ACTIVITY OF CERTAIN MICROORGANISMS ON ALLEVIATING THE BLIGHT DISEASE (Stemphylium vesicarium) OF ONION PLANT

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ABSTRACT: Onion (Alluim cepa L.) is a principal component of the Egyptian table. Stemphylium vesicarium causes tip necrosis and large purple blotch on leaves of onion (Blight disease) which reduce the yield quality. Biologists oriented their efforts to control the disease biologically to limit the manipulation of chemical pesticides. This investigation uses some microorganisms such as the mixtures of Serratia + Bacillus and Serratia + Pseudomonas, as well as the metabolites of Anabaena oryzae + Nostoc muscorum, Azolla pinnata, and Pleurotus columbinus. The most effective biological treatments were the combination between Serratia and Bacillus followed by Pleurotus filtrate which diminished the disease severity to an extent comparable to the chemical pesticide. High content of phenolic compounds was found in Pleurotus filtrate. Cyanobacteria followed by Pleurotus filtrates were the auspicious treatments in respect to the total yield, marketable yield and bulbs weights of onion plants being 16.56 ton/fed., 14.18 ton/fed. and 104 g for cyanobacterial filtrate treatment and 16.43 ton/fed., 13.95 ton/fed. and 100.33 g for Pleurotus filtrate treatment compared to the control (14.7, 12.86 and 96.0 in the second season, respectively).

Key words: Allium cepa L., biocontrol, Stemphylium vesicarium, Bacillus, Serratia, Pseudomonas, Anabaena, Nostoc, azolla, Pleurotus.

INTRODUCTION

In Egypt, onion (Allium cepa L.) is the main important and oldest vegetable crop. The Egyptian onion is famous allover the world because of its early appearance and superior quality. Onion is cultivated mainly for food, but also for traditional medicine. It is attacked by many diseases, which vary according to region, season and variety. production. Diseases can affect the harvesting, processing and marketing, which reduce the quality and yield, and thereby increase the cost of production, and the export potential. Tomaz and Lima (1988) reported that purple blotch and Stemphylium blight are the most important diseases in Northern India, which causes considerable losses in seed crops as well as bulb crops. It can cause severe damage, especially to the onion seed crop and losses of about 80 to 85% of the crop by affecting leaves and seed stalk. Hussein et al. (2007) stated that onion plants (Allium cepa) cv. Giza 6 grown

in several commercial fields in Upper Egypt (Assiut Governorate) exhibited symptoms of blight on the leaves and seed-stalks. Initial symptoms on leaves consisted of tip necrosis followed by small white and/or large purple spots. Such observed symptoms resembled Stemphylium blight symptoms on onion leaves, which are caused by S. vesicarium (Wallr.) Simmons. The disease is wide spread in Asia and Europe and has been recorded previously on onion plants in South Africa and in New York, Many attempts were carried out for controlling Stemphylium blight on onion using cultural practices and chemical control (Aveling and Snyman, 1993). Certain chemicals, such as salicylic acid (SA) and dichloroisonicotinic acid (INA), potassium salts, amino butyric acid (BABA) and Bion were reported to induce systemic acquired resistance (SAR) in plants (Oostendorp et 2001). **Biologists** found microorganisms play an effective role in

controlling and reducing the disease impact. Nalisha et al. (2006) and Okay et al. (2013) noticed that Bacillus subtilis produces antifungal substances which have inhibitory effects on a wide range of fungi and Serratia has chitinase which degrade the chitin composing the cell wall of fungi. Beside , cyanobacteria such as nitrogen fixation Anabaena and Nostoc isolated from various aguatic and terrestrial habitats showed antimicrobial activity (Yadav et al., 2012). Wang and Ng (2004) reported that some antifungal peptides were produced from fruiting bodies of the edible mushroom like Pleurotus eryngii. Some microorganisms were used by Hussein et al. (2007) as bioagents to inhibit the growth of Stemphylium vesicarium mycelia, those Bacillus subtilis, Pseudomonas fluorescens, Trichoderma harzianum, Gliocladium sp. and Saccharomyces cerevisiae.

The present work aims to throw the light on the biological control of *Stemphylium* blight on onion by some cyanobacterial filtrates besides some other bacterial strains compared to chemical pesticide.

MATERIALS AND METHODS

Two field experiments were conducted on onion crop (*Allium cepa* cv, Giza 20) at Nubaria Agricultural Research Station Farm, during 2009/2010 and 2010/2011 winter seasons. Soil samples, from the experimental site, were determined for pertinent characters (Page *et al.*, 1982). Results of the analyses are presented in Table (1).

Table (1): Mechanical and chemical properties of the experimental soil

	Soil Characters	Value				
Mechanical analysis	Sand %	58.57				
	Silt %	24.84				
	Clay %	16.59				
	Soil textural class	Sandy loam				
Organic matter (%)		0.19				
	CaCO3 (meq/l)	22.43				
Soil pH (1:2.5 soil/water susp.)		8.3				
EC (dS/m) (1:2.5 soil/ water extr.)		2.17				
	Ca ⁺²	6.69				
Soluble cation (meq/l)	Mg ⁺²	1.73				
	K⁺	4.29				
	Na [⁺]	10.37				
Soluble anions Soluble cations (meq/l)	CO ⁻²	-				
	HCO ₃	5.71				
	Cl	11.72				
	SO ₄ -2	5.57				
Soluble micronutrient (meq/l)	Zn	1.14				
	Fe	1.37				
	, Mn	1.09				
	Cu	0.67				

Onion seedlings were transplanted on December 23th, 2009 and December 21st,2010. The experimental plot area was 7 m²(1/600 fed.), each plot consisted of four ridges, 50 cm wide and 3.5 m long. Seedlings were planted on both sides of the ridges and spaced at 10 cm. The recommended doses of phosphorus fertilizer 45 kg P₂O₅/fed. as superphosphate and 100 kg K₂SO₄/fed. as potassium fertilizer were applied during soil preparation for all plots. Nitrogen fertilizer (90 kg N/fed.) as ammonium nitrate was added in case of control, monthly in three equal doses after All other cultural practices transplanting. including irrigation and control of weed, pathogen diseases and pests of onion production in the calcareous soils at Nubaria region were applied.

1. Chemical pesticide:

Traditional (commercial) name: Mitor Cu 50% WP.

2. Cvanobacteria and growth conditions:

Cyanobacterial species, namely strains of Anabaena oryzae, Nostoc muscorum, and Azolla pinnata were obtained from the Microbiology Department, Soils, Water and Environ. Res. Inst., (ARC), Giza. The strains were maintained in BG11 medium (Rippka et al., 1979). Cultures were incubated in a growth chamber under continuous illumination (2000 lux) and temperature of 25 °C ± 2 °C. Azolla pinnata was grown on modified Yoshida medium (Yoshida et al., 1976).

3. Preparation of algal culture and Azolla pinnata aqueous filtrates:

After 30 days of incubation, each algal biomass, in its medium, was put in an electric mixer, then the mixture of biomass with its medium was filtered and the extracted free cells were kept at 4°C till field application. Azolla pinnata was harvested from the culture medium and mixed well with distilled water and diluted by 1:1 (w/v) using an electric mixer, then filtered to obtain the extracted free cells and kept at 4 °C till field applications.

4. Bioagent sources and growth conditions:

Bacillus subitlis, Pseudomonas fluorescence and Serratia sp. were obtained from the Microbiology Department, Soils Water and Environ. Res. Inst., ARC, Giza. The bacterial strains were grown on King Medium (King et al., 1954), gently shaked on a rotary shaker incubator at 30 +/- 2 °C up to reach the log phase (ca. 10⁷cfu ml⁻¹).

5. Pleurotus columbinus source and growth condition:

The white rot fungus (Pleurotus columbinus) was obtained from the Unit of Production , Faculty Mushroom Agriculture Shams Ain University .Pleurotus columbinus was grown on potato dextrose medium, in a rotary shaker at 200 rpm for 7 days (Martin, 1950) . After propagation, the growth and its medium was put in an electric mixer, then filtered and kept at 4 °C till field application.

A randomized complete block design (RCBD) with four replicates was used as an experimental layout. The allocated treatments were as follows:

- T1: Control (without chemical or biological treatment).
- T2: Chemical pesticide.
- T3: Mixture of Anabaena oryzae and Nostoc muscorum filtrate.
- T4: Azolla pinnata filtrate.
- T5: Mixture of Serratia sp and Bacillus subtilis .
- T6: Mixture of Serratia sp and Psceudomonas fluorescens.
- T7: Pleurotus colombinus filtrate.

parameters were recorded :

1- Disease severity:

The first reading of infection density was taken after 75 days of transplanting, the second reading was after 119 days of transplanting then the third reading was after 140 days of transplanting. The infection density was estimated microscopically. Ten plants were taken from each plot to calculate disease severity according to Tarabulsi et al (1998), applying the following equation:

% Ds = Σ (n × r)×100/5N where: n= number of plants in each category. r= numerical number of each category. N= total number of examined plants.

Plants were rated for disease severity (Ds), as follows:

0 = no infection; 1= very weak infection; 2 = weak infection (tiny necrotic lesion; 3 = moderate infection (medium size lesions with corky tissue); 4 = severe infection (well developed large necrotic lesions and 5 = very severe infection (complete death of the leaf plant).

2-Vegetative growth characters:

After 120 days from transplanting, 15 plants were selected from each experimental plot for measuring plant height and number of tubular leaves per plant.

3- Days of maturity:

Number of days from transplanting to maturity was recorded. Maturity was determined by both neck softening and 50 % top down.

4- Yield and its components:

At harvest, all plants in plot were uprooted and determined for:

- a- Total yield (t/fed.).
- b- Marketable yield (t/fed.): represents the weight of single bulbs.
- c- Average bulb weight (g).

5- Bulb quality traits encompassing:

- a- Percentage of single, double bulbs and bolters: It was estimated by dividing number of single, double bulbs and blotters by the total number of bulbs per plot x 100.
- b- Percentage of total soluble solids (TSS %) (that include mainly soluble sugars , amino acids and soluble minerals): It was determined using the hand refractometer .

Nitrogen, phosphorus and potassium were determined according to Jakson (1976). Total phenols were determined spectrophotometrically as described by Swain and Hillis (1959). Total carbohydrates were determined by the method of Dubois *et al.*(1956).

Soil biological activity: CO₂ evolution was determined according to Gaur *et al.* (1971).

Total bacterial counts were estimated on nutrient agar using the spread plate method (Allen,1956).

Statistical analysis:

A combined analysis for the two seasons was calculated by MSTAT program. Combined average values from the four replicates of each treatment were interpreted using the analysis of variance (ANOVA).

RESULTS AND DISSCUSION

Fig. (1) reveals the effect of different biological treatments, compared to chemical pesticide, on disease severity of onion plant that infested by Stemphylium vesicarium throughout two agricultural seasons. The chemical pesticide was the best treatment in lowering the disease severity for the three readings in both seasons. All biological treatments had positive effects on disease severity, compared to the control but, at lower extent, compared to the chemical pesticide. The treatments of Serratia plus Bacillus followed by Pleurotus filtrate were the best in diminishing the disease severity comparable to the chemical pesticide in the second season among the biological treatments. Badalyan (2008) found that some species of Pleurotus had antagonistic effect and excrete antifungal substances against phytopathogens like Fusarium and Trichoderma. Ghasemi et al (2010)confirmed that chitinase isolated from Bacillus pumilus had antifungal activity and inhibited all tested fungi like Rhizoctonia solani, Verticillium sp., Nigrospora sp., Stemphyllium botryosum and Bipolaris sp. Belal et al . (2013) found that Bacillus subtilis produced antimicrobial metabolites against two fungal and one bacterial strains; Pseudoperonospora cubensis, Sphaerotheca fuliginae and Staphylococcus aureus that infect cucumber. antimicrobial metabolites mostly included acids and made collapse sporangiophor and sporangia of P. cubensis and conidiophore and conidia of S. fuliginae and also had antagonistic activity against Staphylococcus aureus . Nandakumar et al. (2002) reported that some strains of Pseudomonas fluorescens exudated some

substances like 2,4-diacetyl phloroglucinol, iron chelating siderophore, hydrogen cyanide, lytic enzymes, such as chitinase and β -1,3-glucanase in cultures acting as antibiotics against the rice sheath blight pathogen *Rhizoctonia solani*.

Fig. (2) Shows slight differences in total bacterial counts in rhizosphere regions of the infected onion plants by *Stemphylium vesicarium* between the two seasons, in the second season counts were higher than the first. The chemical pesticide recorded the lowest bacterial numbers, compared to the biological treatments which were higher than both of the control and pesticide treatments. *Serratia* plus *Bacillus* treatment was the superior among the biological treatments, followed by *Anabaena* plus *Nostoc* filtrate treatment, while *Serratia* plus *Pseudomonas* and *Pleurotus* filtrate treatments were the lowest.

Fig. (3) Exhibits variations among all treatments in respect to the rate of CO_2 evolution in the rhizosphere of onion plants infected by *Stemphylium vesicarium*. The chemical pesticide treatment was the lowest among all treatments, due to its mortal action on soil microflora.

Cyanobacterial filtrate followed by Pleurotus filtrate were the uppermost treatments for CO₂ evolution among biological treatments. This might attributed to excreting some regulating substances having the ability to control the fungal disease and promote the biological activity in the soil. Unexpectedly, the control treatment recorded high value of CO₂ rate, due to the biological activity of microflora in the plant rhizosphere. The importance of the evolved CO2 estimation was ascertained to the soil respiration as a result to decomposition of green manures and increase the numbers of bacteria followed the carbon dioxide evolution (Haney et al., 2008). Trognitz et al. (2012) reported that the beneficial soil microorganisms stimulate the plant to resist stresses during developmental stages and could occupy niches inside the plant, which were in concurrence with pathogens, whereas deleterious microorganisms might be used for the biocontrol of weeds. Plant-associated microbial communities are essential for growth parameters like plant nutrition, resistance to biotic and abiotic stresses, plant survival and distribution.

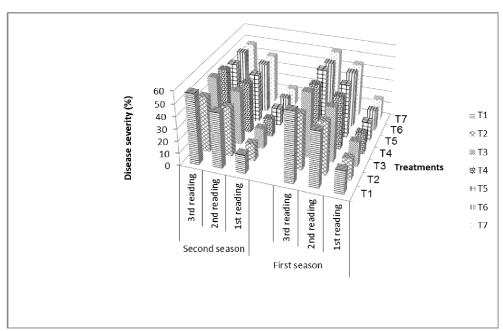


Fig. (1): Disease severity percentage of *Stemphylium vesicarium* during cultivation of onion crop due to the different treatments in the two seasons.

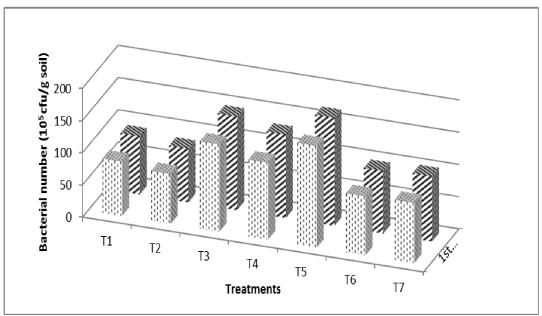


Fig (2): Total bacterial numbers in rhizosphere of infected onion plants of both seasons due to the different treatments.

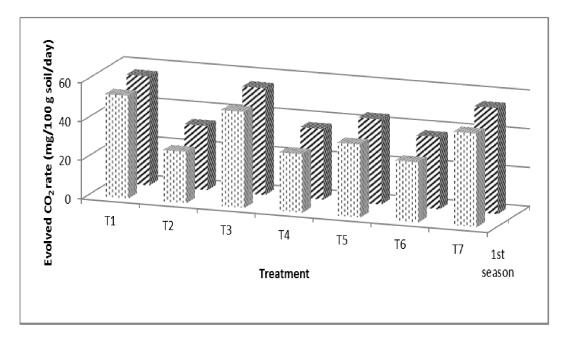


Fig. (3): CO₂ evolution in the rhizosphere of onion plant infected by *Stemphylium* vesicarium in both seasons as affected by the different treatments.

Fig. (4 a, b and c) brightens up the effect of the applied treatments on nitrogen, phosphorus and potassium contents in plants. Generally, the second season was better than the first one. The cyanobacterial filtrate followed by azolla and *Pleurotus* filtrates boosted the nitrogen absorption by the plant. In case of phosphorus, all the

biological treatments were lower contents than chemical pesticide treatment. The pesticide and the control treatments were not affected by the infection by *Stemphylium vesicarium*. *Per contra* in the case of

phosphorus, all the biological treatments were better in absorption of potassium than the chemical pesticide and control treatments, where cyanobacteria and azolla filtrates were the eclecticism.

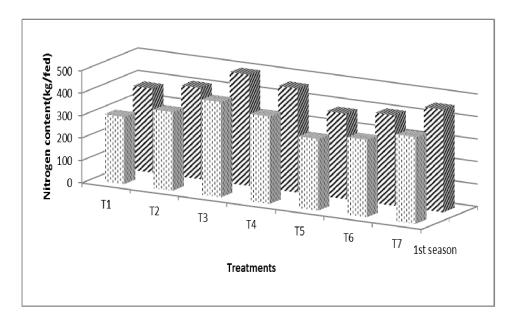


Fig. (4a): Nitrogen contents in onion plants treated by chemical pesticide and biological agents in the presence of *Stemphylium vesicarium* during the two seasons.

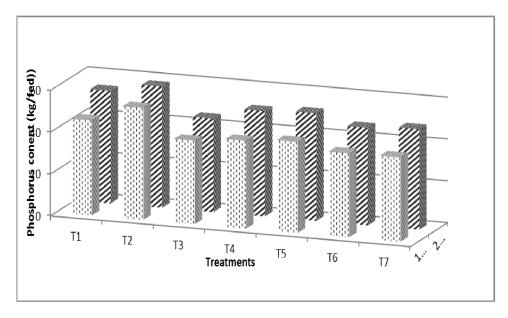


Fig. (4b): Phosphorus contents in onion plants treated by chemical pesticide and biological agents in the presence of *Stemphylium vesicarium* during the two seasons.

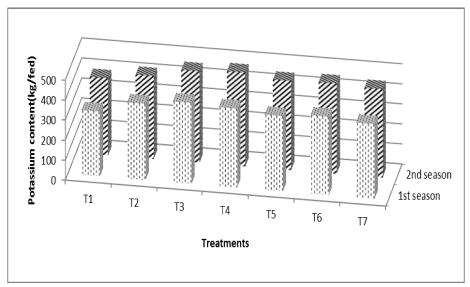


Fig. (4c): Potassium contents in onion plants treated by chemical pesticide and biological agents in presence of *Stemphylium vesicarium* during two cultivation seasons.

Fig. (5) clarifies the influence of chemical pesticide and some different biological agents on onion plants in the presence of Stemphylium vesicarium during the two cultivation seasons. High content of phenol was observed in the Pleurotus filtrate treatment, while chemical and biological treatments showed lower values. Total phenols were not affected by by Stemphylium vesicarium infection and recorded highest value in the control treatment comparable the Pleurotus filtrate treatment. Cheng et al.(2013) studied the relationship between the phenolic and flavonoid contents and antioxidant, and the correlation among them. The increasing of antioxidants is due to elevating the contents of the total phenol and flavonoids that have scavenging activity of free radicals.

Fig. (6) Illustrates the effect of some bioagents on total carbohydrates in onion plants infected by *Stemphylium vesicarium* during the two seasons. All the biological treatments had positive effects on total carbohydrates. The promising treatments were cyanobacterial, azolla and *Pleurotus* filtrate treatments. The low values of total carbohydrates appeared in the control and chemical pesticide treatments. Ashraf *et al.* (2013) reported that exoploysaccharides

released from some microorganisms played a role in fertility and enhancing the formation of soil micro-aggregates and contribute to building up soil physical structure, regulated nutrients and water flow from rhizosphere soil to the plant, promoted the vegetative growth, and protect the root against pathogens.

Table (2) presents some quantitative and qualitative characteristics of onion plants infected by Stemphylium vesicarium and chemical and treated with biological treatments, during the two seasons. The cyanobacteria followed by Pleurotus filtrates were the auspicious treatments in regard to the total yield, marketable yield and average of bulb weight of the onion plants in the respective seasons, where records were 16.56 ton/fed., 14.18 ton/ fed. and 104 g for the cyanobacterial filtrate treatment and 16.43 ton/fed., 13.95 ton/fed. and 100.33 g for the Pleurotus filtrate treatments. Time of maturity decreased as a result of the chemical pesticide reaching 161.7 days and the mixture of Serattia and Pseudomonas reached 161 days followed by the cyanobacterial and Pleurotus filtrates reached to 162 days for both last treatments in the first season while in the second season, the time of maturity diminished

significantly by the chemical pesticide (153.3 days). The chemical pesticide and azolla filtrate increased the percentage of single bulb reaching 91 and 90.67 % respectively, followed by the cyanobacterial

filtrate, mixture of *Serratia* and *Bacillus* and *Pleurotus* filtrate, which reached 89.67, 90 and 90 % in the second season, respectively.

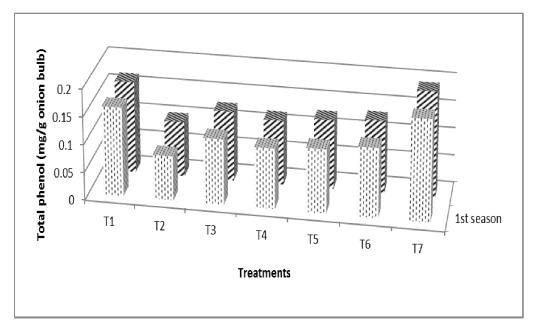


Fig. (5): Total phenols in onion plants infected by *Stemphylium vesicarium* and treated by chemical pesticide and different bioagents during the two seasons.

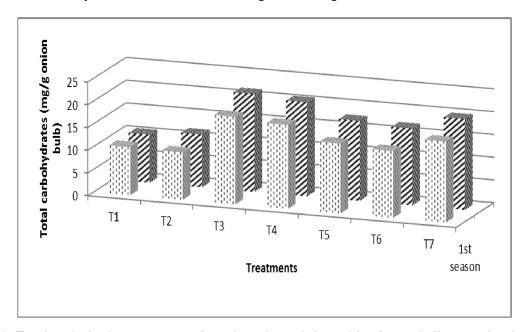


Fig (6): Total carbohydrate contents in onion plants infected by *Stemphylium vesicarium* and treated with chemical pesticide and different bioagents during the two cultivation seasons.

Table (2): Some quantitative and qualitative characteristics of infected onion plant by Stemphylium vesicarium that treated with some chemical and biological treatments through two seasons.

	ıreaiii	nents thro		seasons.							
Treatment	Total yield (ton/fed)	Marketable yield (ton/fed)	Average of weight bulb (g)	Time of maturity (days)	% of single bulb	% of double bulb	% of bolters	No. of leaves/plant	Plant height (cm)		
2010-2011 season											
T1	12.76	11.60	75.00	164.7	91.0	9.667	3.333	6.667	62.33		
	G	D	G	A	A	DE	A	EF	ABC		
T2	14.97	13.89	93.33	161.7	88.33	12.67	1.0	6.767	62.33		
	CDE	AB	C	ABCD	ABC	BC	C	EF	ABC		
T3	15.11	13.84	94.33	162.0	83.0	15.67	1.333	7.167	61.33		
	CDE	AB	A	ABC	D	A	BC	CDE	BC		
T4	13.75	12.70	86.25	164.0	85.33	14.0	1.333	7.233	63.67		
	FG	ABCD	D	AB	CD	AB	BC	BCDE	ABC		
T5	13.14	12.03	84.33	162.7	87.67	9.667	2.667	6.533	63.0		
	G	CD	G	ABC	ABC	DE	AB	F	ABC		
T6	13.72	12.07	85.66	161.0	87.67	12.33	1.333	7.033	60.33		
	FG	CD	D	ABCD	ABC	AB	BC	DEF	C		
T7	14.28 EF	12.68BCD	88.66 D	162.0ABC	86.33BCD 1 season	13.33AB	1.333BC	7.1DEF	61.67BC		
T1	14.70	12.86	96.0	155.7	89.0	9.333	3.333	7.1	64.0		
	DEF	ABCD	EF	BCD	ABC	DE	A	DEF	ABC		
T2	15.89	13.97	95.67	153.3	91.0	7.333	2.333	7.8	64.33		
	ABC	AB	F	D	A	E	ABC	AB	AB		
Т3	16.56	14.18	104.0	158.0	89.67	10.33	1.667	7.833	63.67		
	A	A	E	ABCD	AB	CD	BC	A	BC		
T4	15.90	13.06	97.33	154.7	90.67	8.333	2.333	7.567	65.67		
	ABC	ABCD	EF	CD	A	DE	ABC	ABCD	A		
T5	15.26	13.07	94.33	156.0	90.0	8.333	2.333	7.167	64.33		
	CED	ABCD	B	BCD	AB	DE	ABC	CDE	A		
Т6	15.52	13.46	92.33	157.0	89.0	8.333	2.333	7.6	63.0		
	BCD	ABC	FG	ABCD	ABC	DE	ABC	ABCD	ABC		
T7	16.43	13.95	100.33	155.7	90.0	8.333	2.333	7.7	64.0		
	AB	AB	EF	BCD	AB	DE	ABC	ABC	ABC		
LCD at 0.05	1.032	1.493	8.133	8.474	3.932	2.559	1.634	0.5739	3.696		
				Main effec	ct of years			•			
Mean	13.96	12.69	149.0	162.6	87.05	12.48	1.762	6.929	62.10		
	B	A	A	A	B	A	A	B	AB		
Mean	15.75	13.51	138.7	155.8	89.9	8.667	1.524	7.533	64.14		
	A	A	B	B	A	B	A	A	A		
LCD at 0.05	1.462	1.085	8.818	4.767	1.525	3.718	0.7932	0.2918	1.879		
				Main effect of	of treatments						
T1	13.73	12.23	85.5	160.2	90.0	9.5	3.333	6.883	63.17		
	D	D	E	A	A	BC	A	BC	AB		
T2	15.43	13.93	235.5	157.5	89.67	10.0	1.667	7.283	63.33		
	AB	AB	B	A	A	BC	B	AB	AB		
Т3	15.84	14.01	375.8	160.0	86.33	13.0	1.5	7.5	62.5		
	A	A	A	A	B	A	B	A	AB		
T4	14.82	12.88	225.3	159.3	88.0	11.17	1.833	7.4	64.67		
	BC	BCD	C	A	AB	B	B	A	A		
T5	14.20	12.55	234.8	159.3	88.83	9.0	3.0	6.850	63.67		
	CD	CD	B	A	AB	C	A	C	AB		
T6	14.62	12.77	219.5	159.0	88.33	10.5	1.833	7.317	61.67		
	C	CD	D	A	AB	BC	B	A	B		
T7	15.35	13.32	225.2	158.8	88.17	10.83	1.833	7.400	62.83		
	AB	ABC	CD	A	AB	B	B	A	AB		
LCD at 0.05	0.7297	1.056	5.751	5.992	2.780	1.809	1.155	0.4058	2.613		

The percentage of double bulb was low in the chemical pesticide reaching 7.33 %, while all the biological treatments scord 8.33 % except the cyanobacterial filtrate gave 10.33 % in the second season. The percentage of bolters decreased by the chemical pesticide to reach 1.0 %, while all the biological treatments reached 1.33 %, except the mixture of Serratia and Bacillus which reached 2.67 % in the first, season while in the second season, the algal extract gave 1.67 %, while all the chemical and biological treatments acheived 2.33 %. The biological treatments were the concerning the number of onion leaves. Their values were arranged, in the second season, as follows: algal extract (7.83), chemical pesticide (7.8), Pleurotus filtrate (7.7), mixture of Serratia and Pseudomonas azolla extract (7.57), mixture of Serratia and Bacillus (7.17) and the last was control (7.1), in a descending Metabolite exudates excreted microorganisms like cyanobacteria and yeasts can improve the quality and quantity of onion yield throughout the excrtion of vitamins, amino acids and plant hormones (Abdel-Raouf et al., 2012 and Shehata et al. , 2012). Vitamins like glutathione and amino acids like sulpher amino acids could significantly promote the growth and quality of green onion criteria such as shoot length, white part length, bulb diameter, leaf photosynthetic pigment, number of leaves, fresh and dry weight of onion plant (El-Awadi and Abd El Wahed, 2012). Bioactive compounds, such as auxins, gibberellins cytokinins produced and were cyanobacteria and act as bioregulators for plant growth. These regulators can improve the total carbohydrates, proteins chlorophyll in the plant and also increase organic conditions, improve water holding structure. capacity and improve bufferina capacity against fluctuation in pH levels of the soil (Subramaniyan and Malliga, 2011).

Conclusion:

This investigation aims at controling the blight disease on the leaves and stalks of onion plant caused by *Stemphylium vesicarium*. The biological treatments had a

distinctive position in enhancing the quality and quantity properties of infested plants. The second season was often better than the first one as a result of cultivation at the same location, due to the residual effect of the applied treatments. The cyanobacterial and Pleurotus filtrates had an obvious role in overcoming the disease and in raising the rate of CO2 evolution in soil, content of nitrogen in leaves, crop yield, marketing yield and weight of bulbs of the onion plant, while Pleurotus filtrate was the best in increasing the phenol content in the leaves of onion plant. In general, biological treatments were supposed to replace the chemical pesticide, as to control the blight disease and improve the crop yield of onion quantitatively and qualitatively.

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النشاط الحيوى لبعض الكائنات الحيه الدقيقه لمكافحة مرض اللفحه (ستيميفيليوم فيسيكاريوم) لنبات البصل

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الملخص العربي

يعتبر البصل عنصرا اساسيا على المائدة المصريه ، و نظرا لما يسببه فطر ستيميفيليوم فيسيكاريوم تليفا و بقعا زرقاء على اوراق البصل التى تقلل من المحصول و تسبب صفاتا غير مرغوب فيها . و لذا وجه علماء البيولوجيا جهودهم لمكافحة المرض بيولوجيا للحد من استخدام المبيدات الكيميائيهز وقد استخدم في هذا البحث مجموعات من الكائنات الحيه الدقيقه مثل خليط من بكتريا السيراتيا مع الباسياس ، خليط من السيراتيا مع السيدوموناس ، و راشح خليط من سلالتي السيانوبكتريا انابينا اوريزا مع نوستوك ماسكورم ، راشح الازولا بيناتا ، و راشح فطر البلوروتس كولومبينوس لمكافحة المرض.

و اظهرت النتایج ان اکثر المعاملات البیولوجیه تأثیرا کان من خلال استخدام خلیط من السیراتیا مع الباسیاس یتبعها استخدام راشح فطر البلوروتس حیث کان تأثیرهما علی شدة الاصابه یقترب من المبید الکیمیائی . کما وجد ارتفاع ملحوظ فی محتوی الفینولات عند المعامله براشح فطر البلوروتس . و احتات المعامله براشح السیانوبکتریا و یتبعها راشح الفطر مکانه ممیزه فی تأثیرهما علی المحصول و المحصول التسویقی و متوسط وزن البصله التی کانت نتائجها کالتالی : ۱۲.۵۱ طن الفدان ، ۱۲.۱۸ طن الفدان ، ۱۲.۶۲ جم علی التوالی بالنسبه لراشح فطر السیانوبکتریا ، ۱۲.۶۳ طن الفدان ، ۱۳.۹۰ جم علی التوالی بالنسبه لراشح فطر البلوروتس بالمقارنه بالکنترول الذی کانت نتائجه ۱۶.۷۷ طن الفدان ، ۱۲.۸۲ طن الفدان ، ۹۲ جم علی التوالی و ذلك من خلال الموسم الثانی.