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RESEARCH PAPER

Biological control of date palm diseases with native antagonistic fungi of Oued Righ region (Algerian Sahara)

Lutte biologique contre les maladies fongiques du palmier dattier avec des champignons antagonistes de la région d'Oued Righ (sahara algérien)

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Abstract

The date palm (*Phoenix dactylifera* L.) is one of the principal species in the south of Algeria. However, this speculation is confronted to some problems including plant health. Several fungal and bacterial antagonists were used to control plant diseases. The present study was undertaken to investigate the interaction of antagonistic fungi (*Trichoderma harzianum, Pythium sp. and Aspergillus niger*) and pathogenic fungi (*Fusarium sp. and Alternaria sp.*) of date palms. The confrontation test of two groups of microorganisms with the antagonists on nutrient agar revealed the inhibition of mycelia growth of the two fungi. The *Fusarium sp.* was inhibited with 68% by *T. harzianum*, 63% by *Pythium sp.* and 46% by *A. Niger*. As for the *Alternaria sp.*, inhibition rates of 63% and 54% were exercised by *T. harzianum* and *Pythium sp.*, respectively. Starting from the sixth day of the cultivation, the antagonistic fungi invaded the colonies of the pathogenic fungi and began to sporulate marking a very important myco-parasitic power.

Keywords: Date palm, biological control, Fusarium sp., Alternaria sp., antagonistic fungi, Algeria.

Résumé

Le palmier dattier (<u>Phoenix dactylifera</u> L.) est l'une des principales espèces cultivées dans le sud algérien. Cette spéculation a été confrontée à quelques problèmes phytosanitaires. Plusieurs champignons et bactéries antagonistes sont utilisés en lutte biologique. La présente étude a été menée afin d'étudier l'interaction entre des champignons antagonistes (<u>Trichoderma harzianum</u>, <u>Pythium sp</u>. et <u>Aspergillus niger</u>) et des champignons phytopathogènes (<u>Fusarium sp. et Alternaria sp.</u>) du palmier dattier. Le test de confrontation des deux groupes de microorganismes dans un milieu de culture artificiel (PDA) a révélé un effet inhibiteur très marqué. Le <u>Fusarium sp.</u> a été inhibé de 68% par <u>T. harzianum</u> 63% par <u>Pythium sp.</u> et 46% par <u>A. Niger</u>. Quant à l'Alternaria sp., des taux d'inhibition de 63% et 54% ont été exercés par <u>T. harzianum</u> et <u>Pythium sp.</u>, respectivement. A partir du sixième jour de la mise en culture, les champignons antagonistes envahissent les colonies des champignons phytopathogènes et commencent à sporuler et marquer un pouvoir myco-parasitaire très important.

Mots-clés: Palmier dattier, lutte biologique, <u>Fusarium sp., Alternaria sp.</u>, champignon antagonist, Algérie.

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1.INTRODUCTION

The date palm is one of the well adapted species to the arid, semi-arid and hot areas. It has the ultimate fruit of the desert where it plays at the same time an economic role thanks to the production of the dates which constitutes a base of human and animal consumption, and an ecological role since it confers on the oasis. Its structure makes protection against the wind and the projection of the desert while creating under its cover a microclimate favorable to the development of many additional cultures that condition the saving of the oasis and the stability of the populations which live there.

Date palms can be subject to many diseases or complexes of diseases, among which some are serious, can result an inescapable death. Many review works were published on the pathology of date palm; by many authors. Thus, laboratory studies were conducted into the pathology laboratory of the station of INRAA, 13 fungi were found on different parts of the dead palm trees. The folowing fungal types were identified: *Diplodia phoenicum*, *Theilaviopsis paradoxa*, *Phytophthora sp.*, *Helminthosporium sp.*, *Stemphylium botryosum*, *Alternaria sp.*, *Aspergillus niger*, *Aspergillus sp.*, *Cladosporium sp.* and Fusa*rium sp.* (Lakhdari et al., 2013).

In order to look for other alternatives methods by using biological control against aggressive pathogenic fungi of date palm, in this work we studied the interaction between three antagonistic fungi isolated from potato culture of Oued Righ region in the experimental station of Sidi Mehdi (INRAA) against two pathogenic fungi which cause diseases on the date palm of the southeastern Algeria. This work therefore consists by studying *in vitro* the interaction between these antagonistic and pathogenic fungi of palm trees by direct contact.

2.MATERIAL AND METHODS

2.1. Fungal agents

Infected vascular tissues from stem, leaflet and root parts of date palm cultivar showing symptomswere collected separately from farmer's field in the regions ofMegarine and Beldet Omar which are situated in the southeastern of Algeria (OuedRigh) (Fig. 01).

The analytical techniques used werethose usually practiced in a laboratory of plant pathology. Fragments of reached parts from 5 to 10 mm presenting of the typical symptoms are cut out then planted in a suitable culture medium after disinfection, rinsing with sterile distilled water, and then drying (Fig. 02). Incubation takes place at a temperature between 24-26 °C. Once well differentiated colonies, they will be then re-inoculated in the same medium of sowing (for obtaining the purified cultures).

The identification of fungal flora is not only carried out by the color and the shape of the colony but also from an examination of mycelia and conidia on the nutrient medium. We used the following identification keys: Barnett(1972) and Ellis (1971).

2.2. Antagonistic agents

For the isolation of the antagonist *Trichoderma harzianum*, we collected the leaves of the potato which have morbid symptoms were cut into small pieces and then subjected to disinfection with sodium hypochlorite (5%) for 3 min, then rinsed with sterile distilled



Figure 01: Infected palm trees

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water. This operation aimed to eliminate saprophytic flora. After these, fragments were dried on a paper sterile filtered and aseptically placed on an agar medium (PDA) into Petri dishes with 3 fragments per dish. Incubation took place at 20 °C under a photoperiod of 12 hours (Fig. 03).

The other antagonistic fungi were isolated from soil (*Pythium sp.*) and palm trees (*Aspergillus niger*).

2.3. Method of mycelia growth measurement

The direct confrontation between the antagonistic fungiand the pathogens agent was studied by the method of Rapilly(1968). This method is consisted to put on the same Petri, two agar pellets (5 mm diameter) one strain of the antagonist and other of pathogen are positioned along a diametrical axis 3 cm away (Figs. 04, 05). We have tested every antagonist on every pathogen (*T. harzianum vs Fusarium sp, T. harzianum vs Alternaria sp.; Pythium sp. Vs Fusarium sp., Pythium sp. vs Alternaria sp. and A. niger vs Fusarium sp.*) with three repetitions in every test; incubation is performed at 25°C for six days in the dark. The evolution of mycelia growth wasperformed every 24 hours by measuring the diameter of the colony of the pathogen and the antagonist. The valuation of inhibition by the antagonist is estimated by calculating the percentage of inhibition of mycelia growth by the following formula:

I %=[1-Cn/Co]x 100

Cn: average diameter of colonies of pathogen in the presence of the antagonist.

Co: average diameter of colonies of control.

2.4. Statistical analyses

For the *in vitro* (on Petri plates) assays, the data were collected as mean colony diameter (average of two perpendicular diameters) values in each replication. These analyses were calculated by statistical software (STAT BOX 6.0.4.,GRIMMERSOFT). The device is held in total uni-factorial randomization by Newman and Keuls test at 5% and 1% (P \leq 0.05 and P \leq 0.01). The antagonistic treatments (fungi species and untreated control) and *Fusarium sp., Alternaria sp.* were the two fixed factors.



Figure 02: Sowing fungi



Figure 03: Sowing and purification of indigenous fungi





Figure 04: Antagonism method







Figure 06: Fungal pathogens (A: Fusariumsp.; B: Alternaria sp.)

3. Results

3.1. Fungal antagonists and pathogens

Among the isolated fungi from the palm trees, we have observed that the most important fungi were *Fusarium sp. and Alternaria sp.* (Fig. 06) with an abnormal symptoms on the date palms; and according to the antagonism effects of several fungi, we were isolated from the soil of the experimental station of Sidi Mehdi (INRAA) some fungi which contain an antagonist characters like *T. harzianum and Pythium sp.*, with an addition one (A. niger) which were isolated from palm trees (Fig. 07).

3.2. Direct confrontation between *Fusarium sp.* and *Trichoderma harzianum*

Direct confrontation of *T. harzianum and Fusarium sp*.showed a faster mycelia growth of T. harzianum than of Fusariumsp. (Fig. 08). After three days of incubation, the Petri was completely filled by the antagonist, while the *Fusarium sp*. only occupied a surface of 12,33 mm diameter; which corresponded to an inhibition of mycelia growth greater than 68% (Tab. 01). The control of *Fusarium sp*. occupied an area of approximately 39 mm in diameter until the fifth day(Fig. 08).

Table 1: Direct confrontation	between	Fusarium sp	. and
Trichoderma harzianum			

Days	Pathogen <i>Fusarium sp.</i>	Antagonist <i>T. harzianum</i>	Control Fusarium sp.	Inhibition percentage
1	4,5 mm	8 mm	5 mm	10 %
2	9,66mm	26,33 mm	15,16 mm	36 %
3	12,33 mm	39 mm	24,16 mm	50 %
4	12,33 mm	39 mm	34,16 mm	63 %
5	12,33 mm	39 mm	39 mm	68 %
6	12,33 mm	39 mm	39 mm	68 %

Table 2: Direct confrontation between *Alternaria sp.* and*Trichoderma harzianum*

Days	Pathogen <i>Alternaria sp.</i>	Antagonist <i>T. harzianum</i>	Control Alternaria sp.	Inhibition percentage
1	-	-	-	-
2	3,16 mm	3,83 mm	3,5 mm	10 %
3	5,75 mm	7 mm	10,25 mm	44 %
4	7,66 mm	10,66 mm	15,5 mm	51 %
5	8,66 mm	14,83 mm	18,16 mm	53 %
6	9 mm	18,33 mm	24,16 mm	63 %

3.3. Direct confrontation between *Alternaria sp.* and *T. harzianum*

Direct confrontation of *Trichoderma harzianumand Alternaria sp.* showed a faster mycelia growth of *T. harzianum* than that of *Alternaria sp.* (Fig. 09). After six days of incubation, the Petri was filled by the antagonist, with a remarkable growth rate. Whereas, *Alternaria sp.* occupied a surface area of 9 mm in diameter; with more or less aerial mycelium and a low growth rate compared to the antagonist. This correspond an inhibition of mycelia growth by 63% (Tab. 02).

3.4. Direct confrontation between *Fusarium sp.* and *Pythium sp.*

The antagonism test between *Pythium sp. and Fusarium sp.* showed a faster mycelia growth of *T. viride* than that *of Alternaria sp.* (Fig. 10). Fo ur days after incubation, the Petri was completely filled by the antagonist, with anamazing growth rate. Whereas, *Fusarium sp.* occupied a surface area of 2,25 mm in diameter (Fig. 10). This antagonist correspond an inhibition of mycelia growth by 63%(Tab. 3).

3.5. Direct confrontation between *Alternaria sp.* and *Pythium sp.*

The results presented in the table below, the direct confrontation of *Pythium sp.* and *Alternaria sp.* showed a faster mycelia growth of Pythium sp. than of Alternariasp. (Fig. 11). After four days of incubation, the Petri was completely filled by the antagonist, while the *Alternaria sp.* only occupied a surface of 9 mm diameter; which reached to an inhibition of mycelia growth greater than 54% (Tab. 4). The control of *Alternaria sp.* occupied an area of approximately 28,5 mm in diameter until the sixth day (Fig. 11).

Table 3: Direct confrontation between *Fusarium sp.* and*Pythium sp.*

-	-			
Days	Pathogen Fusarium sp.	Antagonist Pythium sp.	Control Fusarium sp.	Inhibition percentage
1	-	-	-	-
2	1 mm	13 mm	3 mm	67%
3	2,25 mm	32,25 mm	4,75 mm	53%
4	2,25 mm	40,75 mm	6,62 mm	66%
5	3,45 mm	43,5 mm	8,5 mm	59%
6	3,75 mm	45 mm	10,25 mm	63%



Figure 07: Fungal antagonists (A: T. harzianum; B: Pythium sp.; C: A. niger)

Figure 08: Direct confrontation between *Fusarium sp.* and *Trichoderma harzianum* after six days of incubation





Figure 09: Direct confrontation between *Alternaria sp.* and *Trichoderma harzianum* after six days of incubation

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Table 4: Direct confrontation between *Alternaria sp.* and*Pythium sp.*

-	1			
Days	Pathogen <i>Alternaria sp.</i>	Antagonist Pythium sp.	Control Alternaria sp.	Inhibition percentage
1	-	-	-	-
2	3,83 mm	18,33 mm	5,66 mm	32%
3	8,66 mm	32 mm	12 mm	28%
4	9 mm	38 mm	17,83 mm	50%
5	12,16 mm	47,16 mm	23,5 mm	48%
6	13 mm	47,5 mm	28,5 mm	54%

Table 5: Direct confrontation between *Fusarium sp.* andAspergillus niger

Days	Pathogen Fusarium sp.	Antagonist <i>A. niger</i>	Control Fusarium sp.	Inhibition percentage
1	-	-	-	-
2	1,66 mm	2 mm	3 mm	45%
3	2,25 mm	3,16 mm	4,75 mm	53%
4	3,33 mm	3,66 mm	6,62 mm	50%
5	5,5 mm	29,66 mm	8,5 mm	35%
6	5,5 mm	45,16 mm	10,25 mm	46%

3.6. Direct confrontation between *Fusarium sp.* and *Aspergillus niger*

The data of the table below showed us a great inhibition between the antagonist *Aspergillus niger* and the pathogen *Fusarium sp.*(Fig. 12). After six days of incubation, the Petri was completely filled by the antagonist, while the *Fusarium sp.* occupied a surface of 5,5 mm diameter; which corresponds to an inhibition of mycelia growth greater than 46% (Tab. 5).

3.7. Statistical analyses

The statistical analyses showed that all treatments tested on the pathogenic fungal of palm trees had a significant effect. These results confirm that there is an inhibition caused by the antagonistic fungi against the two pathogenic fungi.



Figure 10: Direct confrontation between Fusarium sp. and Pythium sp. after six days of incubation



Figure 11: Direct confrontation between Alternaria sp. and Pythium sp. after six days of incubation



Figure 12:Direct confrontation between *Fusarium sp.* and *A. niger* after six days of incubation

4. Discussion

Whipps and McQui lken (1993) reported that A. niger, *A. terreus, P. citrinum, T. harzianum* and species of Bacillus control soil-borne diseases (In Alwathnani, 2012). According to the last author, biological control of *Fusarium oxysporum f. sp. lycopersici* by *A. niger* revealed a rate of 35,6% and by *T. harzianum* was about 44,4%.

The *Trichoderma* have been known a long time ago by their antagonistic activities against many fungi such as: *Fusarium solani var coerulum, Fusarium roseum var sambucinum and Fusarium roseum var graminearum* (Daami-Remadi, 2001); *Alternaria alternata* (Gveroska & Ziberoski, 2011); *Fusarium oxysporum f. sp.* Lentis (Essalmani & Lahlou, 2004).

Also Sundaramoorthy and Balabaskar (2013) have found that the efficacy of *Trichoderma spp.* against wilt of tomato caused by *Fusarium oxysporum f. sp. Lycopersici* note an important rate of 57%. Therefore, the antagonist *T. harzianum* is chosen to be the most promising bio-control agent for *F. oxysporum f. sp. lycopersici*.

The same author said that the mycelia growth inhibition of *F. oxysporum* (60%) at the 5th day in the confrontation plate assay was negatively correlated with mycelia growth of *F. oxysporum* in the presence of *T. harzianum*.

Allowing to Jat & Agalave (2013), in an investigation *Trichoderma harzianum* hampered the growth of *Alternaria alternata* by 48.33% but Hussainet al., (2009) found that *Trichoderma harzianum* reduced the growth of *Alternaria alternata* by 67.7%.

Chanchal (2014), in antagonistic tests, found that *T. harzianum* inhibited 70% growth of *Fusarium moni-liforme* and 55% of *Fusarium sacchari*.

The results of Bouziane (2011), showed that the inhibition of mycelia growth in the direct confrontation between *T. harzianum* and *Alternaria sp.* is about 34%.

Alkatatnyet al. (2007) found that the fungal antagonist *Pythium oligandrum* inhibited the fungal pathogen *F. oxysporum* by 100%.

Bouneghou (2011) noted that the mycelium growth of *Alternaria alternata* was inhibited by *Pythium sp.* with an inhibition percentage of 50.3%, unlike, *Fusarium roseum* which was inhibited less than the first pathogen with 40%.

Conclusion

The present evaluation gave clear indication that the isolates of *Trichoderma harzianum* (isolated from leaves of potato), *Pythium sp.* (isolated from soil) and *Aspergillus niger* (isolated from palm trees) are strong and virulent antagonists.

Indeed, the direct confrontation tests between the pathogen (*Fusarium sp.* and *Alternaria sp.*) and antagonists (*T. harzianum, Pythium sp. and A. niger*) on a medium culture (PDA) have shown an inhibition of the pathogen mycelia growth tested. the Confrontation of the fungal pathogen with the antagonists revealed the inhibition of mycelia growth of the three fungi: *T. harzianum* with (*Fusarium sp.* = 68% and *Alternaria sp.* = 63%); *Pythium sp.* with (*Fusarium sp.* = 63% and *Alternaria sp.* = 54%) and A. niger with *Fusarium sp.* = 46%).

These results allowed us to believe that these three fungal antagonists haveability to remove *Fusarium sp.* and *Alternaria sp.* from date palm and can be tested *in vivo* and have practical interest in the biological control against diseases of this culture.

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BIBLIOGRAPHY

Alwathnani HA, Kahkashan P (2012) Biological control of *Fusarium* wilt of tomato by antagonist fungi and cyanobacteria. African Journal of Biotechnology 11(5): 1100-1105.

Barnett HL, Hanter BB (1972) Illustraed genera of imperfect fungi Burgess. Pub. Co. Minneapolis, USA pp 241.

Bouneghou SB (2011) L'effet inhibiteur de Pythium sp. sur la croissance mycélienne de Fusarium roseum et d'Alternaria alternata. Mémoire Master. Univ. Constantine 50 p.

Bouziane Z, Dehimat L, Abdelaziz W, Benabdelkader M, Kacem Chaouche N (2011) The antagonism between *Trichoderma viride* and other pathogenic fungal strains in *Zea mays*. Agriculture and Biology Journal of North America 4: 584-590.

Chanchal G, Sharmita G (2014) Antagonistic effect of *Trichoderma viride* and *Trichoderma harzianum* against plant pathogenic fungi and its growth on different agro-waste substrates. International Journal of Engineering and Technical Research 23-25.

Daami-Remadi M (2001) Lutte biologique contre les *Fusarium spp.* agents pathogènes responsables de la pourriture sèche des tubercules de pomme de terre. Mémoire de Diplôme d'Études Approfondies en Protection des Plantes et Environnement. Tunisie: École supérieure d'Horticulture et d'Élevage de Chott Mariem 72 p.

El-Katatny MH, Abdelzaher HMA, Mahmoud ShoulkamyA (2007) Antagonistic actions of *Py*thium oligandrum and *Trichoderma harzianum* against phytopathogenic fungi (*Fusarium oxysporum and Pythium ultimum var. ultimum*). Phytopathology and Plant Protection 39 (4): 289-301

Ellis MB (1971) More Dematiaceous hyphomycetes commonwealth, Myccological Institute.

Essalmani H, Lahlou H (2004) Induction par *Trichoderma harzianum*, de la résistance des plantes de lentille contre *Fusarium oxysporum f. sp. lentis*. Actes Inst. Agron. Vet. (Maroc) 24 (1&2): 51-58.

Gveroska B, Ziberoski J (2011) *Trichoderma harzianumas* a biocontrol agent against Alternaria alternata on tobacco. Appl. Techn. & Innov 7(2):.67-76

Jat JG, Agalave HR (2013) Antagonistic properties of *Trichoderma* species against oilseed-borne fungi. Science Research Reporter 3 (2):171-174.

Rapilly F (1968) Les techniques de mycologie en pathologie végétale. Annales des Epiphytes. INRA 19: 102.

Sundaramoorthy S, Balabaskar P (2013) Biocontrol efficacy of Trichoderma spp. against wilt of tomato caused by *Fusarium oxysporum f. sp. Lycopersici.* Journal of Applied Biology & Biotechnology 1(03):036-040.

الأخضري و، أفروخ ع، دهليز ع، حمي ح، سود ع، بوشكيمة ن (2013) .عزل و تشخيص الفطريات الممرضة لأشجار نخيل التمر بمرتفع وادي ريغ ورقلة- الجزائر (مجلة Phoenix dactylifera L زراعة الرافدين . المجلد 41. الملحق 01. ص. 323-333