

Rhizospheric Pseudomonas fluorescens as rejuvenating and root proliferating agents in black pepper

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ABSTRACT: Six strains of Pseudomonas fluorescens, which had been found efficient in root-rot suppression in black pepper (Piper nigrum L.) caused by Phytophthora capsici were tested for their ability in rejuvenating Phytophthora infected black pepper cuttings. Among the treatments, which had only bacterial application, HSR-6 ensured survival of 67% of the infected cuttings where as only 22% of the untreated plants survived 3 months after treatment. The combined application of HSR-51, HSR-6+HSR-13 and HSR-51+HSR-11 with Metalaxyl -Mancozeb resulted in 100% survival of the infected plants, showing an additive effect. Experiments conducted for root-proliferation activity of the P. fluorescens showed that the bacterial strains helped increase the root biomass of the plants (30-135%). All the strains increased the root length and root area in the treated plants (12-127%, 43-200%, respectively). Strains were also found to produce plant growth hormones, viz. IAA & GA.

KEY WORDS: Black pepper, root-rot, Phytophthora capsici, Pseudomonas fluorescens, rejuvenation, root proliferation

INTRODUCTION

Black pepper is an important export oriented crop. Foot-rot of black pepper caused by Phytophthora capsici is a very serious disease, which caused severe economic loss to the farmers (Sarma, 2003). It is inevitable that fewer pesticides will be used in the future and that greater stress would be towards disease management using rhizosphere microorganisms. Fluorescent pseudomonads are some of the effective candidates for biological control of soil-borne plant pathogens owing to their mizosphere competence, plant growth promotion and disease suppression (Kloepper and Schroth, 1981).

Lynch and Whipps (1991) proved plant growth promotion by rhizobacteria due to chemical and physical stimulation of plant roots resulting in more rapid emergence, higher chlorophy! I level and increased stature. Pluorescent pseudomonads produce plant growth regulators and even suppress deleterious soil-borne pathogens (Dubeikovsky et al., 1993). Liu et al. (2000) observed longer shoots/

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roots in plants (reated with strains of Plant Growth

strains of PGPRs were tested for their effectiveness

in rejuvenating black pepper cuttings infected with

P. capsici in order to adopt the technology for

nursery management, if found feasible. Studies were

also conducted with 5 strains of rhizobacteria for

their efficacy in root proliferation in black pepper.

MATERIALS AND METHODS

different treatments in completely randomized

design (CRD). The P. fluorescens strains used for

the study were HSR-51, HSR-13, HSR-18, HSR-11,

IISR-8 and IISR-6. The strains were multiplied in

nutrient broth at 28°C for 48 h with constant agitation

(150 rpm) and bacteria were harvested by pelleting

at 7000 rpm for 10 min. The cells were resuspended

in 10 mM MgSO₄. This inoculum was used to treat

were used for the study. There were 40 treatments

altogether (Table 1), which also included a fungicide,

Ridomil Mancozeb (2.5g/L), drenched in soil @

Table 1. Treatment details

Treatment

P. fluorescens -HSR 51

P. fluorescens - HSR/H

IISR 51+ Phorate

(2g/L)+Phorate

HSR 51+ phorate

P. fluorescens - IISR 6

(2g/L)+ Phorate

HSR 6+ Phorate

Infected black pepper cuttings from a nursery

HSR 51+ Metalaxyl - Mancozeb (2g/L)

HSR 51+ Metalaxyl - Mancozeb (2g/L)

HSR 51+ Metalaxyl - Mancozeb

HSR 51+ Metalaxyl - Mancozeb

HSR 6+ Metalaxyl - Mancozeb (2g/L)

The experiments were conducted with 40

In the present study, short fisted efficient

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Treatment

(2g/L) +

HSR 6+ Metalaxyi-Mancozeb (2g/L)+

IISR 13+ Metalaxyl - Mancozeb (2g/L)

HSR 18+ Metalaxyl-Mancozeb (2g/L)

HSR 8 + Metalaxyl - Mancozeb (2g/L)

HSR 8+ Metalaxyl - Mancozeb (2g/L)+

HSR51+HSR11+Metalaxyl-Mancozeb

HSR 6+HSR 13 + Metalaxyl-Mancozeb

HSR 51+ HSR 11 + Metalaxyl -

HSR 51+ HSR 11+ Phorate

HSR 6 + HSR 13+ Phorate

Mancozeb (2g/L)+ Phorate

IISR 6 + IISR 13+ Metalaxyl -

HSR 18+ HSR 8 + Metalaxyl -

HSR 18+ HSR 8 + Metalaxyl-Mancozeb

Metalaxyl-Maneozeb (2g/L)+Phorate

HSR 18 + HSR 8 + Phorate

Metalaxyl-Mancozeb (2g/L)

HSR 13+ Metalaxyl-Mancozeb

Phorate

IISR18+ Metalaxyl-Mancozeb

P. fluorescens - IISR 13

P. fluorescens -IISR 18

P. fluorescens -IISR 8

HSR 13+ Phorate

IISR 18+ Phorate

(2g/L)+Phorate

HSR 8 + Phorate

HSR 51+HSR 11

Mancozeb (2g/L)

(2g/L)+ Phorate

HSR 6+HSR 13

HSR 18+ HSR 8

Mancozeb (2g/L)

(2g/L)+ Phorate

Control

Phorate

(2g/L)

Phorate

Promoting Rhizobacteria (PGPR),

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250ml/bag and Phorate @ 5g/bag (all standard recommended dosages). Treatments with combination of fungicide and bacterial strains also were maintained. Five replicates were maintained for each treatment with a plot size of 3 black pepper cuttings. Before planting, the roots were tested for P. capsici (by plating), Radopholus similis and Meloidogyne incognita (enumeration under microscope). The roots were thoroughly washed in running tap water to remove adhering soil and dipped in the bacterial suspension (10¹⁰ efu / ml) for 30 min. and planted in polythene bags containing 5kg of potting mixture. Then the soil was drenched with the bacterial suspension (108) cells/g of soil). The plants were watered regularly. Observations were made on the survival of the infected plants. The values obtained were mean of 5 replicates and means were separated by Duncans' Multiple Range Test.

Effect of P. fluorescens strains in root proliferation

In another experiment, five strains of P. fluorescens (HSR-51, HSR-13, HSR-11, HSR-8, and IISR-6) were evaluated in the greenhouse for root initiation and proliferation in black pepper plants. Two-node cuttings of black pepper were prepared and dipped in bacterial suspension (10⁸ cells/ml) for 30 min. These cuttings were planted in sterile coir compost and the bags were drenched with bacterial suspension (Log-8 cfu/ml). These cuttings were uprooted after 60 days and thorough examination of the root system viz. root length, total number of roots, the total area of roots and the root biomass were estimated after scanning and analysis using the software, GS-Root® (PP Systems, Winter Street, USA) and compared with that of the untreated plants. The values obtained were mean of 5 replicates and means were separated by Duncans' Multiple Range Test.

In order to find out the mechanism by which the *Pseudomonas* strains enhance the growth in black pepper, the strains were tested for the production of Indole Acetic Acid (IAA) and Gibberellic Acid (GA), which are growth-promoting hormones in plants. The assay was performed as per the protocol of Hasan (2002).

RESULTS AND DISCUSSION

P. capsici was detected in the roots of the infected plants before planting. Population of R. similis was detected (15-25/g root) in the roots and M. incognita was not detected. Observations were made on percentage survival of the infected cuttings, which was recorded over a period of 3 months. Strains of P. fluorescens could rejuvenate the infected black pepper cuttings to varying levels. After 3 months, out of the bacteria-only treatments. the plants treated with IISR-6 resulted in maximum number of surviving plants (67%), followed by IISR-13 (56%) whereas untreated set had only 22 per cent survival (Table 2). Earlier studies showed that the antagonism by these strains of P. fluorescens against P. capsici is by different modes viz. production of volatile and non-volatile inhibitory compounds including hydrogen cyanide. production of siderophores and mycolytic enzymes which reduced the proliferation of the pathogen (Diby and Sarma, 2005a & 2005b). This could also be explained as a means of systemic resistance offered by the bacterial strains to the host plant (Diby and Sarma, 2005). Metalaxyl - Mancozeb. showed an additive effect with the bacterial strains in protecting the plant. Survival of 100 per cent of the infected plants was obtained when bacterial treatment (HSR-11, HSR-51+HSR-11 and HSR-6+IISR-13) was combined with Metalaxyl-Mancozeb (Table 2). When applied alone, IISR-6 was found to be superior to other strains in rejuvenating the infected plants. The ability of certain fluorescent pseudomonads in protecting black pepper cuttings from infection by P. capsici in an artificially inoculated system has been demonstrated earlier wherein the antagonists acted as preventive agents (Jubina and Grija, 1998; Anith et al., 2002), while the present study demonstrated the ability of P. fluorescens strains as rejuvenating agents in black pepper cuttings, which had already been infected naturally.

Where combination of bacterial strains was used, there was no additive or synergistic effect. Phorate had no effect on the survival of the plant since the population of *R. similis* was only very less and the major pathogen being *P. capsici*.

Table 2. Survival of infected cuttings after three months of treatment

SLno	Treatment	Survival (%)	SI.no.	Treatment	Survival (%)
1	P. fluorescens AISR 51	14.4 bold	21	P. fluorescens -HSR 8	22.2 def
2	HSR 51+ Metalaxyl - Mancozeb (2g/L)	77.8 abc	22	HSR 8+ Metalaxyl - Mancozeb (2g/L)	88.9 ab
3	IISR 51+ Phorate	22.2 Gel	23	HSR8+ Phorate	00.0 r
4	HSR 51+ Metalaxyl - Mancozeo (2g/L)+ Phorate	33,3 edet	24	HSR 8 + Metalaxyt - Mancozeb (2g/L) + Phorate	88,9 ab
5	P fluorescens - HSR 11	22.2 edef	25	IISR 51+IISR 11	11.1 er
6	HSR (1+ Metalaxy) - Mancozeb (2g/L)	100,0 °	26	IISR 51+ IISR 11 + Metalaxyl - Mancozeb (2g/L)	100.0 *
7	HSR 11+ Phorate	22.2 cdef	27	HSR 51+ HSR 11+ Phorate	22.2 ^{cdef}
8	HSR 11+ Metalaxyl - Mancozeb (2g/L)+ Phorate	22.2 ^{eder}	28	IISR51+ IISR11+ Metalaxyl- Mancozeb (2g/L)+ Phorate	55.6 bede
9	P. fluorescens - HSR 6	66.7 ^{alkd}	29	IISR 6+IISR 13	44.4 bedef
10	HSR 6+ Metalaxyl - Mancozeb (2g/L)	88,9 ^{ab}	30	HSR 6+HSR 13 + Metalaxyl- Mancozeb (2g/L)	100.0 ¤
11	HSR 6+ Phorate	44,4 octo	31	IISR 6 + IISR 13+ Phorate	11.1 et
12	HSR o+ Meialaxyl - Mancozeb (2g/L)+ Phorate	77.8 ak	32	HSR 6 + HSR 13+ Metalaxyl - Mancozeb (2g/L)+ Phorate	44.4 beder
13	P. fluorescens – IISR 13	55.5 lxde	33	IISR 18+ IISR 8	11.1 cdef
14	HSR 13+ Metalaxyl - Mancozeb (2g/L)	66.6 ^{abcd}	34	IISR 18+ IISR 8 + Metalaxyl - Mancozeb (2g/L)	88.9 ab
15	IISR 13+ Phorate	55.5 lede	35	IISR 18 + IISR 8 + Phorate	66.7 ^{alsod}
16	HSR 13+ Metalaxyl - Mancozeb (2g/L)+ Phorate	60,7 also	36	HSR 18+ HSR 8 + Metalaxyl- Mancozeb (2g/L)+ Phorate	66.7 ^{abc}
17	P. fluorescens -IISR 18	22.2 ^{cdel}	.37	.Control	22.2 ^{cdef}
	HSR 18+ Metalaxyl - Mancozeb (2g/L)	44,4 bedet	38	Meti,laxyl-Mancozeb (2g/L)	66.7 bede
19	HSR 18+ Phorate	11.1 ef	39	Phorate	22.2 ef
	HSR 18+ Metalaxyl - Mancozeb (2g/L)+ Phorate	44,4 bedet	40	Metalaxyl-Mancozeh (2g/L)+ Phorate	44.4 beder

In the experiment for root proliferation activities, the bacterial strains significantly increased the total root biomass (30-135%) of the black pepper plants (Table 3). The number of roots have been significantly increased (82-137%) apart from increasing the root length (12-127%) and

thereby root area (43-200%). These beneficiary attributes can be corroborated with the hormonal and nutritional factors by which the rhizobacteria influence the plant (Diby and Sarma, 2005c). Plant growth regulators, *viz.* IAA and GA were produced by these strains, as detected in chromatographic

Table 3. Different root-growth parameters of plants under treatment

P. fluorescens strains	Root biomass (g)	Total root length (c/n)	Total root area (mm²)	Total number of roots
IISR-6	46.95 ah	1048.046 a	2410.088*	506.7 "
IISR-8	48.39 ah	0754.308 ^{bc}	1349,492 ^{kd}	415.9 ab
IISR-11	47.4() ^{ah}	0962.685 ^{ah}	2004.929 ab	480.94
IISR-13	33,06 tc	0518.401 ^{cd}	1155.100 ^{ed}	389.8 .b
IISR-51	59.63 ^a	0917.521 ^{ah}	1668.761 ™	490.2 4
Control	25.35°	0461.294 d	0803.061 ^d	213.9 h

studies. The TLC plate was observed in an Alpha Imager-Image Analysis system and in a UV-trans illuminator, GA showed greenish fluorescens. IAA showed violet red colour in visible light and orange in UV light. GA was detected with all the strains and IAA, with strains, IISR-6. IISR-11, IISR-13 and IISR-51. Earlier studies showed that these strains could solubilize complex forms of P in the soil thus making it available to the plant. The intake of other minerals such as N and P also was found to be more with P. fluorescens treated black pepper plants (Diby et al., 2005c). These factors not only stimulated the root for higher absorption of nutrients and minerals but also affected better root health.

Most root-promoting bacteria synthesis IAA and it has been clearly demonstrated that they stimulate the formation of lateral and adventitious roots (Barbieri and Galli, 1993). The other evidences include that of *P. putida* GR12-2 cells that produced wild type levels of IAA stimulated the formation of many short adventitious roots on mung bean cuttings and in IAA over producing mutant stimulated the formulation of even more adventitious roots than the wild type strain (Mayak *et al.*, 1997).

The results showed the effectiveness of *P. fluorescens* strains in rejuvenation of infected black pepper cuttings, which could be explored for effective nursery management. The study also proved the rhizobacteria-mediated root proliferation in black pepper.

ACKNOWLEDGEMENTS

Financial support from the Department of Biotechnology, Government of India. New Delhi is gratefully acknowledged.

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