoot length (cm) | Shoot fresh mass (g) | Root fresh mass (g) |
| L | <i>Tagetes</i> | 6.2±1.1a | 3.5±2.9a | 4.9±1.4a |
| | Grass | 5.1±0.8b | 1.8±0.5b | 2.0±0.8b |
| S | <i>Tagetes</i> | 7.1±1.1a | 3.0±0.6a | 3.1±0.8a |
| | Grass | 4.7±1.3b | 1.6±0.8b | 1.7±0.8b |
| C | <i>Tagetes</i> | 6.5±1.4a | 2.3±0.7a | 4.3±1.5a |
| | Grass | 4.9±1.1b | 1.5±0.6b | 3.6±1.5b |
| M | <i>Tagetes</i> | 7.3±0.8a | 3.1±0.3a | 4.2±1.4a |
| | Grass | 5.3±0.8b | 2.1±0.3b | 2.5±0.8b |
| J | <i>Tagetes</i> | 6.6±0.7a | 2.3±1.0a | 2.9±1.0a |
| | Grass | 5.3±0.9b | 1.3±0.6b | 2.1±0.6b |

¹ Mean ± SD (n = 10). Different letters indicate significant differences between *Tagetes* and grass-treated ARD plots revealed by ANOVA for each farm separately.

Shoot fresh mass and shoot length were significantly larger when plants were inoculated with nematodes from the *Tagetes* treated ARD plots compared to inoculation of nematodes from the grass treated ARD plots ($P = 0.0001$, Table 5). The factor FARM had a significant effect on shoot fresh mass ($P = 0.0089$), but not on shoot length ($P = 0.33$). The respective interactions of FARM and TREATMENT were not significant. Contrasts revealed for each farm a significant *Tagetes* effect on shoot fresh mass ($P < 0.006$), while the difference between treatments was most pronounced for farm L, followed by farm S (Table 5).

4. Discussion

In this study, the efficacy of *Tagetes* preculture to improve apple plant growth in ARD soils has been shown in field trials of a diverse selection of orchards and tree nurseries. All studied farms had major problems with ARD, especially the tree nurseries that typically replant apple rootstocks every or every second year, depending on their specialization on rootstock production or grafting, respectively. The seasonal increase in shoot base diameter on *Tagetes* plots was on average 30% higher than on adjacent control plots where grass was grown instead (Fig. 1). This effect was pronounced for the tree nurseries, while the apple-growing farm J showed the same trend. The other apple-growing farm experienced unfavourable conditions during the season, so that the stems did not show detectable increase in diameter. Therefore, an effect of preceding *Tagetes* cultivation on stem growth could not be detected there. However, when nematodes were extracted from these plots and inoculated to susceptible M26 apple plantlets, the effect of the preceding *Tagetes* cultivation on how the nematodes affected apple roots became very clear for all farms (Fig. 3). In contrast to the nematode fraction from *Tagetes* plots, the nematodes from ARD plots with preceding grass cultivation caused the discolored and stunted roots that are typical for ARD (Fig. 2). This confirms our previous finding that the nematode fraction contained one main driver of ARD [31]. As in this study, the nematode fraction was obtained from soil by floatation on a dense MgSO_4 solution, and collection and washing on a 20 μm -sieve. The nematode fraction thus contains the microbes that are associated with the nematode bodies. These microbes were shown to synergistically enhance ARD symptoms together with the nematode fraction, but they had hardly any effect when inoculated to apple plantlets alone [31]. While floatation is a standard technique to retrieve nematodes from soil [48], and the fraction on the 20 μm -sieve mainly contains nematodes when microscopically analysed, it might contain other small organisms and microbes associated with small organic particles, which may play a role in ARD. This needs to be ruled out in future studies. Neither in the previous nor in the present study played plant-parasitic nematodes a significant role in ARD. They were only detected in low numbers in the ARD soils. None of the genera was detected in all farms and could be associated with ARD. The genus *Pratylenchus* was only detected in two tree nurseries. It has been frequently reported in association with ARD [14,29,49–51], with a damage thresholds of 50 *Pratylenchus* per 100 ml soil [52]. However, in other studies the reduction of *Pratylenchus* by nematicides did not improve tree growth in ARD affected orchards [32], or was hardly correlated with the gain in plant growth ($R^2 = 0.186$) [53], or ARD symptoms were observed despite reduction of *Pratylenchus* by soil treatment [30,52]. In an orchard in Washington, a heavy ARD infestation was treated either by fumigation or *Brassica* seed meal [54]. While *Pratylenchus* reached high densities in apple roots in the second year after fumigation but not in the *Brassica* seed meal treatment, this difference was not reflected by trunk increase or apple yield. Plant-parasitic nematodes thus can eventually increase damage of ARD-affected roots but did not have a relevant contribution to ARD at least in our study. Nevertheless, the *Tagetes* treatment in our study further reduced the relative abundance of phytoparasites within the nematode community, which is an added benefit. The count data showed that *Tagetes* changed the structure of the nematode community, while the total number of nematodes in soil tended to increase (Table 4).

In the pot experiment, both *T. patula* and *T. tenuifolia* significantly reduced plant damage by the nematode fraction from the treated ARD soil that was inoculated to apple plantlets. For the *T. patula* treatment, nematode-mediated ARD was abolished, as shoot and root growth did not significantly differ from the uninoculated control. The *T. tenuifolia* treatments were not equally efficient. Shoot growth was less improved compared to the *T. patula* treatments, and roots also showed this trend. The farms applied mainly *T. patula* to reduce ARD, but farm L used *T. erecta* instead. Notably, farm L showed the most pronounced reduction of nematode-mediated ARD among the farms in the biotest (Fig. 3). In contrast, the effect of *T. erecta* preculture on stem growth in the field was not significant for this tree nursery, while the other two nurseries significantly improved stem growth after *T. patula* preculture. This coincided with the overall lower plant growth in farm L

compared to the other two tree nurseries. It should be considered that the rootstock A2 cultivated by farm L may be more susceptible to ARD than M9. In farm M, lack of trunk growth in the field completely prevented the detection of a *Tagetes* effect, while the biotest clearly revealed a significant treatment effect. The farms differed in the type of cultured rootstock, soil type, soil texture, and other parameters, so that the differential effect of the *Tagetes* species might be better reflected by the growth of apple plants in the more controlled biotest. *T. patula* was reported to be attractive for root invasion by *P. penetrans* and other endoparasites, but the nematodes cannot multiply in the roots [55]. Cultivation of *Tagetes* as pre-crop reduced populations of root-knot and lesion nematodes and substantially increased yield in the subsequent crop (melon, tomato, or potato) [56,57]. This led to the view that *Tagetes* acts as a trap crop. However, metabolites released from the roots of mature *Tagetes* plants act against diverse herbivorous and non-herbivorous nematodes, especially thiophene compounds like α -terthienyl [58–60]. A study on the amount of nematocidal thiophenes in roots of *Tagetes* species revealed the highest concentration in *T. tenuifolia*, followed by *T. patula* and *T. erecta* [59]. Interestingly, when comparing the amount of thiophenes produced by the roots per area of cultivated *Tagetes*, then probably *T. minuta* has the highest effect on nematodes, followed by *T. patula*, *T. erecta*, and *T. tenuifolia* (320 / 39 / 14 / 12 mg m⁻², respectively). This coincides with the slightly better performance of *T. patula* compared to *T. tenuifolia* in our pot experiment.

At the farms, *Tagetes* was incorporated into soil either in autumn or in spring. It is unclear whether soil incorporation of *Tagetes* contributes to the effect on ARD. In the pot experiment, incorporation of *T. patula* had no effect on growth of apple plantlets in the subsequent biotest with the inoculated nematode fraction. The treatment with *T. tenuifolia* showed a trend for increased shoot growth when the soil was incubated with the chopped and incorporated plants compared to the treatment without incorporation. This might be explained by a green manure effect on nematodes, rather than an introduction of nematocidal compounds from the shoot, because actively growing roots of *Tagetes* act most effective on nematodes while root extracts or other parts of the plant are less efficient [61,62].

Tagetes cultivation is already applied in tree nurseries to control *P. penetrans*. However, farmers are not yet aware that it is also a management option against ARD. *Tagetes* cultivation is less expensive compared to physical soil disinfection, and less damaging to the environment compared to chemical soil fumigation. However, it needs to be investigated for how long *Tagetes* cultivation suppresses the ARD causing biota, and to what extent production losses during *Tagetes* cultivation is outweighed by the gain in soil health.

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References

1. Mai, W.F.; Merwin, I.A.; Abawi, G.S. Diagnosis, etiology and management of replant disorders in New York cherry and apple orchards. *Acta Hortic.* **1994**, *363*, 33–42, doi:10.17660/ActaHortic.1994.363.5.
2. Mai, W.F.; Abawi, G.S. Controlling replant diseases of pome and stone fruits in Northeastern United States by preplant fumigation. *Plant Dis.* **1981**, *65*, 859–864.
3. Winkelmann, T.; Smalla, K.; Amelung, W.; Baab, G.; Grunewaldt-Stöcker, G.; Kanfra, X.; Meyhöfer, R.; Reim, S.; Schmitz, M.; Vetterlein, D.; et al. Apple replant disease: causes and mitigation strategies. *Curr. Issues Mol. Biol.* **2019**, *30*, 89–106, doi:10.21775/cimb.030.089.
4. Mazzola, M.; Manici, L.M. Apple replant disease: role of microbial ecology in cause and control. *Annu. Rev. Phytopathol.* **2012**, *50*, 45–65, doi:10.1146/annurev-phyto-081211-173005.
5. Grunewaldt-Stöcker, G.; Mahnkopp, F.; Popp, C.; Maiss, E.; Winkelmann, T. Diagnosis of apple replant disease (ARD): microscopic evidence of early symptoms in fine roots of different apple rootstock genotypes. *Scientia Hortic.* **2019**, *243*, 583–594, doi:10.1016/j.scienta.2018.09.014.
6. Mazzola, M. Elucidation of the microbial complex having a causal role in the development of apple replant disease in Washington. *Phytopathology* **1998**, *88*, 930–938, doi:10.1094/PHYTO.1998.88.9.930.
7. Yim, B.; Smalla, K.; Winkelmann, T. Evaluation of apple replant problems based on different soil disinfection treatments — links to soil microbial community structure? *Plant Soil* **2013**, *366*, 617–631, doi:10.1007/s11104-012-1454-6.
8. Klaus, H. Das Problem der Bodenmüdigkeit unter Berücksichtigung des Obstbaus. *Landw. Jahrb.* **1939**, *89*, 413–459.
9. Bronsart, H. von. Der heutige Stand unseres Wissens von der Bodenmüdigkeit. *Zeitschrift für Pflanzenernährung, Düngung, Bodenkunde* **1949**, *45*, 166–193.
10. Manici, L.M.; Caputo, F.; Saccà, M.L. Secondary metabolites released into the rhizosphere by *Fusarium oxysporum* and *Fusarium* spp. as underestimated component of nonspecific replant disease. *Plant Soil* **2017**, *415*, 85–98, doi:10.1007/s11104-016-3152-2.
11. Otto, G. Investigations on the cause of soil sickness in fruit trees VII. An actinomycete isolated from rootlets of apple seedlings, the probable cause of specific apple replant disease. *Journal of Cultivated Plants* **2017**, *69*, 175–179, doi:10.1399/JfK.2017.05.04.
12. Tewoldemedhin, Y.T.; Mazzola, M.; Labuschagne, I.; McLeod, A. A multi-phasic approach reveals that apple replant disease is caused by multiple biological agents, with some agents acting synergistically. *Soil Biol. Biochem.* **2011**, *43*, 1917–1927, doi:10.1016/j.soilbio.2011.05.014.
13. Tewoldemedhin, Y.T.; Mazzola, M.; Botha, W.J.; Spies, C.F.J.; McLeod, A. Characterization of fungi (*Fusarium* and *Rhizoctonia*) and oomycetes (*Phytophthora* and *Pythium*) associated with apple orchards in South Africa. *Eur. J. Plant Pathol.* **2011**, *130*, 215–229, doi:10.1007/s10658-011-9747-9.
14. Nyoni, M.; Mazzola, M.; Wessels, J.P.B.; McLeod, A. The efficacy of semiselective chemicals and chloropicrin/1,3-dichloropropene-containing fumigants in managing apple replant disease in South Africa. *Plant Dis.* **2019**, *103*, 1363–1373, doi:10.1094/PDIS-10-18-1844-RE.
15. Popp, C.; Wamhoff, D.; Winkelmann, T.; Maiss, E.; Grunewaldt-Stöcker, G. Molecular identification of Nectriaceae in infections of apple replant disease affected roots collected by Harris Uni-Core punching or laser microdissection. *J. Plant Dis. Prot.* **2020**, *127*, 571–582, doi:10.1007/s41348-020-00333-x.
16. Manici, L.M.; Kelderer, M.; Caputo, F.; Saccà, M.L.; Nicoletti, F.; Topp, A.R.; Mazzola, M. Involvement of *Dactylonectria* and *Ilyonectria* spp. in tree decline affecting multi-generation apple orchards. *Plant Soil* **2018**, *425*, 217–230, doi:10.1007/s11104-018-3571-3.

17. Tewoldemedhin, Y.T.; Mazzola, M.; Mostert, L.; McLeod, A. *Cylindrocarpon* species associated with apple tree roots in South Africa and their quantification using real-time PCR. *Eur. J. Plant Pathol.* **2011**, *129*, 637–651, doi:10.1007/s10658-010-9728-4.
18. Lucas, M.; Balbín-Suárez, A.; Smalla, K.; Vetterlein, D. Root growth, function and rhizosphere microbiome analyses show local rather than systemic effects in apple plant response to replant disease soil. *PLoS One* **2018**, *13*, e0204922, doi:10.1371/journal.pone.0204922.
19. Weiß, S.; Liu, B.; Reckwell, D.; Beerhues, L.; Winkelmann, T. Impaired defense reactions in apple replant disease-affected roots of *Malus domestica* 'M26'. *Tree Physiol.* **2017**, *37*, 1–14, doi:10.1093/treephys/tpx108.
20. Reim, S.; Rohr, A.-D.; Winkelmann, T.; Weiß, S.; Liu, B.; Beerhues, L.; Schmitz, M.; Hanke, M.-V.; Flachowsky, H. Genes involved in stress response and especially in phytoalexin biosynthesis are upregulated in four *Malus* genotypes in response to apple replant disease. *Front. Plant Sci.* **2020**, *10*, 1724, doi:10.3389/fpls.2019.01724.
21. Weiß, S.; Bartsch, M.; Winkelmann, T. Transcriptomic analysis of molecular responses in *Malus domestica* 'M26' roots affected by apple replant disease. *Plant Mol. Biol.* **2017**, *94*, 303–318, doi:10.1007/s11103-017-0608-6.
22. Yeates, G.W.; Bongers, T.; Goede, R.G.M. de; Freckman, D.W.; Georgieva, S.S. Feeding habits in soil nematode families and genera - an outline for soil ecologists. *J. Nematol.* **1993**, *25*, 315–331.
23. Elling, A.A. Major emerging problems with minor *Meloidogyne* species. *Phytopathology* **2013**, *103*, 1092–1102, doi:10.1094/PHYTO-01-13-0019-RVW.
24. Jones, J.T.; Haegeman, A.; Danchin, E.G.J.; Gaur, H.S.; Helder, J.; Jones, M.G.K.; Kikuchi, T.; Manzanilla - López, R.; Palomares - Rius, J.E.; Wesemael, W.M.L. Top 10 plant - parasitic nematodes in molecular plant pathology. *Mol. Plant Pathol.* **2013**, *14*, 946–961, doi:10.1111/mpp.12057.
25. Singh, S.; Singh, B.; Singh, A.P. Nematodes: a threat to sustainability of agriculture. *Procedia Environ. Sci.* **2015**, *29*, 215–216, doi:10.1016/j.proenv.2015.07.270.
26. Stirling, G.R. *Biological control of plant parasitic nematodes*, 2nd; CABI: Wallingford, UK, 2014, ISBN 1780644159.
27. Back, M.A.; Haydock, P.P.J.; Jenkinson, P. Disease complexes involving plant parasitic nematodes and soilborne pathogens. *Plant Pathol.* **2002**, *51*, 683–697, doi:10.1046/j.1365-3059.2002.00785.x.
28. Morris, K.A.; Langston, D.B.; Dutta, B.; Davis, R.F.; Timper, P.; Noe, J.P.; Dickson, D.W. Evidence for a disease complex between *Pythium aphanidermatum* and root-knot nematodes in cucumber. *Plant Health Progr.* **2016**, *17*, 200–201, doi:10.1094/PHP-BR-16-0036.
29. Dullahide, S.R.; Stirling, G.R.; Nikulin, A.; Stirling, A.M. The role of nematodes, fungi, bacteria, and abiotic factors in the etiology of apple replant problems in the Granite Belt of Queensland. *Aust. J. Exp. Agric.* **1994**, *34*, 1177–1182, doi:10.1071/EA9941177.
30. Jaffee, B.A.; Abawi, G.S.; Mai, W.F. Role of soil microflora and *Pratylenchus penetrans* in an apple replant disease. *Phytopathology* **1982**, *72*, 247–251.
31. Kanfra, X.; Liu, B.; Beerhues, L.; Sørensen, S.J.; Heuer, H. Free-living nematodes together with associated microbes play an essential role in apple replant disease. *Front. Plant Sci.* **2018**, *9*, 1666, doi:10.3389/fpls.2018.01666.
32. Covey, R.P., Jr.; Benson, N.R.; Haglund, W.A. Effect of soil fumigation on the apple replant disease in Washington. *Phytopathology* **1979**, *69*, 684–686.
33. Auvil, T.D.; Schmidt, T.R.; Hanrahan, I.; Castillo, F.; McFerson, J.R.; Fazio, G. Evaluation of dwarfing rootstocks in washington apple replant sites. *Acta Hort.* **2011**, 265–271, doi:10.17660/ActaHortic.2011.903.33.
34. Onkendi, E.M.; Kariuki, G.M.; Marais, M.; Moleleki, L.N. The threat of root - knot nematodes (*Meloidogyne* spp.) in Africa: a review. *Plant Pathol.* **2014**, *63*, 727–737, doi:10.1111/ppa.12202.

35. Zasada, I.A.; Halbrendt, J.M.; Kokalis-Burelle, N.; LaMondia, J.; McKenry, M.V.; Noling, J.W. Managing nematodes without methyl bromide. *Annu. Rev. Phytopathol.* **2010**, *48*, 311–328, doi:10.1146/annurev-phyto-073009-114425.
36. Mazzola, M.; Hewavitharana, S.S.; Strauss, S.L. Brassica seed meal soil amendments transform the rhizosphere microbiome and improve apple production through resistance to pathogen reinfestation. *Phytopathology* **2015**, *105*, 460–469, doi:10.1094/PHYTO-09-14-0247-R.
37. Yim, B.; Hanschen, F.S.; Wrede, A.; Nitt, H.; Schreiner, M.; Smalla, K.; Winkelmann, T. Effects of biofumigation using *Brassica juncea* and *Raphanus sativus* in comparison to disinfection using Basamid on apple plant growth and soil microbial communities at three field sites with replant disease. *Plant Soil* **2016**, *406*, 389–408, doi:10.1007/s11104-016-2876-3.
38. Agbenin, N.O. Biological control of plant parasitic nematodes: prospects and challenges for the poor Africa farmer. *Plant Protect. Sci.* **2012**, *47*, 62–67, doi:10.17221/46/2010-PPS.
39. Fourie, H.; van Aardt, W.J.; Venter, C.; Tiedt, L.R. The effects of CropGuard® on the motility, ultrastructure, and respiration of two *Meloidogyne* species. *Nematopica* **2014**, *44*, 85–92.
40. Reim, S.; Cestaro, A.; Siewert, C.; Wöhner, T.; Mahnkopp-Dirks, F.; Winkelmann, T.; Hanke, M.-V.; Flachowsky, H. Evaluation of tolerance to apple replant disease (ARD) in *Malus* germplasm. *Acta Hortic.* **2021**, 327–334, doi:10.17660/ActaHortic.2021.1307.50.
41. Karakas, M.; Bolukbasi, E. A review: using marigolds (*Tagetes* spp.) as an alternative to chemical nematicides for nematode management. *IJAEMS* **2019**, *5*, 556–560, doi:10.22161/ijaems.59.3.
42. XU, L.; Juan, C.; QI, H.; SHI, Y. Phytochemicals and their biological activities of plants in *Tagetes* L. *Chinese Herbal Medicines* **2012**, *4*, 103–117, doi:10.3969/j.issn.1674-6384.2012.02.004.
43. Supradip, S.; Suresh, W.; Kundu, A.; Kumar, B.; Decksha, J. Antifungal acetylinic thiophenes from *Tagetes minuta*: potential biopesticide. *J. Appl. Bot. Food Qual.* **2012**, *85*, 207–211.
44. Hooks, C.R.R.; Wang, K.-H.; Ploeg, A.; McSorley, R. Using marigold (*Tagetes* spp.) as a cover crop to protect crops from plant-parasitic nematodes. *Appl. Soil Ecol.* **2010**, *46*, 307–320, doi:10.1016/j.apsoil.2010.09.005.
45. Yim, B.; Nitt, H.; Wrede, A.; Jacquioud, S.; Sørensen, S.J.; Winkelmann, T.; Smalla, K. Effects of soil pre-treatment with Basamid® granules, *Brassica juncea*, *Raphanus sativus*, and *Tagetes patula* on bacterial and fungal communities at two apple replant disease sites. *Front. Microbiol.* **2017**, *8*, 1604, doi:10.3389/fmicb.2017.01604.
46. Mahnkopp, F.; Simon, M.; Lehdorff, E.; Pätzold, S.; Wrede, A.; Winkelmann, T. Induction and diagnosis of apple replant disease (ARD): a matter of heterogeneous soil properties? *Scientia Hortic.* **2018**, *241*, 167–177, doi:10.1016/j.scienta.2018.06.076.
47. Hooper, D.J.; Hallmann, J.; Subbotin, S.A. Methods for extraction, processing and detection of plant and soil nematodes. In *Plant parasitic nematodes in subtropical and tropical agriculture*, 2nd; Luc, M., Sikora, R.A., Bridge, J., Eds.; CABI: Wallingford, 2005; pp 53–85, ISBN 9780851997278.
48. European and Mediterranean Plant Protection Organization. PM 7/119 (1) Nematode extraction. *EPPO Bull.* **2013**, *43*, 471–495, doi:10.1111/epp.12077.
49. Santo, G.S.; Wilson, J.H. Effects of fenamiphos on *Pratylenchus penetrans* and growth of apple. *J. Nematol.* **1990**, *22*, 779–782.
50. Moein, S.; Mazzola, M.; Ntushelo, N.S.; McLeod, A. Apple nursery trees and irrigation water as potential external inoculum sources of apple replant disease in South Africa. *Eur. J. Plant Pathol.* **2019**, *153*, 1131–1147, doi:10.1007/s10658-018-01631-9.
51. Isutsa, D.K.; Merwin, I.A. Nematodes and fungi associated with apple replant disorder in sampled New York State orchards. *Global J. Bioscience Biotechnol.* **2014**, *3*, 174–180.

-
52. Hoestra, H. Replant disease of apple in the Netherlands. *Meded. Landbouwhogeschool Wageningen* **1968**, 68, 1–105.
 53. Oostenbrink, M.; Hoestra, H. Nematode damage and “specific sickness” in *Rosa*, *Malus* and *Laburnum*. *T. Pl.-ziekten* **1961**, 67, 264–272.
 54. Wang, L.; Mazzola, M. Field evaluation of reduced rate Brassicaceae seed meal amendment and rootstock genotype on the microbiome and control of apple replant disease. *Phytopathology* **2019**, 109, 1378–1391, doi:10.1094/PHYTO-02-19-0045-R.
 55. Nježić, B.; Sutter, N. de; Moens, M. Interaction of *Tagetes patula* cv. Single Gold with the life cycle of the plant-parasitic nematodes *Meloidogyne chitwoodi* and *Pratylenchus penetrans*. *Russ. J. Nematol.* **2014**, 22, 101–108.
 56. Ploeg, A.T. Effects of selected marigold varieties on root-knot nematodes and tomato and melon yields. *Plant Dis.* **2002**, 86, 505–508, doi:10.1094/PDIS.2002.86.5.505.
 57. Kimpinski, J.; Arsenault, W.J.; Gallant, C.E.; Sanderson, J.B. The effect of marigolds (*Tagetes* spp.) and other cover crops on *Pratylenchus penetrans* and on following potato crops. *J. Nematol.* **2000**, 32, 531–536.
 58. Hamaguchi, T.; Sato, K.; Vicente, C.S.L.; Hasegawa, K. Nematicidal actions of the marigold exudate α -terthienyl: oxidative stress-inducing compound penetrates nematode hypodermis. *Biol. Open* **2019**, 8, doi:10.1242/bio.038646.
 59. Marotti, I.; Marotti, M.; Piccaglia, R.; Natri, A.; Grandi, S.; Dinelli, G. Thiophene occurrence in different *Tagetes* species: agricultural biomasses as sources of biocidal substances. *J. Sci. Food Agric.* **2010**, 90, 1210–1217, doi:10.1002/jsfa.3950.
 60. Uhlenbroek, J.H.; Bijloo, J.D. Investigations on nematicides: I. Isolation and structure of a nematicidal principle occurring in *Tagetes* roots. *Recl. Trav. Chim. Pays-Bas* **1958**, 77, 1004–1009, doi:10.1002/recl.19580771103.
 61. Wang, K.-H.; Hooks, C.R.; Ploeg, A. Protecting crops from nematode pests: using marigold as an alternative to chemical nematicides. *Plant Dis.* **2007**, PD-35, 1–6.
 62. Natarajan, N.; Cork, A.; Boomathi, N.; Pandi, R.; Velavan, S.; Dhakshnamoorthy, G. Cold aqueous extracts of African marigold, *Tagetes erecta* for control tomato root knot nematode, *Meloidogyne incognita*. *Crop Prot.* **2006**, 25, 1210–1213, doi:10.1016/j.cropro.2006.03.008.