INTEGRATED MANAGEMENT OF MAJOR FUNGAL DISEASES OF TOMATO

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ABSTRACT

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The experiment was conducted at Bangladesh Agricultural University, Mymensingh during the period from October 2004 to March 2005 to determine an eco-friendly management practices against major fungal diseases of tomato. The treatments were: $T_1 = BAU$ - Biofungicide + Sanitation + Neem (3), T_2 = MOC (Mustard oil cake) + Neem (2) + Karmacha, $T_3 = BAU$ -Biofungicide + Neem + Karmacha (2), $T_4 = BAU$ -Biofungicide + Karmacha (2) + Mahogony, $T_5 = BAU$ -Biofungicide + MOC + Neem + Karmacha + Mahogony, $T_6 = MOC$ + Karmacha + Mahogony (2), $T_7 = BAU$ - Biofungicide + MOC + Neem + Mahogony (2), $T_8 = MOC$ + Sanitation + Neem (3), $T_9 = BAU$ - Biofungicide + MOC + Neem + Mahogony + Sanitation and $T_{10} =$ control. In case of late blight treatment T_7 gave the lowest value but it showed statistically insignificant with rest treatments except T_{10} . Regarding early blight T_6 , T_7 , T_8 and T_9 exhibited more or less equally effective against the disease and they were statistically similar. As high as 33% wilt infection was recorded in T_{10} while no wilt infection was detected in all the rest treatments.

Key words: Tomato, BAU-Biofungicide, Fungal diseases

INTRODUCTION

Tomato (*Lycopersicum esculentum* Mill) is the most popular vegetable in the world because of its taste, colour and high nutritive value and also for its diversified use (Bose and Som, 1986). In Bangladesh the average yield of tomato is 2.71 metric tons per acre (B.B.S, 2004) which is lamentably low as compared to the other leading tomato producing countries (FAO, 1999). There are many factors involved in such low yield of tomato in Bangladesh; among them are infestations by fungi, bacteria, nematodes or viruses and the competing weeds are predominant (Villaral, 1980).

Over 200 diseases have been reported to affect the tomato plants in the world (Watterson, 1986). Among the fungal diseases early blight (Alternaria solani), late blight (Phytophthora infestans) and fusarial wilt (Fusarium oxysporum) are major. Both late and early blight can be effectively controlled by using fungicides but it is costly as well as not easily available to farmers' door. Wilt control has been restricted to use of wilt resistant cultivars, grafting on wilt resistant root stalk, crop rotation, deep ploughing of land and also use of different soil amendments. Removal of infected plants from the field will help limiting the disease spread. Considering the above points the most urgent need is to develop varieties of tomato that can resist the ravage of important fungal disease like early blight, late blight and wilt. But none of the cultivated tomato varieties in the country are found to be horizontally resistant to these diseases. Therefore, the general control of disease by employing Integrated Disease Management (IDM) program has drawn special attention to the researchers. It can reduce the cost of healthy cropping and the farmers can easily apply them in the field. The IDM practices not only save the crop from the referred field diseases but also reduce the possibility of attack by the other pathogens (fungi, viruses, bacteria and nematodes) to tomato crop in a cropping season. There is a great need to carry out farmer level research aiming to develop a holistic disease management model to manage the major diseases of tomato. In these circumstances, the present study has been undertaken to develop an eco-friendly management practices against major fungal diseases of tomato.

MATERIALS AND METHODS

The experiment was conducted at Bangladesh Agricultural University, Mymensingh during the period from October 2004 to March 2005. Seeds of tomato variety, Oxball (susceptible to diseases) were sown in seedbed on 20 October, 2004. Cow dung 10 tons per ha was applied and no chemical fertilizer was used in this experiment. The unit plot size was 1m x 1m. Row to row and plant to plant spacing was 50 cm. The experiment was laid out in the Randomized Complete Block Design (RCBD) having three replications. Distance between the blocks was 1m and between the plots was 0.5m. Apparently healthy seedlings of 35 days old were transplanted in the experimental field. There were 10 treatments as follows:

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 $\begin{array}{l} T_1 = BAU- \ Biofungicide + Sanitation + Neem (3)\\ T_2 = MOC \ (Mustard oil cake) + Neem \ (2) + Karamcha\\ T_3 = BAU- \ Biofungicide + Neem \ + Karamcha \ (2)\\ T_4 = BAU- \ Biofungicide + Karamcha \ (2) + Mahogony\\ T_5 = BAU- \ Biofungicide \ + MOC \ + Neem \ + Karamcha \ + Mahogony\\ T_6 = MOC \ + Karamcha \ + Mahogony \ (2)\\ T_7 = BAU- \ Biofungicide \ + MOC \ + Neem \ + Mahogony \ (2)\\ T_8 = MOC \ + Sanitation \ + Neem \ (3)\\ T_9 = BAU- \ Biofungicide \ + MOC \ + Karamcha \ + Mahogony \ + Sanitation\\ T_{10} = Control \end{array}$

The BAU-Biofungicide was added in the assigned pits @ 50g/pit and mixed well. MOC was decomposed in water for seven days after well grinding. After decomposing, it was diluted by adding plain water and applied @ 50g per plant in ring placement in soil around the base of the seedling, after 30 days of transplanting of seedlings. Sanitation was done 2 times after 30 days and 60 days of transplanting. The diseased leaves, which were 25% infected or more were removed. The dried and dead leaves were also removed from the plot. Plant extract was prepared as suggested by Sharmin (2003).

Neem (*Azadirachta indica*) Mahogony (*Swietenia Mahogony*)) and Karamcha (*Carissa carandas*) extract were applied @ 2g/l at 15days interval. First spray was given ten days after transplanting (DAT). Intercultural operations were done as and when necessasary. Data were taken on late blight infected plant, disease severity of late and early blight (0-6 scale for late blight and 0-5 scale for early blight, Vakalounakis, 1983) and wilted plants. Data were taken at 25, 40, 55 and 70 DAT. Percent data were transformed following Arcsine transformation.

RESULTS AND DISCUSSION

Late blight

Significant variation among the treatments becomes evident on percent late blighted plants regardless of data recording after days after transplanting (DAT) of tomato except 40 DAT. The late blight infected plants ranged 11.11-40.79%. At 25 DAT the treatment T_7 (Bio-fungicide+MOC+Neem+Mehogoni-2) appeared best one in reducing late blight infected plants and showed significantly better compared to rest treatments except T6 (MOC+Karamcha+Mehogoni-2), T_8 (MOC+Sanitatin+Neam-3) and T_9 (Bio-fungicide + MOC + Karamcha + Mehogoni + Sanitation). (Table 3). Significantly higher late blight infected plants was recorded by T_{10} (Control) and it differed significantly with all the rest treatments. The effect of treatments on late blight infection was insignificant at 40 DAT. At 55 DAT late blight infected plants ranged 48.15-81.48%. Although treatment T_7 gave the lowest late blight infection numerically but it showed statistically similar to all the treatments expect T_{10} . More than 92% tomato plants became infected plant was recorded by T_7 and it showed statistically similar T_5 (Bio-fungicide + MOC + Neem + Karamcha + Mehogoni) and T_9 .

While disease severity of late blight was considered, there were no significant variation among the treatments whatever they were assessed at 25, 40, 55 and 70 DAT (Table 2). Late blight infection with lower disease severity prevailed at 25 DAT and it increased gradually with increasing of plant age.

Early blight

Significant variation among the treatments became well pronounced in controlling early blight of tomato in all four observations. It was evident that at 25 DAP, more than 59 percent plant became infected due to the disease in control treatment (T_{10}). The lowest early blighted plants of 18.51% were recorded in T_7 and it showed statistically insignificant with only T_6 (MOC+Karamcha+Mehogoni), T_8 and T_9 . The treatment, T_7 also proved its affectivity on observations at 40, 55 and 70 DAT by exhibiting the lowest plant infection due to early blight. While statistical analysis was performed, T_7 gave significantly lower infected plants followed by T_9 , T_8 and T_6 and they were statistically similar in all four times of observations.

Fusarium wilt

The results showed that all the tested treatments effectively controlled the wilt disease where none of plants died due to the disease except T_{10} (Control). In control more than 33% plant wilted during the entire growth period, most of which happened within 40 days of transplanting. Result revealed that integration of treatment combinations (T_1 to T_9) efficiently suppressed the causal agent of tomato wilt in the experimental field (Table 1). The results of present investigation indicate that the incidence of late blight infected plants and disease

severity due to *Phytophthora* infection were rather low at the period between 25 and 40 DAT. The occurrence of late blight attained an *epiphytotic* momentum when the plants interred into their reproductive phase that means between 40 and 55 DAT. This may be happened due to congenial environmental condition of the fungus. During this period the minimum and the maximum air temperature were 12.49° C and 22.72° C respectively, cocepled with more than 80% relative humidity. This was in accordance with Dey et al (1998) who worked with late blight of potato. The treatment T_7 appeared the best against late blight where Bio-fungicide (*T. harzianum*) integrated with MOC, Neem and Mehogoni but results were not, so much encouraging. This was close agreement with the findings of Slusarski and Pieter (2003) and Dey (2004). Dey (2004) screened a good number of antagonists including T. harzianum and T. viride against late blight under artificial inoculation of P. infestans in net house and concluded that the antagonists have the ability to reduce the late blight infection as prophylactic, not a curative. Integration of treatments with sanitation had some positive influence against late blight which is corroborate with the findings of Cohen (1987), Tumwine (1990), Begum (2001) and Islam (2002). But all of then suggested that sanitation with fungicide spray is more effective in controlling late blight of tomato. Regarding early blight T7 also exhibited better in controlling the disease compared to other treatment combinations. The effectiveness of Trichoderma against Alternaria spp. has been reported by Slusarski and Pieter (2003). Under the study all the integrating treatments (T_1 to T_9) performed excellent against wilt (Fusariam oxysporum) disease of tomato. The findings of the present study clearly supported those obtained by many researchers throughout the world (Ehteshmul et al. 1990, Parveen and Ghaffar, 1995; Mukherjee et al. 1995; Raj and Kapoor, 1996; Hossain and Fakir, 2001; Banu, 2003 and Dey, 2004) who worked on the biocontrol potentiality of different species of Trichoderma both in vitro and in vivo against wide range of soilborne pathogens including Fusarium oxysporum Elad et al. (1982) claimed that T. harzianum excreted 1.3glucanase and chitinase that showed antagonistic activity to control soil-borne pathogens. The affectivity of Mustard oil cake became reflected against. F. oxyporum under the present study which was in line with the findings Raj and kapoor (1996).

Treatments	% Wilt (up to 70 DAT)		
T_1	0		
T_2	0		
T ₃	0		
T_4	0		
T ₅	0		
T_6	0		
T_7	0		
T_8	0		
T ₉	0		
T_{10}	33.33		

Table 1 Effect of treatments on % Wilt during the growth period under field condition

 $T_1 = BAU$ - Bio-fungicide +Sanitation + Neem (3)

 T_2 = Mustard oil cake + Neem (2) + Karamcha

 T_3 = BAU- Bio-fungicide + Neem + Karamcha (2)

 $T_4 = BAU$ - Bio-fungicide +Karamcha (2) + Mehogoni

 $T_5 = BAU \text{-} Bio\text{-} fungicide + Mustard oil cake + Neem + Karamcha + Mehogoni$

 T_6 = Mustard oil cake + Karamcha + Mehogoni (2)

 $T_7 = BAU$ - Bio-fungicide + Mustard oil cake + Neem + Mehogoni (2)

 T_8 = Mustard oil cake + Sanitation +Neem (3)

 $T_9 = BAU \text{-} Bio\text{-} fungicide + Mustard oil cake + Karamcha + Mehogoni + Sanitation}$

 $T_{10} = Control \\$

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Treatments	% Early blighted plants				
	Days after transplanting (DAT)				
	25	40	55	70	
т	44.443 b	59.260 b	62.963 b	66.670 b	
T_1	(41.78)	(50.30)	(52.48)	(54.70)	
т	40.737 bc	55.560 bc	59.263 bc	70.373 ab	
T_2	(39.64)	(48.16)	(50.30)	(56.98)	
т	40.737 bc	59.263 b	62.967 b	70.373 ab	
T ₃	(39.64)	(50.30)	(52.48)	(56.98)	
т	40.737 bc	51.853 bc	55.557 bc	62.967 bc	
T_4	(39.64)	(46.03)	(48.16)	(52.48)	
т	37.033 bcd	48.147 bcd	55.557 bc	62.967 bc	
T ₅	(37.47)	(43.91)	(48.16)	(52.48)	
т	25.923 de	40.740 cde	44.443 cd	55.557 bc	
T_6	(30.59)	(39.64)	(41.78)	(48.16)	
т	18.517 e	25.930 e	29.627 d	33.330 d	
T_7	(25.48)	(30.59)	(32.96)	(35.24)	
т	29.627 cde	40.737 cde	44.443 cd	48.147 cd	
T_8	(32.96)	(39.64)	(41.78)	(43.91)	
т	25.923 de	33.330 de	37.033 d	37.033 d	
T ₉	(30.59)	(35.24)	(37.47)	(37.47)	
т	59.263 a	77.780 a	81.483 a	85.187 a	
T ₁₀	(50.30)	(61.82)	(64.45)	(67.29)	
evel of significance (p=0.05)	**	**	**	**	

Table 2 Effect of treatments on the occurrence of early blight infection at different growing periods under field condition

*Figures in parenthesis indicate the transformed value, ** =Significant at 1% level.

Table 3 Effect of treatments on the occurrence of late blight infection at different growing periods under fie	ld
condition	

	% late blighted plants				
Treatments	Days after transplanting (DAT)				
	25	40	55	70	
T ₁	33.330 b	37.033	70.373 ab	74.077 b	
	(35.24)	(37.47)	(56.98)	(59.34)	
т	29.627 bc	33.330	66.670 ab	74.077 b	
T_2	(32.96)	(35.24)	(54.70)	(59.34)	
T ₃	29.627 bc	37.033	66.670 ab	74.077 b	
13	(32.96)	(37.47)	(54.70)	(59.34)	
T_4	29.627 bc	33.330	66.670 ab	66.670 bc	
14	(32.96)	(35.24)	(54.70)	(54.70)	
T_5	25.923 bcd	29.627	55.557 b	59.263 bcd	
15	(30.59)	(32.96)	(48.16)	(50.30)	
т	14.813 de	25.923	55.557 b	66.670 bc	
T ₆	(22.63)	(30.59)	(48.16)	(54.70)	
т	11.110 e	18.517	48.150 b	44.443 d	
T ₇	(19.46)	(25.48)	(43.91)	(41.78)	
т	18.517 cde	25.923	62.967 ab	62.967 bc	
T ₈	(25.48)	(30.59)	(52.48)	(52.48)	
т	14.813 de	22.220	48.147 b	55.557 cd	
T ₉	(22.63)	(28.11)	(43.91)	(48.16)	
т	40.737 a	48.147	81.483 a	92.593 a	
T ₁₀	(39.64)	(43.91)	(64.45)	(74.11)	
Level of hificance (0.05)	**	NS	*	**	

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significance (0.05) *Figures in parenthesis indicate the transformed value, ** =Significant at 1% level.

* = Significant at 5% level, NS = Non significant

	Disease severity % Leaf spot/leaf blighted symptom bearing leaves				
Treatments					
		Infected			
	Days After Transplanting (DAT)				
	25	40	55	70	
т. 	22.220	33.330	59.263	92.593	
T_1	(28.11)	(35.24)	(50.30)	(74.11)	
т	33.330	48.147	48.147	81.483	
T_2	(35.24)	(43.91)	(43.91)	(64.45)	
T	22.220	37.033	51.853	85.187	
T_3	(28.11)	(37.47)	(46.03)	(67.29)	
т	22.220	37.037	44.447	77.777	
T_4	(28.11)	(37.47)	(41.78)	(61.82)	
т	22.220	37.037	37.037	70.370	
T ₅	(28.11)	(37.47)	(37.47)	(56.98)	
т	18.517	33.330	62.967	96.297	
T_6	(25.48)	(35.24)	(52.48)	(78.76)	
т	11.110	25.923	40.740	74.077	
T_7	(19.46)	(30.59)	(39.64)	(59.34)	
т	18.517	29.627	44.447	77.777	
T_8	(25.48)	(32.96)	(41.78)	(61.82)	
T ₉	14.813	25.923	40.740	74.077	
	(22.63)	(30.59)	(39.64)	(59.34)	
T ₁₀	25.923	55.557	74.077	97.407	
	(30.59)	(48.16)	(59.34)	(80.19)	

Table 4 Effect of %leaf infection/plant due to late blight under different treatments as different days after transplanting in the field

*Figures in parenthesis indicate the transformed value

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