Evaluation of Trichoderma spp. against Phytophthora spp.

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SUMMARY

A dual culture technique was used to evaluate the efficacy of *Trichoderma* spp. in inhibiting the growth of *Phytophthora* spp. At 72 hrs. of incubation the maximum per cent inhibition of mycelial growth of *Phytophthora nicitianae* was noted in treatment of T. harzianum (48.71 %). The maximum per cent inhibition of *Phytophthora citrophthora* was noted in treatment of *T. viride* (45.30 %) at 72 hrs. of incubation followed by T. lignorum (40.88 %) and *T. harzianum*(38.12 %).

Key words : Trichoderma, Phytophthora, Inhibition.

In the light of present day constraints on plant disease control practices, biological control is increasingly capturing the imagination of many plant pathologists and is gaining stature as a possible practical agricultural method for soil borne pathogen control. Biological control of soil borne plant pathogens by the addition of antagonistic micro-organisms to soil is a potential non-chemical means for plant disease control. The species *Trichoderma*, capable of hyperparasitizing pathogenic fungi are highly efficient antagonists.

MATERIALS AND METHODS

The antagonistic potential of *T. viride*, *T. harzianum*, T. lignorum, T. koningii and T. hamatum was assessed against Phytophthora spp. by dual culture technique on PDA medium. For this 20 ml of sterilized and cooled medium (PDA) was poured in each Petriplate (90 mm diameter) and was allowed to solidify. A 5 mm disc of Phytophthora sp. was placed at one end of the medium with the help of sterilized cork borer. Just opposite to it 5 mm disc of the Trichoderma sp. was placed. For this a week old culture of *Phytophthora* sp. and *Trichoderma* spp. in Petridishes on sterilized PDA medium were used. Four replications for *Phytophthora* sp. and control i.e. without inoculation of the Trichoderma spp. were maintained. Petriplates were incubated at 28 + 2°C temperature in inverted position. After 24, 48 and 72 hours the mycelial growth of Phytophthora sp. was measured in treated and controlled plate and per cent inhibition was calculated by the formula suggested by Vincent (1947)

$$I = \frac{C - T}{C} \times 100$$

Where,

I = Per cent inhibition in mycelial growth
C = Growth of mycelium in control (mm)
T = Growth of mycelium in treatment (mm)

RESULTS AND DISCUSSION

To evaluate the efficacy of *Trichoderma* spp. in inhibiting the growth of *Phytophthora* sp. a laboratory testing was done by using dual culture technique and the relevant data so obtained are presented in Table 1 and 2.

The data from Table 1 revealed that there was a significant reduction in the mycelial growth of Phytophthora nicotianae after 24 hrs. of incubation by T. viride (24.25 mm) as compared to other treatments. It was followed by the reduction brought about by T. harzianum (26.00 mm), T. koningii (28.00 mm), T. lignorum (31.75 mm) and T. hamatum (32.00 mm). After 48 hrs of incubation, the significant reduction in the mycelial growth of Phytophthora nicotianae was observed in treatment of T. viride (33.00 mm) followed by T. harzianum (33.25), T. koningii (35.25 mm), T. hamatum (37.25 mm) and T. lignorum (38.75 mm). Maximum mycelial growth of Phytophthora was observed in control plates (75.75 mm). After 72 hrs of incubation, the significant reduction in the mycelial growth of *Phytophthora nicotianae*. was observed in the treatment of T. harzianum (44.75 mm) followed by T. viride (45.75 mm), T. koningii (51.75 mm), T. hamatum (53.25 mm) and *T. lignorum* (54.75 mm). The maximum mycelial growth of Phytophthora nicotianae was observed in control plates (87.25 mm). At 72 hrs. of incubation the maximum per cent inhibition of mycelial growth of Phytophthora nicotianae was noted in treatment of *T. harzianum* (48.71 %) followed by *T. viride* (47.56 %), T. koningii (40.68 %) and T. hamatum (38.96 %). The minimum per cent inhibition of mycelial growth

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