

SCREENING OF *TRICHODERMA VIRIDE* AS A BIO-CONTROL AGENT AGAINST DIFFERENT SOIL-BORNE INFECTIOUS AND PHYTOPATHOGENIC FUNGI

Paras Shah^{1*}, Mohammad Abid¹, Alia Abbas¹, Noreen Basheer¹, Abdul Hakeem Sheikh¹, Nusrat Jabeen² and Uzma Sitara³

¹A.G. Lab of Plant Pathology and Aerobiology, Department of Botany, Federal Urdu University of Arts, Science and Technology, Gulshan-e-Iqbal Campus, Karachi-75300, Pakistan.

²Department of Microbiology, University of Karachi, Karachi, Pakistan.

³Food Quality and Safety Research Institute, Southern Zone Agriculture Research Centre, Karachi, Pakistan.

*Corresponding Author E-mail: paras.shah@yahoo.com

ABSTRACT

Phytopathogenic fungi such as *Fusarium solani*, *F. oxysporum*, *Alternaria alternata*, *Rhizoctonia solani*, *Drechslera biseptata*, *Penicillium purpurogenum* etc. were isolated from wastewater irrigated agricultural soil at the site of Malir river near Quaidabad and Lyari river near old Sabzi mandi. Soil samples were collected from the rhizosphere of various vegetable crops i.e. Ridged gourd, Round gourd, Bottle gourd, Corn, Okra, Chili, Tomato, Eggplant and Cabbage etc. Besides these fungal pathogens, bio-control agent was also sequestered from soil samples. *In-vitro*, *Trichoderma viride* was tested against all isolated infectious and pathogenic fungi of plant by dual culture technique. In present study, *T. viride* significantly suppressed the pathogens by inhibiting its mycelial growth. It showed the antagonistic activity on most common and economically important phytopathogenic fungi i.e. *F. solani* (82.91%), *F. oxysporum* (82.38%), *A. alternata* (77.19%), *R. solani* (62.25%), and *P. purpurogenum* (50.79%). It was also very effective against various infectious soil-borne fungi such as *Aspergillus fumigatus* (79.92%), *A. parasiticus* (74.49%) *Humicola* sp. (66.67%) and *A. terreus* (47.62%). With the help of present studies, it is evaluated that *T. viride* comprises the higher inhibitory response towards several soil-borne infectious and pathogenic fungi that harm economically important food crops.

KEY-WORDS: Antagonism, Soil-borne fungi, pathogens

INTRODUCTION

Several fungal and bacterial phytopathogens cause serious diseases in crops that lead to massive losses to the economy of any country. To conquer this problem farmers use various methods such as crop rotation (Ikeda *et al.* 2015), disease-resistant varieties of crops (Witek *et al.* 2016) and many other strategies. Beside these, farmers use different agrochemicals against phytopathogens to prevent crop losses (Srivastava and Sharma 2014), but these chemical pesticides that are used against various plant diseases cause serious environmental hazards. They not only pollute our land but also inhibit the growth of beneficial microorganisms of our environment, cause reproductive toxicity, respiratory disorders, and carcinogenesis in mammals and adversely affect the health of humans when they consume the affected mammals. Due to these reasons, various biological control agents such as plant materials, antagonistic microbes etc. are gaining more importance against phytopathogens.

The bio-control agents not only inhibit phytopathogens but they also act as growth promoter of plant (Sharon *et al.*, 2011). Various fungi and bacteria have been scrutinized as biological control for many years as they have antagonistic properties against several phytopathogenic microbes. Antagonistic microorganisms produce some antibiotics (Glick *et al.*, 2007) to inhibit the growth of various phytopathogens (Dilantha *et al.*, 2005). The biotic agents include numerous plant growth promoting microorganisms such as *Pseudomonas* sp. (Bakker *et al.*, 2007), *Bacillus* sp. (Jourdan *et al.*, 2009; Kloepper *et al.*, 2004), *Serratia* sp. (Press *et al.*, 1997), *Piriformospora indica* (Shores *et al.*, 2010), *Trichoderma* sp. (Koike *et al.*, 2001; Segarra *et al.*, 2009), *Penicillium simplicissimum* (Elsharkawy *et al.*, 2012), *Phoma* sp. (Sultana *et al.*, 2009), non-pathogenic *Fusarium oxysporum* (Fravel *et al.*, 2003) and Arbuscular mycorrhizal fungi (Pozo *et al.*, 2009).

Bio-control agents compete with pathogen for space and nutrition to grow. Some antagonists are also able to produce enzymes, antibiotics and toxic compounds by which they impede the growth of pathogen. In case of *Trichoderma* sp., it obstructs pathogen propagules to germinate and execute the pathogen cells by producing metabolites. It also acidify the medium to inhibit the growth of pathogen. Correspondingly, *Trichoderma* sp. can use as bio-fertilizer that it produce positive effects on plant growth by stimulating the mechanism of plant-defense. According to Vyas and Vyas (1995), *Trichoderma* spp. are very effective to control numerous phytopathogens like *Rhizoctonia solani*, *Sclerotium rolfsii*, *Pythium ultimum*, *Fusarium oxysporum* and *Macrophomina phaseolina* etc. so that it alter the use of chemical fungicides such as captan, benomyl, methyl bromide.

Trichoderma spp. are accompanied with the rhizosphere and rhizoplane of plants as they are capable to colonize the roots of plants and make symbiotic association like mycorrhizae so that they protect the roots of plant against pathogenic infection by stimulating the plant-defense mechanism as well as enhance the growth of plant (Harman *et al.*, 2004). Certain species of *Trichoderma* colonize the roots of plants on long-term basis and enter the epidermal tissues where, they secrete some compounds inside the plant body (such as non-ribosomal peptides, terpenoids, pyrones and indolic-derived compounds) that stimulate the resistance of plant against pathogens attack (Harman *et al.*, 2004; Vinale *et al.*, 2008; Contreras-Cornejo *et al.*, 2016). According to McIntyre *et al.* (2004), some antagonists change the morphology of pathogen body by coiling and form appressorium-like structures that penetrate the host body and comprise the high concentrations of osmotic solutes like glycerol. In case of *Trichoderma* sp., it attaches to the pathogen lectins by its cell-wall carbohydrates. After binding, it starts coiling around the body of pathogen and develop its appressoria to infect (Howell, 2003). The aim of the present study was to determine inhibition of various phytopathogenic soil-borne fungi via *Trichoderma viride* under *in-vitro* conditions.

MATERIALS AND METHODS

In the month of May 2018, the soil samples were collected from the rhizosphere and rhizoplane of different vegetable crops which were irrigated with wastewater at the site of Malir and Lyari River. The fungal species were isolated from soil samples by serial dilution plate method with the incubation period of 5-7 days at 28 ± 2 °C. Isolated fungal species were identified by the standard references and manual books of Barnett and Hunter (1972), Raper and Fennell (1965) and Ellis (1971; 1976). The isolated phytopathogens and bio-control agent were multiplied on PDA (Potato Dextrose Agar) plate for further study.

T. viride was selected as bio-control agent from isolated fungal species of soil samples to evaluate its antagonistic activity against different sequestered phytopathogens. The agar disc of 6mm of each pathogen from pure culture was inoculated at the periphery of the PDA plates 0.5cm away from edge of the plate, as well as same sized disc of antagonist was placed opposite to the different pathogen inoculated plates. In the same manner agar disc of test pathogens were placed near the edge of fresh PDA plate for each pathogen lonely and marked as control. Plates were inoculated for 5-7 days at 28 ± 2 °C to evaluate the interaction of antagonist with each pathogen. The interaction was determined by growing colonies of antagonist and pathogen towards each other as shown in Fig. 1. The radius of each colony with control plate were measured in cm. The inhibition percentage of pathogens by antagonist was calculated by following formula (Royse and Ries, 1977; Whips, 1987; Reddy and Hynes, 1993).

$$\text{Inhibition percentage (\%)} = \frac{R_1 - R_2}{R_1} \times 100$$

Where; R_1 was symbolized as the radius of pathogen from control plate and R_2 was the radius of treated pathogen with antagonist.

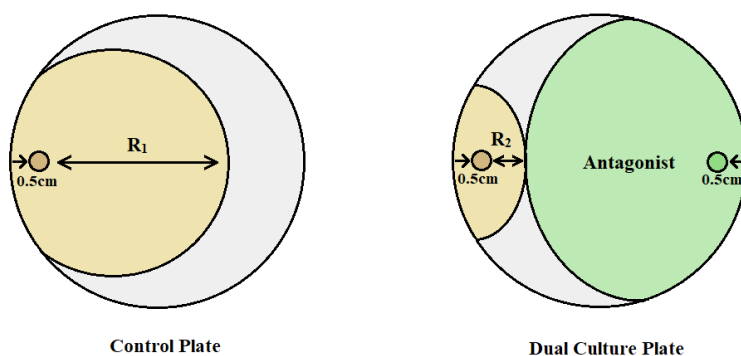


Fig. 1. R_1 indicates mycelial growth of pathogen alone as control (Control Plate) and R_2 is the inhibited growth of pathogen by antagonist (Dual Culture Plate, Skidmore and Dickinson, 1976).

Analysis of variance was performed by Minitab 17.

RESULTS

The growth of the soil borne phytopathogenic fungal species tested was significantly and differentially suppressed by the *T. viride* (Table 1). The inhibition of various fungi by *T. viride* is presented in Fig. 2 and 3. *F.*

solani and *F. oxysporum* were suppressed with the highest inhibition percentage that is $82.90 \pm 0.43\%$ and $82.38 \pm 0.48\%$, these are economically important pathogenic fungi to cause disease in many plants. *A. alternata* was controlled by $77.20 \pm 0.88\%$. *R. solani* was inhibited by $62.2 \pm 1.06\%$ of inhibition as well as colony growth of *D. biseptata* was suppressed with the $60.90 \pm 0.64\%$ of inhibition percentage by *T. viride*. It was also effective against the colony growth of *P. purpogenum* with the inhibition percentage of $50.79 \pm 1.59\%$. Besides, *P. digitatum* was controlled by the percentage of $47.62 \pm 2.28\%$.

T. viride also has antagonistic property against the colony growth of *A. parasiticus* and *A. fumigatus* that is $74.48 \pm 0.41\%$ and $79.92 \pm 0.40\%$. These are infectious soil-borne fungi for plant and animal as well that produce aflatoxins. The growth of *Humicola* sp. was controlled by the inhibition percentage of $66.67 \pm 1.93\%$ whereas *A. terreus* was suppressed by $47.62 \pm 1.19\%$.

Table 1. F- Value derived from ANOVA for inhibition percentage of different pathogenic fungi by *T. viride*.

Source	DF	SS	MS	F-Value	P-Value
Pathogenic Fungi	10	5665.74	566.574	126.99	0.001***
Error	22	98.15	4.461		
Total	32	5763.89			

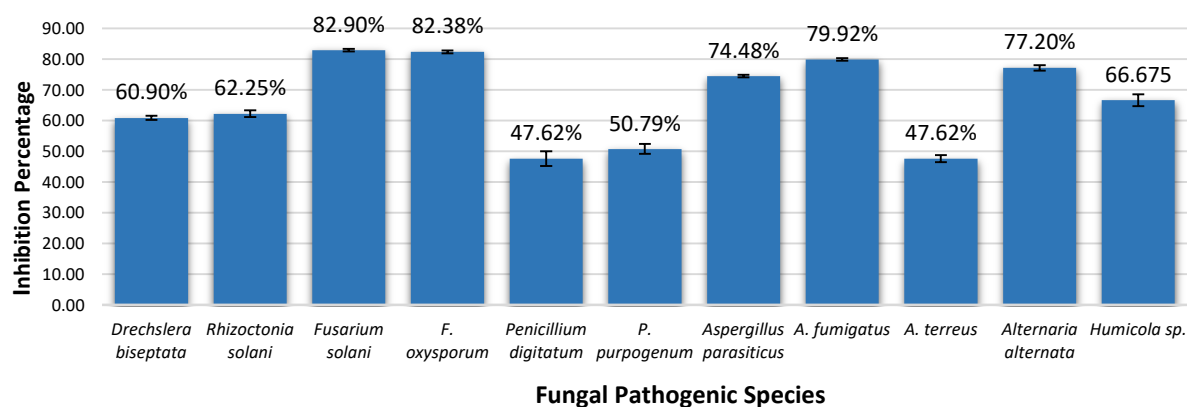


Fig 2. Inhibition percentage of various phytopathogens by *Trichoderma viride*.

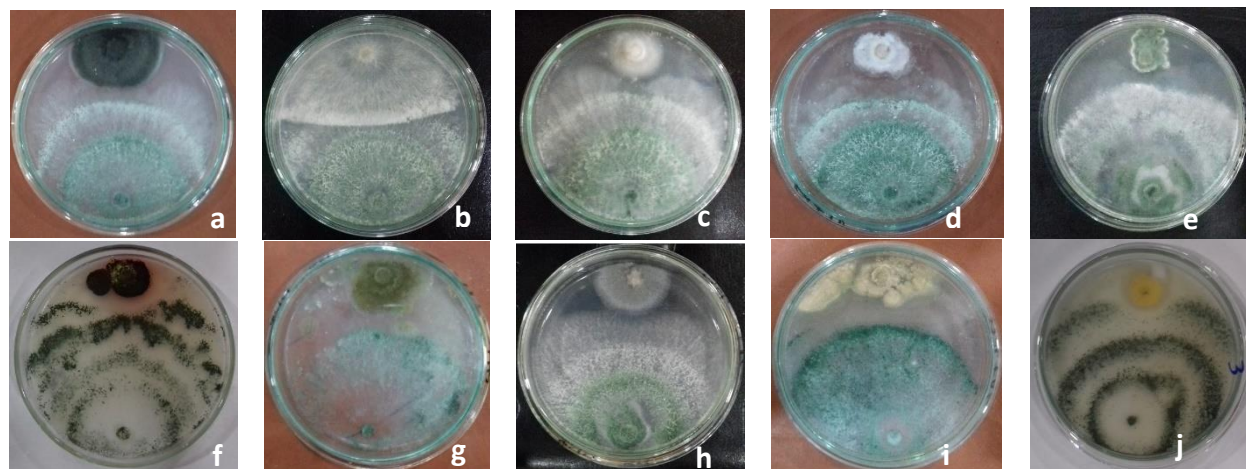


Fig. 3. Antagonistic activity of *Trichoderma viride* against (a) *Drechslera biseptata* (b) *Rhizoctonia solani* (c) *Fusarium solani* (d) *F. oxysporum* (e) *Penicillium digitatum* (f) *P. purpogenum* (g) *Aspergillus parasiticus* (h) *A. fumigatus* (i) *A. terreus* and (j) *Humicola* sp.

DISCUSSION

Almost all the species of *Trichoderma* have been considered as prospective bio-control agents as they impede the plant pathogenic soil-borne fungi that cause severe diseases in plants (Elad and Freeman, 2002; Howell, 2003;

Ziedan *et al.*, 2015). Several mechanisms that are performed by *Trichoderma* spp. to obstruct the growth of phytopathogenic fungi i.e. they compete with pathogen for nutrition and space, myco-parasitism, secretes antibiotics, stimulate the plant defense system towards pathogen (Vey *et al.*, 2001; Harman *et al.*, 2004; Shores *et al.*, 2010). This study was carried out to investigate the inhibition percentage by *T. viride* on various infectious soil-borne and phytopathogenic fungi viz. *A. alternate*, *A. fumigatus*, *A. parasiticus*, *A. terreus*, *D. biseptata*, *F. solani*, *F. oxysporum*, *Humicola* sp., *P. digitatum*, *P. purpogenum*, and *R. solani*. The present research indicated that *T. viride* has caused the highest inhibition percentage against wilt pathogen *F. solani* ($82.90 \pm 0.43\%$) and *F. oxysporum* ($82.38 \pm 0.48\%$). Almost similar result was observed by Ibrahim and Abdelaziz (2017) that *in-vivo*, *Fusarium* root rot infected plants were controlled by 81% with bio-control agent *Trichoderma* sp. Correspondingly, Gil *et al.* (2008) observed the positive results in obstructing peanut root rot caused by *F. solani* with the combination of *Trichoderma* spp. and *Gliocladium* spp. Naglot *et al.* (2015) also obtained similar results of *Trichoderma* sp. against the growth of *F. solani* as well as Sundaramoorthy and Balabaskar (2013) reported antagonistic activity of *T. harzianum* towards the growth of *F. oxysporum*. Pandey and Hussain (2006) reported *T. viride* and *T. harzianum* against *R. solani* which shows similar results as those of the present study that is inhibition around $62.25 \pm 1.06\%$. *Drechslera biseptata* was inhibited by $60.90 \pm 0.64\%$ as well as *in-vitro*, Pandey and Hussain (2010) also observed that two species of *Trichoderma* (*T. viride* and *T. harzianum*) had almost equal antagonistic activity against the growth of *D. tetramera* producing disease on capsicum.

Beside *Fusarium* spp., present study also confirmed the findings of antagonistic property of *T. viride* against *A. alternata*. Ganie *et al.* (2013) observed inhibition activity of *Trichoderma* spp. in contradiction of early blight pathogen *A. solani*. Our results are similar to Patale and Mukadam (2011) who studied the antagonistic activity of three different strains of *Trichoderma* against *A. solani*, *P. notatum*, *R. solani*, *F. oxysporum*, *A. flavus*, *A. niger* and *Phytophthora* sp. Amongst these species, *A. alternata* was inhibited by $77.20 \pm 0.88\%$, *P. digitatum* was inhibited by $47.62 \pm 2.38\%$ and *P. purpogenum* by $50.79 \pm 1.59\%$.

The cosmopolitan *Aspergillus* spp. colonize the roots of many crops such as cereals, legumes, nuts, vegetables etc. It is the opportunistic pathogen for many important crops. According to Calistru *et al.* (1997), the growth of *A. flavus* and *F. moniliforme* was suppressed by culture filtrates of four different strains of *Trichoderma* spp. The results of our study also showed inhibition of *A. fumigatus* by $79.92 \pm 0.40\%$, *A. parasiticus* by $74.48 \pm 0.41\%$ and *A. terreus* by $47.62 \pm 1.19\%$ impede by *T. viride*.

Doi and Mori (1994) have reported inhibition of fungal hyphal growth of various pathogenic fungi on culture plate, due to the volatile compounds secreting from *Trichoderma* spp. The volatile compounds secreted by *T. viride* have potential to impede the growth of wood decay fungi like *Lentinus lepidus* and *Coriolus versicolor*. Alkyl pyrones is one of the volatile compounds produced from *T. harzianum* (Fravel, 1988).

According to Elad *et al.* (1983), *Trichoderma* spp. penetrates its hypha inside pathogen, This penetration is facilitated by the enzymatic activity of *Trichoderma* sp. *R. solani* is parasitized by *T. harzianum* via producing chitinase (Benhamou and Chet, 1993).

CONCLUSION

It was found that *T. viride* has potential to impede the growth of several phytopathogenic fungi which cause serious diseases in important crops. This antagonistic isolate can be used as prospective bio-control agent in agriculture and forestry.

REFERENCES

- Bakker, P.A., C.M. Pieterse and L.C. Van Loon (2007). Induced systemic resistance by fluorescent *Pseudomonas* spp. *Phytopathol.*, 97: 239-243.
- Barnett, H. and B.B. Hunter (1972). *Illustrated Genera of Imperfect Fungi*. Burgess Publishing Co., Minneapolis, Minnesota. pp. 24.
- Benhamou, N. and I. Chet (1993). Hyphal interactions between *Trichoderma harzianum* and *Rhizoctonia solani*: ultrastructure and gold cytochemistry of the mycoparasitic process. *Phytopathol.*, 83: 1062-1062.
- Calistru, C., M. McLean and P. Berjak (1997). *In vitro* studies on the potential for biological control of *Aspergillus flavus* and *Fusarium moniliforme* by *Trichoderma* species. *Mycopathologia*, 137: 115-124.
- Contreras-Cornejo, H.A., L. Macías-Rodríguez, E. del-Val and J. Larsen (2016). Ecological functions of *Trichoderma* spp. and their secondary metabolites in the rhizosphere: interactions with plants. *FEMS microbial. ecol.*, 92: 036.
- Dilantha, W.G., S. Nakkeeran and Y. Zhang (2005). Biosynthesis of antibiotics by PGPR and its relation in biocontrol of plant diseases. *Biocontrol and Biofertilization*, pp. 67-109.

- Doi, S. and M. Mori (1994). Antifungal properties of metabolites produced by *Trichoderma* isolates from sawdust media of edible fungi against wood decay fungi. *Material und Organismen*, 28: 143–51.
- Elad, Y. and S. Freeman (2002) Biological control of fungal plant pathogens. In: *The Mycota, A Comprehensive Treatise on Fungi as Experimental Systems for Basic and Applied Research, XI. Agricultural Applications*. (Kempken F., Ed). Springer, Heidelberg, Germany, pp. 93–109.
- Elad, Y., I. Chet, P. Boyle and Y. Henis (1983). Parasitism of *Trichoderma* spp. on *Rhizoctonia solani* and *Sclerotium rolfii*-scanning electron microscopy and fluorescence microscopy. *Phytopathol.*, 73: 85-88.
- Ellis, M.B. (1971). *Dematiaceous Hyphomycetes*. Commonwealth Mycological Institute, Kew Surrey, England, pp. 608.
- Ellis, M.B. (1976). *More Dematiaceous Hyphomycetes*. Commonwealth Mycological Institute, Kew Surrey, England. pp. 507.
- Elsharkawy, M. M., M. Shimizu, H. Takahashi and M. Hyakumachi (2012). Induction of systemic resistance against Cucumber mosaic virus by *Penicillium simplicissimum* GP17-2 in Arabidopsis and tobacco. *Plant Pathol.*, 61: 964-976.
- Fravel, D., C. Olivain and C. Alabouvette (2003). *Fusarium oxysporum* and its biocontrol. *New Phytologist*, 157: 493-502.
- Fravel, D.R. (1988). Role of antibiosis in the biocontrol of plant diseases. *Annual Review of Phytopathol.*, 26: 75–91.
- Ganie, S.A., M.Y. Ghani, Q. Nissar and Shabir-u-Rehman (2013). Bioefficacy of plant extracts and biocontrol agents against *Alternaria solani*. *African Journal of Microbiol. Res.*, 7: 4397-4402.
- Gil, S.V., R. Pedelini, C. Oddino, M. Zuza, A. Marinelli and G.J. March (2008). The role of potential biocontrol agents in the management of peanut root rot in Argentina. *J. Plant Pathol.*, 35-41.
- Glick, B. R., Z. Cheng, J. Czarny, and J. Duan (2007). Promotion of plant growth by ACC deaminase-producing soil bacteria. In: *New perspectives and approaches in plant growth-promoting Rhizobacteria research*, pp. 329-339. Springer, Dordrecht.
- Harman, G.E., C.R. Howell, A. Viterbo, I. Chet and M. Lorito (2004). *Trichoderma* species-opportunistic, avirulent plant symbionts. *Nat. Rev. Microbiol.*, 2: 43.
- Howell, C. R. (2003). Mechanisms employed by *Trichoderma* species in the biological control of plant diseases: the history and evolution of current concepts. *Pl. Dis.*, 87: 4-10.
- Ibrahim, M. and A. Abdelaziz (2017). Antagonistic Fungi, Soil Amendment and Soil Solarization as an Integrated Tactics for Controlling *Fusarium* Root Rot of Lupine (*Lupinus termis*). *American J. of Microbiol. Res.*, 5: 7-14.
- Ikedu, K., S. Banno, A. Furusawa, S. Shibata, K. Nakaho and M. Fujimura (2015). Crop rotation with broccoli suppresses *Verticillium* wilt of eggplant. *J. of general pl. pathol.*, 81: 77-82.
- Jourdan, E., G. Henry, F. Duby, J. Dommès, J.P. Barthelemy, P. Thonart and M.A.R.C. Ongena (2009). Insights into the defense-related events occurring in plant cells following perception of surfactin-type lipopeptide from *Bacillus subtilis*. *Mol. Pl-Microbe Interactions*, 22: 456-468.
- Kloepper, J. W., C.M. Ryu and S. Zhang (2004). Induced systemic resistance and promotion of plant growth by *Bacillus* spp. *Phytopathol.*, 94: 1259-1266.
- Koike, N., M. Hyakumachi, K. Kageyama, S. Tsuyumu and N. Doke (2001). Induction of systemic resistance in cucumber against several diseases by plant growth-promoting fungi: lignification and superoxide generation. *Euro. J. of Pl. Pathol.*, 107: 523-533.
- McIntyre, M., J. Nielsen, J. Arnau, H. van der Brink, K. Hansen and S. Madrid (eds) (2004). *Proceedings of the 7th European Conference on Fungal Genetics*. Copenhagen, Denmark. pp. 68.
- Naglot, A., S. Goswami, I. Rahman, D.D. Shrimali, K.K. Yadav V.K. Gupta, A. J. Rabha, H. K. Gogoi and V. Veer (2015). Antagonistic potential of native *Trichoderma viride* strain against potent tea fungal pathogens in North East India. *The Pl. Pathol. J.*, 31: 278-289.
- Pandey, A. and M.A. Hussain (2006). Antagonism between *Trichoderma* spp. and *Rhizoctonia solani* on *Capsicum frutescens*. *Environ. Biol. and Conser.*, 11: 11-12.
- Pandey, A. and M.A. Hussain (2010). Comparative antagonistic profiling of different *Trichoderma* species against *Drechslera tetramera*, pathogenic to *Capsicum frutescens*. *J. Phythol.*, 2: 28–29
- Patale, S.S. and D.S. Mukadam (2011). Management of plant pathogenic fungi by using *Trichoderma* species. *Biosci. Discovery J.*, 2: 36-37.
- Pozo, M.J., A. Verhage, J. García-Andrade, J.M. García and C. Azcón-Aguilar (2009). Priming plant defence against pathogens by arbuscular mycorrhizal fungi. In *Mycorrhizas-functional processes and ecological impact*, pp. 123-135. Springer, Berlin, Heidelberg.

- Press, C.M., M. Wilson, S. Tuzun and J.W. Kloepper (1997). Salicylic acid produced by *Serratia marcescens* 90-166 is not the primary determinant of induced systemic resistance in cucumber or tobacco. *Mol. Pl-Microbe Interactions*, 10: 761-768.
- Raper, K.B. and D.I. Fennell (1965). *The genus Aspergillus*. Williams and Wilkins Company.
- Reddy, M.C. and R.K. Hynes. (1993). Relationship between In vitro growth inhibition of pathogens and suppression of pre-emergence damping-off and post emergence root rot of white bean seedlings in the green house by bacteria. *Can. J. Microbiol.*, 40: 113- 199.
- Royse, D.J. and S.M. Ries (1977). The influence of fungi isolated from peach twigs on the pathogenicity of *Cytospora cinata*. *Phytopathol.*, 63: 603-607.
- Segarra, G., S. Van der Ent, I. Trillas, and C.M.J. Pieterse (2009). MYB72, a node of convergence in induced systemic resistance triggered by a fungal and a bacterial beneficial microbe. *Pl. Biol.*, 11: 90-96.
- Sharon, E., I. Chet and Y. Spiegel (2011). *Trichoderma* as a biological control agent. In *Biol. Contr. of Pl-Parasitic Nematodes*, pp. 183-201.
- Shoresh, M., G. E. Harman and F. Mastouri (2010). Induced systemic resistance and plant responses to fungal biocontrol agents. *Annual Review of Phytopathol.*, 48: 21-43.
- Skidmore, A. M. and C.H. Dickinson (1976). Colony interactions and hyphal interference between *Septoria odorum* and phylloplane fungi. *Trans British Mycological Society*, 66: 57-64.
- Srivastava, M.P. and S. Sharma (2014). Potential of PGPR bacteria in plant disease management. *Biol. Cont. for Preventing Food Deterioration*, pp. 87-116.
- Sultana, F., M.M. Hossain, M. Kubota and M. Hyakumachi (2009). Induction of systemic resistance in *Arabidopsis thaliana* in response to a culture filtrate from a plant growth-promoting fungus, *Phoma* sp. GS8-3. *Pl. Biol.*, 11: 97-104.
- Sundaramoorthy, S. and P. Balabaskar (2013). Biocontrol efficacy of *Trichoderma* spp. against wilt of tomato caused by *Fusarium oxysporum* f. sp. *lycopersici*. *J. App. Biol. & Biotechnol.* 1: 36-40.
- Vey, A., R.E. Hoagland and T.M. Butt (2001). Toxic metabolites of fungal biocontrol agents. In: Butt TM, Jackson C, Magan N (eds) *Fungi as biocontrol agents: Progress, problems and potential*. CAB International, Bristol, pp. 311-346
- Vinale, F., K. Sivasithamparam, E.L. Ghisalberti, R. Marra, M.J. Barbetti, H. Li , S.L. Woo and M. Lorito (2008). A novel role for *Trichoderma* secondary metabolites in the interactions with plants. *Physiol. and Mol. Pl. Pathol.*, 72: 80-86.
- Vyas, S.C. and S. Vyas (1995). Integrated Control of Dry Root of Soybean. In: *Modern Fungicides and Antifungal Compounds*, Lyr, H., P.E. Russel and H.D. Sisler (Eds.). Intercept Ltd., Andover, pp: 562-572.
- Whips, J.M. (1987). Effect of media on growth and interactions between a range of soil- borne glass house pathogens and antagonistic fungi. *New Phytol.*, 107: 127-142.
- Witek, K., F. Jupe, A.I. Witek, D. Baker, M.D. Clark and J.D. Jones (2016). Accelerated cloning of a potato late blight-resistance gene using RenSeq and SMRT sequencing. *Nat. Biotechnol.*, 34: 656.
- Ziedan, E.S.H.E., E.S.H. Farrag and A.F. Sahab (2015). Effect of *Trichoderma harzianum* against *Thielaviopsis paradoxa* and their pathological potential on date palm seedlings. *Int. J. of Agricul. Technol.*, 11: 913-923.

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