

Prevalence of *Trichoderma* spp in Different Cropping Systems of Guntur District, Andhra Pradesh

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ABSTRACT

Seven different cropping systems of Guntur district, Andhra Pradesh, *viz.*, tobacco nurseries, groundnut, vegetables, cotton, chillies, citrus and rice-pulse systems were assessed for the prevalence of *Trichoderma* spp. Direct soil plating method was found better in enumeration of Trichoderma population compared to dilution plate method. Fungal population was more in tobacco and rice – pulse cropping systems, while it is less in chillies and citrus cropping systems.Direct plating of soil on the *Trichoderma* selective medium resulted in appearance of *Trichoderma* colonies only from cotton, chillies, citrus and rice-pulse system and not from tobacco nurseries, groundnut and vegetables.

Keywords : Cropping systems, Fungal population, Prevalence, Soil physicochemical properties, *Trichoderma* spp

Trichoderma spp. were extensively studied biocontrol agents in the management of soil borne plant pathogens (Papavizas, 1985; Upadhyay and Mukapadhyay, 1986). In majority of cases, success was achieved mainly in vitro but not in vivo. Of the several factors that determine the success of biological control, support of the soil and the cropping system practiced play an important role in the establishment of the applied antagonist. Further, prevalence of antagonistic soil microflora in a definite cropping system also indicates health of the soil. Accordingly, the present investigation was undertaken to assess the prevalence of native antagonistic microflora, viz., Trichoderma spp. in different cropping systems of Guntur district, Andhra Pradesh, India.

MATERIALS AND METHODS

Collection of soil samples

For the present study soil samples were collected from seven different cropping systems, *i. e.*, tobacco nurseries, groundnut, vegetables, chillies, cotton, citrus and rice-pulse systems in Guntur district, Andhra Pradesh (Table 1). For each cropping system three villages were chosen. In each village one field was selected from which five soil samples were collected and a composite sample was prepared following quartering technique.

Soil testing

Soil testing was done for pH, EC and organic carbon (Jackson, 1973), available nitrogen (Subbaiah and Asija, 1956), available phosphorus (Watanabe and Olsen, 1965), available potassium (Jackson, 1973) and available copper (Lindsay and Norvell, 1978). Standard soil test ratings were followed for nitrogen (Ramamoorthy and Bajaj, 1969), organic carbon, available phosphorus, available potassium, soil reaction and electrical conductivity (Anonymous, 1989).

Population dynamics of *Trichoderma* spp. and fluorescent Pseudomonads:

Soil samples collected from different cropping systems were assessed for the microbial load, *i.e.*, total fungal and total bacterial colony forming units (cfu) on Potato dextrose agar (PDA) and Nutrient agar (NA) media respectively by using dilution plate technique. Isolation of Trichoderma spp. from soil samples was done by dilution plate method and also directly spreading 0.25 g of each test soil sample on to the petriplates containing Trichoderma selective medium (TSM) under aseptic conditions and incubated at 28 ± 1°C for three days. Identification of species of Trichoderma was done based on cultural characteristics such as colony colour, medium pigmentation and microscopic observations of conidial characteristics as described by Gams and Bisset (1998).

RESULTS AND DISCUSSION

Of the twenty one soil samples analyzed in the present study, six were of loamy sand type (tobacco and groundnut systems) and remaining fifteen were of clay loam type (Table 2). In the loamy sand type, pH was found to be acidic (4.2 - 5.8)

| S. No | Cropping system surveyed | Sample / location number | Village | Area |
|-------|-----------------------------|--|--|------------|
| 1 | Tobacco nursery | | Subbareddy Palem Vedullapalli Kethapalam | Bapatla |
| 2 | Groundnut | | Kothapalem Stuvertpuram Kavurivaripalem Gavinivaripalem | Bapatla |
| 3 | Vegetables | | Devinuthala Narakoduru Budampadu | Narakoduru |
| 4 | Chilies | Ľ ₃ Ľ ₁ Ľ ₂ | Chebrolu Varagani Nagulapadu | Kakumanu |
| 5 | Cotton | -3 L ₁ L ₂ L ₂ | Kommuru Nagulapadu Pedanandipadu | Kakumani |
| 6 | Citrus orchards | -3 L ₁ L ₂ L | Kakumanu Kolakalur Revendrapadu | Tenali |
| 7 | Rice-pulse system | لہ بہ بہ ہے | Chiluvur Gudiwada Kolakalur Halfpet | Tenali |

Table 1. Details of soil samples collected from different cropping systems in Guntur district, Andhra Pradesh.

with normal EC (<1.0 dSm⁻¹) and low OC (<0.5%). These soils were found low in N (<280 kg ha⁻¹) and high in P_2O_5 (>50 kg ha⁻¹) and K_2O (>300 kg ha⁻¹) content. Copper content varied from 0.9 to 2.4ppm in different loamy sand type soils collected. Among the fifteen clay loam soils, ten were found to be weakly alkaline (7.5-8.5 pH), normal EC (<1.0 d Sm⁻¹), low OC (<0.5%), low N (<280 kg ha⁻¹), medium to high P_2O_5 (>20 kg ha⁻¹) and K_2O (>150 kg ha⁻¹). In the remaining five samples, four samples were found to be neutral in pH (6.0-7.5) and one sample was in acidic pH (<6.0) (sample L₂ of rice-pulse system).

Of all the samples analysed, highest mean fungal population was recorded in tobacco $(1.78 \times 10^5 \text{ cfu g}^{-1})$ and rice - pulse system $(1.57 \times 10^5 \text{ cfu g}^{-1})$ (Table 3). Least mean fungal population was recorded in chillies and citrus systems. It may be noted that though tobacco is a loamy sand type with low OC, acidic pH of the soils could be the reason for high fungal cfu. Brady (1990) stated that molds are abundant in acid surface soils where bacteria and actinomycetes offer mild competition. Low population of fungi in chillies and citrus may be attributed to their neutral (one sample each) to alkaline reaction.

In the present study, though mean fungal cfu varied from 0.43×10^5 cfu g⁻¹ to 1.78×10^5 cfu g⁻¹ soil, population of *Trichoderma* was very low, *i.e.*, only <2 colonies per g of soil could be recovered when observed on *Trichoderma* selective medium through direct soil plating. These observations were in contradiction to Srinivasa Rao (1999), who recorded 49.26 × 10⁴ cfu g⁻¹ of *G. virens*, 50.44 × 10⁴ cfu g⁻¹ of *T. harzianum* and 51.82 × 10⁴ cfu g⁻¹ of *T. viride* while working with chillies nursery system of Bapatla region in Guntur district of Andhra Pradesh. A decline in the population of *Trichoderma* in the soils analysed was observed in the present studies.

Further in the present experiment, though higher fungal population was observed in tobacco nursery soils, *Trichoderma* prevalence was found nil as no colonies were obtained even when the soil was directly plated on the TSM (Table 4). Similar

| S. No | Cropping system | Sample number | Texture | pН | EC (dSm ⁻¹⁾ | OC(%) | | Kg ha⁻¹ | | Cu |
|-------|--------------------|------------------|------------|-----|---------------------------|-------|-------|----------|------------------|-----|
| | System | number | | | (uSIII 7 | | N | P_2O_5 | K ₂ O | ppm |
| 1 | Tobacco | L₁ | Loamy sand | 5.8 | 0.23 | 0.12 | 37.6 | 47.9 | 499.5 | 2.4 |
| | | L_2 | Loamy sand | 5.1 | 0.14 | 0.33 | 40.3 | 100.8 | 335.9 | 1.1 |
| | | L_3 | Loamy sand | 4.2 | 0.10 | 0.16 | 31.3 | 82.2 | 326.8 | 1.0 |
| 2 | Groundnut | L_1 | Loamy sand | 5.8 | 0.19 | 0.17 | 57.3 | 146.8 | 599.2 | 0.9 |
| | | L_2 | Loamy sand | 6.8 | 0.27 | 0.30 | 43.0 | 122.3 | 699.0 | 1.2 |
| | | L_3 | Loamy sand | 5.4 | 0.12 | 0.25 | 41.2 | 208.0 | 726.1 | 0.9 |
| 3 | Vegetables | L_1 | Clay loam | 6.9 | 0.39 | 0.39 | 45.6 | 16.1 | 3999.8 | 1.0 |
| | | L ₂ | Clay loam | 7.3 | 0.26 | 0.35 | 42.1 | 139.5 | 5356.6 | 1.0 |
| | | L_3^{-} | Clay loam | 7.7 | 0.25 | 0.44 | 147.8 | 129.0 | 6899.1 | 1.0 |
| 4 | Chilies | L_1 | Clay loam | 7.7 | 0.26 | 0.33 | 54.6 | 70.9 | 5356.6 | 1.0 |
| | | L ₂ | Clay loam | 7.6 | 1.14 | 0.51 | 46.5 | 99.9 | 8625.0 | 1.1 |
| | | L_3 | Clay loam | 7.8 | 0.53 | 0.58 | 44.8 | 61.3 | 4811.9 | 1.1 |
| 5 | Cotton | L | Clay loam | 7.1 | 2.73 | 0.50 | 168.4 | 128.2 | 7717.0 | 1.1 |
| | | L_2 | Clay loam | 7.6 | 1.10 | 0.51 | 60.9 | 41.1 | 5446.9 | 1.2 |
| | | L_3 | Clay loam | 7.5 | 1.36 | 0.41 | 50.2 | 163.7 | 6264.6 | 1.2 |
| 6 | Citrus | L | Clay loam | 7.4 | 0.46 | 1.05 | 71.6 | 175.8 | 13073.8 | 1.2 |
| | orchards | L ₂ | Clay loam | 7.6 | 0.31 | 0.50 | 50.1 | 134.2 | 4696.6 | 1.5 |
| | | L_3^{-} | Clay loam | 7.5 | 0.22 | 0.94 | 60.0 | 35.3 | 8896.6 | 1.8 |
| 7 | | L | Clay loam | 7.5 | 0.35 | 0.23 | 48.3 | 155.6 | 644.6 | 1.8 |
| | Rice-pulse | L_2 | Clay loam | 6.1 | 0.21 | 0.77 | 59.1 | 249.2 | 853.4 | 2.0 |
| | | L_3^{-} | Clay loam | 7.6 | 0.16 | 0.23 | 50.1 | 100.8 | 3268.5 | 1.8 |

Table 2. Physico-chemical properties of soil samples collected from selected cropping systems.

Table 3. Fungal population in soils of selected cropping systems.

| S. No | Cropping system | Mean fungal population (X 10⁵ cfu g⁻¹ soil) | Species and number of <i>Trichoderma</i> isolates obtained |
|-------|--------------------|---|--|
| 1 | Tobacco | 1.78ª | Nil |
| 2 | Groundnut | 0.48° | Nil |
| 3 | Vegetable | 0.6ª | Nil |
| 4 | Cotton | 0.83 ^b | T. harzianum (3) |
| 5 | Chilli | 0.43 ^c | T. virens (2) T. longibrachiatum (1) |
| 6 | Citrus | 0.43° | T. virens (3) |
| 7 | Rice-pulse | 1.57ª | T. virens (2) T. longibrachiatum (1) |
| | CV (%) | 12.8 | 0 |
| | CD (P=0.01) | 0.25 | |
| | | | |

Figures represent mean values of three locations. Figures with similar alphabets do not differ significantly. Figures in parenthesis indicate number of Trichoderma isolates obtained.

result with zero population of Trichoderma was also obtained with groundnut soil which is also loamy sand type, and also from vegetable (clay loam) system. Low OC and application of fungicides such as copper group (in case of tobacco) and Mancozeb (in case of groundnut as seed treatment and in vegetables as spray) were contributed to nil population of Trichoderma in case of tobacco, groundnut and vegetables. Warcup (1950) stated that colony forming units obtained through direct soil plating probably reflect the number of conidia lying dormant in the soil, rather than active mycelial mass. Thus in the present investigation, low population of *Trichoderma* (2 cfu g⁻¹) obtained from different cropping systems indicated that Trichoderma is not actively proliferating in these soils, rather it may be in dormant state. This may be due to two reasons, *i. e.*, No crop situation and unfavourable soil conditions. As the samples were collected from all the cropping systems when crop is in standing stage, the low prevalence is not due to no crop situation. Instead, it appears that these soils (high pH and low OC) and the cropping practices followed are not favouring proliferation of Trichoderma.

Only twelve *Trichoderma* isolates could be recovered in the present study from cotton, chillies, citrus and rice-pulse systems (Table 4). All the three isolates recovered from cotton system were identified as *T. harzianum* and all the three isolates recovered from citrus system were identified as *T. virens*. From chillies and rice-pulse systems two isolates each of *T. virens* and one isolate each of *T. longibrachiatum* were recovered.

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