Consortial Effect of Endophytic and Plant Growth Promoting Rhizobacteria for the Management of Early Blight of Tomato Incited by Alternaria Solani

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Abstract

Tomato, early blight caused by Alternaria solani has been known to cause severe yield losses. Hence, attempts were made to develop an effective eco-friendly strategy to manage the disease using endophytic and plant growth promoting rhizobacteria. Accordingly, the strains of Bacillus subtilis (EPCO16 and EPC5) and Pseudomonas fluorescens (Pf, Py15 and Fp7) were tested individually and in combination for their effectiveness against early blight of tomato incited by A. solani under in vitro and pot culture conditions. The results revealed that the strains of Bacillus subtilis and Pseudomonas fluorescens were compatible. Under in vitro conditions the combined application of EPCO16+Pf1 was found to effectively inhibit the mycelial growth of the pathogen and promote the growth of tomato seedlings when compared to application of individual strains of the antagonists. Further, a significant reduction in early blight incidence of tomato under greenhouse conditions was observed due to the combined application of EPCO16+Pf1. These findings suggest that synergistic consortia of biocontrol agents may be successfully employed as an eco-friendly strategy for the management of early blight of tomato.

Keywords: Alternaria solani; Early blight; Tomato; Pseudomonas fluorescens; Bacillus subtilis

Introduction

The area under Tomato (Lycopersicon esculentum Mill.) cultivation is on the increase due to its fleshy fruits with high nutritive value. It occupies number one position in its nutrient contribution to human diet. In Tamil Nadu, tomato is grown in an area of 22,433 ha, with a production of 2,82,912 tonnes and a productivity of 12,611 kg/ha [1]. The crop is known to be affected by a number of diseases among which, early blight caused by Alternaria solani (Ellis and Martin) causes yield losses up to 80% [2,3]. The disease manifests as leaf spots with dark brown to black concentric rings which later coalesce and results in blighting of leaves; defoliation and shedding of immature fruits. Also, dark spots and sunken lesions appear near the base of stem resulting in stunting and girdling of stem. Presently, the management of tomato early blight disease has been done with the application of chemical fungicides. However, it may not be sustainable in the longer run as early blight disease has been done with the application of chemical fungicides. In chilli plants. Several approaches have been tried for the sustainable management of early blight of tomato. However, no attempts have been made for the management of early blight disease with mixtures of both PGPR and PGPE strains. Therefore, the present study was designed to evaluate protective effect of endophytic bacterial strains, B. subtilis (EPCO16 and EPC5) and rhizobacterial strains P. fluorescens (Pf1, Py15 and Fp7) against tomato early blight disease caused by A. solani.

Materials and Methods

Isolation of pathogen and maintenance of biocontrol agents

A. solani taken from infected tissue of diseased fruits of tomato were inoculated on water agar medium (2%; 2 g agar/100ml of sterile distilled water) and incubated at 25°C for 3 days. After incubation, single spore colony was transferred to the Petri dishes containing Potato Dextrose Agar (PDA) to obtain pure culture of the pathogen. The commercially released endophytic bacterial strains B. Subtilis (EPCO16 and EPC 5) isolated from coconut and cotton respectively and the rhizobacterial strains of P. fluorescens (Pf1, Py15 and Fp7) were obtained from the culture collection of Department of Plant Pathology, Tamil Nadu Agricultural University (TNAU), Coimbatore, India. The strains of B. Subtilis were maintained on Nutrient Agar (NA) slants and P. fluorescens strains were maintained on King’s B (KB) slants at 4°C.

Compatibility among bacterial strains

The isolates of Pseudomonas and Bacillus were tested for their compatibility among each other following the method of Fukui et al. [11]. The compatibility was determined for P. fluorescens and B. subtilis strains using NA medium. The bacterial strains were streaked
horizontally and vertically to each other. The plates were incubated at room temperature (28 ± 2°C) for 72 h and observed for the inhibition zone. Absence of inhibition zone indicates the compatibility with respective bacterial strains and the presence of inhibition zone indicated the incompatibility.

In vitro evaluation of individual and combined biocontrol agents against *A. solani*

The mycelial disc (9 mm) of the tomato early blight pathogen *Alternaria solani* was placed in the centre of the petri plate. Sterile filter paper discs (6 mm) were placed one cm away from the edge at four sides centring the fungal disc. Twenty five micro litres of bacterial broth culture (9x10⁸ cfu ml⁻¹) of each strain were dropped over the filter paper discs. Observation was taken after seven days for the presence of inhibition zone. Control was maintained with sterile distilled water instead of bacterial inoculum. The radial growth of the pathogen and per cent reduction over control was calculated by using the formula,

\[
\text{Percent reduction over control} = \frac{CT}{T} \times 100
\]

Where C−Mycelial growth of the pathogen in control (mm) and T−Mycelial growth of the pathogen in dual plate (mm).

In vitro efficacy of biocontrol agents on growth parameters of tomato seed bacterization

Seeds (One gram) of tomato cv. PKM1 surface-sterilized with 2% sodium hypochlorite for 20 seconds; rinsed in sterile distilled water and dried overnight were treated with Ten ml of biocontrol inoculums containing not less than 3x10⁸ cfu ml⁻¹. One hundred mg of Carboxy Methyl Cellulose (CMC) was added as an adhesive material. The treated seeds were kept for two hours and air-dried overnight in a sterile Petri dish and used for sowing.

Plant growth-promotion

The plant growth-promoting activity of the biocontrol agents was assessed based on the seedling vigour index by the standard roll towel method [12]. Seed bacterization was done as described above. Twenty-five treated seeds were placed on a pre-soaked germination paper. The seeds were held in position with another germination paper strip and gently pressed. The polythene sheet along with the seeds was then rolled up and incubated in a growth chamber for 10 days. Three replications for each treatment and a suitable control were also maintained. After incubation, the root length and shoot length of individual seedlings were measured and the per cent germination was calculated. The seedling vigour index was calculated using the formula, Vigour Index= (mean root length+mean shoot length) ×% germination [13].

Preparation of individual and mixture of PGPE and PGPR bioformulations

A loopful of *P. fluorescens* and *B. subtilis* were inoculated into the sterilized KB and Nutrient broth, respectively and incubated in a rotary shaker at 150 rpm for 48 h at room temperature (28 ± 2°C). After 48 h of incubation, the broth containing 9x10⁸ cfu/ml was used for the preparation of talc-based formulation. To 400 ml of bacterial suspension, 1 kg of talc powder (sterilized at 105°C for 12 h), calcium carbonate 15 g (to adjust the pH to neutral) and Carboxy Methyl Cellulose (CMC) 10g (adhesive) were mixed under sterile conditions, following the method described by [14]. After shade drying overnight, it was packed in polypropylene bag and sealed. At the time of application the population of bacteria in talc formulation was not less than 2.5-3x10⁸ cfu/g. For bacterial strain mixture, the bacterial strains were grown separately and the strains that are going to make up the mixture were added equally (v/v) and finally mixed with talc powder, CaCO₃ and CMC [14].

Efficacy of bio-formulation mixture on the incidence of *A. solani* disease under greenhouse conditions

A pot culture experiment was conducted with individual and combination of rhizosphere and endophytic bacteria treatments. Seeds of tomato cv. PKM1 were sown in earthen pots (Size-0.35m diameter, 0.50m height, volume of soil: 0.04 m³) filled with sterilized potting soil at five seeds per pot. The talc-based product was dissolved in water (20 g/l) and allowed to settle for 1 h, filtered through muslin cloth and the filtrate was used for spray. The fungicide mancozeb 0.2% spray was used as a positive control. The talc-based bioformulation mixtures were sprayed at 45 days after sowing and one day after spray, the plants were inoculated with the spore suspension (5x10⁸ spores/ml) of *A. solani*. The plants inoculated with the pathogen alone served as inoculated control. Ten days after inoculation, observations on development of early blight symptoms were made. Three replications were maintained for each treatment; each replication consisted of five pots and in each pot four plants were maintained in completely a randomized design under glasshouse conditions. The trial was repeated once and the data presented were the pooling of two glasshouse trials. The intensity of the disease was recorded in each treatment following the 0–9 scale score chart (0- Healthy; 1-1 to 5%; 3-6 to 10%; 5-11 to 25%; 7-26 to 50% and 9-51% leaf area infected) proposed by Ramakrishnan et al. [15]. The Percent Disease Index (PDI) was estimated using the formula suggested by Mckinney [16].

Statistical analysis

The data were analyzed by analysis of variance (ANOVA), and treatment means were compared by Duncan’s Multiple Range Test (DMRT) at 5% level. The data on disease severity were arcsine transformed before undergoing statistical analysis [17].

Results

Compatibility among bacterial strains

PGPR strains of *P. fluorescens* (Pf1, Py15 and Fp7) and PGPE strains of *B. subtilis* (EPCO16 and EPC 5) were tested for their compatibility in vitro. None of the antagonistic bacteria inhibited each other, suggesting that these bacterial biocontrol agents were compatible with each other.

Effect of biocontrol agents on radial mycelial growth of *A. solani*

PGPR and PGPE strains were tested individually and in combination to assess the radial growth of *A. solani*. All the treatments were effective in reducing the mycelia growth of the pathogen. However, the combined application of EPCO16+Pf1 resulted in the least mycelia growth of *A. solani*. The mycelial disc (9 mm) of the tomato early blight pathogen *Alternaria solani* was placed in the centre of the petri plate. Sterile filter paper discs (6 mm) were placed one cm away from the edge at four sides centring the fungal disc. Twenty five micro litres of bacterial broth culture (9x10⁸ cfu ml⁻¹) of each strain were dropped over the filter paper discs. Observation was taken after seven days for the presence of inhibition zone. Control was maintained with sterile distilled water instead of bacterial inoculum. The radial growth of the pathogen and per cent reduction over control was calculated by using the formula,

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effectiveness of strains and more consistent disease suppression [4,5,18]. The results of the present study provide evidence that the P. fluorescens strains (Pf1, Py15 and Fp7) and B. Subtilis strains (EPCO 16 and EPC5) were compatible and effectively inhibited the growth of A. solani. Earlier it has been reported that the bio control agents such as Trichoderma viride and P. fluorescens significantly reduced the mycelial growth, spore germination, spore production and germ tube formation of A. solani and A. alternata [19]. Several strains of Pseudomonas and Bacillus spp. have been reported to produce wide array of antibiotics viz., 2, 4, diacetylphloroglucinol, oligomycin, phenazine, pyoluteorin, pyrozin, pyocyanin, iturin, bacillomycin, zwittermycin A and surfactin which are responsible for their antifungal action [20]. Results from the present study clearly indicated maximum reduction in mycelia growth due to the combination of biocontrol strains than individual strains suggesting the synergism among biocontrol agents in reducing the mycelia growth of the pathogen.

Similarly, the treatment with combination of Pf1+EPCO16 increased the plant growth of tomato more than did individual biocontrol strains. Similar results on increased root and shoot length due to combined application of Pf1+Py15+Bx16+Zimmu in tomato [4] and chilli [5] were also reported. In the pot culture studies also the treatment with combination of Pf1+EPCO16 resulted in a significantly lower early blight disease severity than any of the strains treated individually. Earlier mixtures of PGPR strains were reported to suppress sheath blight in rice more than the individual PGPR strains [14]. Although the chemical treatment with Mancozeb (0.2%) recorded the least disease severity it is noteworthy to observe that the treatment with combination of Pf1+EPCO16 also produced almost comparable results in reducing the disease severity. Thus the eco-friendly nature of antagonistic bacterial formulations is advantageous over the use of chemical fungicides.

**Conclusion**

The results of the present study demonstrated that combined application of Pf1+EPCO16 (endophytic and rhizosphere bacterial strains) is a promising approach for the eco friendly management of early blight disease caused by A. solani and enhancing the growth of the tomato plants.

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