

RESEARCH NOTES

Suppression of *Pratylenchus penetrans* and *Tylenchorhynchus dubius* by *Trichoderma viride*

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Trichoderma spp. are common soil fungi which influence the development of plant-pathogenic fungi in the soil (1, 8). Yang et al. (9) found that *T. harzianum* influenced the severity of Fusarium wilt of cotton in the presence of *Meloidogyne incognita*. We report herein that the survival of two other plant-parasitic nematodes also was influenced by addition of comminuted cultures of *Trichoderma viride* to soil.

Trichoderma viride isolated from roots of apple (*Malus sylvestris* Mill.) was grown in 8% casamino acid liquid medium (2) [25 ml/125-ml Erlenmeyer flask on a rotary shaker (10 rpm/min) at room temperature (ca 22C)]. After 7 days, the contents of the flasks were comminuted in a blender for 3 min and then 125 ml of this mixture were diluted with an equal volume of water. Sixty ml of this mixture were mixed with 250 gm of a fine sandy loam naturally infested with 83 *Pratylenchus penetrans* and 21 *Tylenchorhynchus dubius*/100 gm of soil. The 250 gm of soil were placed in a 340-ml styrofoam cup and mixed by shaking. A sterile medium or deionized water was added to infested soil in a similar manner to provide controls. For comparison, the nematicide oxamyl and the fungicide benomyl were used in soil with and without mycelial suspensions at the rate of 10 µg (a.i.)/gm of soil. The cups of soil were randomized in 4 blocks on a laboratory bench, watered as needed, and incubated for 3 weeks at 22C. After 3 weeks, 2 samples (100 gm each) were taken from each replicate. Nematodes were extracted by the paper-filter method (4). Thus only live, motile nematodes were counted.

In the first experiment with infested soil, counts of *P. penetrans* and *T.*

dubius/100 gm of soil after 3 weeks (Table 1) showed that, although oxamyl was most effective, *T. viride*, oxamyl, and benomyl reduced survival of *P. penetrans* and *T. dubius*. The sterile medium had no effect on survival.

In another experiment, 7-day-old liquid cultures of *T. viride* were washed onto a No. 1 Watman filter paper and allowed to drain for 15 min. The mycelium was rinsed twice with deionized water, resuspended in 125 ml of deionized water, and blended for 3 min at high speed in the blender. Five ml of a water suspension containing 65-75 *P. penetrans* were mixed separately with 25 ml of each of the following: chopped mycelial suspension; filtered medium from the 7-day-old liquid cultures of *T. viride*; sterile, nonused medium; and deionized water. Each treatment was replicated 4 times and placed in a completely randomized arrangement in rows on the laboratory bench. After 3 days at room temperature

TABLE 1. Survival of *Pratylenchus penetrans* and *Tylenchorhynchus dubius* as influenced by *Trichoderma* spp.

Medium & soil additive	No. nematodes/100 gm soil*	
	<i>P. penetrans</i>	<i>T. dubius</i>
Oxamyl-treated soil	2 b [†]	1 b
Benomyl-treated soil	14 b	2 b
Infested soil + <i>Trichoderma</i>	7 b	2 b
Oxamyl-treated soil + <i>Trichoderma</i>	0 b	0 b
Infested soil + sterile medium	63 a	21 a
Infested soil + water	74 a	15 a

*Determined by tissue method 3 weeks after inoculation. Initial numbers of nematodes/100 gm soil were: *P. penetrans* - 83; *T. dubius* - 21.

[†]Figures followed by same letter not significantly different from each other ($P = 0.05$) by Duncan multiple range system.

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(ca 22 C), the mixtures were washed onto a tissue filter (4) and the active nematodes that passed through the filter were counted. The test was repeated. None of these treatments affected *P. penetrans* in these short-term tests.

These results indicate that chopped mycelium of *T. viride* mixed with soil is antagonistic to two common parasitic nematodes found in Connecticut. This finding is not surprising since *Trichoderma* spp. have been isolated from decomposing paper, and decaying paper and sawdust were toxic to *P. penetrans* (6). Although Rich and Miller (7) and Edmunds and Mai (3) did not find *T. viride* antagonistic to *P. penetrans*, it is known to be antagonistic to *Rhizoctonia solani* and other fungi. These antagonistic activities against fungi may have been due to antifungal substances produced by *Trichoderma* spp. (1).

Our results suggest that *Trichoderma* spp. in soil are in part responsible for the poor survival of *P. penetrans* and *T. dubius*. Miller (5) found that fungicides which increased hatching of eggs of the tobacco cyst nematode *Heterodera tabacum* also prevented breakdown of the paper cup containers from which *Trichoderma* spp. were isolated (unpublished data). Results presented herein suggest that *Trichoderma* spp. may have played a role in suppressing hatching of eggs of *H. tabacum*. If *Trichoderma* spp. suppress hatching or influence egg development of *H. tabacum*, they may also influence survival of *P. penetrans*.

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