Phytohormones Producing Fungal Endophytes Enhance Nutritional Status and Suppress Pathogenic Fungal Infection in Tomato

M. M. G. Saad\(^1\)*, and H. H. Badry\(^2\)

ABSTRACT

Two endophytic fungi, namely, *Curvularia lunata* and *Nigrospora sphaerica*, were isolated from *Melia azedarach*, an exotic tree introduced in Egypt from Asia. The fungal endophytes were identified by microscopic examination and molecular identification of nucleotide sequence by DNA sequencing of the purified PCR product. Their antagonistic activities against phytopathogenic fungi and their ability to produce important growth hormone and providing some necessary nutrients for plant growth were also evaluated. Both endophytes exhibited antagonistic activities: *C. lunata* caused 56 and 50% growth inhibition of *Alternaria solani* and *Fusarium oxysporum*, while *N. sphaerica* suppressed both pathogenic fungi by 63.4 and 56.6%, respectively. *N. sphaerica* was able to dissolve insoluble phosphorus, produce ammonia, and secrete 40 µg mL\(^{-1}\) of IAA. In contrast, *C. lunata* failed to dissolve phosphorus, secreted less amount of IAA (3 µg mL\(^{-1}\)), but produced ammonia. A greenhouse pot experiment was conducted using phosphorus deficient soil to find out the ability of both endophytes to improve growth of tomato plants. *N. sphaerica* significantly increased shoot fresh weight by 13 and 22% over *C. lunata* and the control, respectively. Concerning the nutritional status of tomato plants, both endophytes led to significant increase in nitrogen concentration in shoots when applying 50% of the recommended mineral fertilizer. *N. sphaerica* enhanced phosphorus concentration in shoots by 13% over the control. Finally, the antifungal activities of both endophytes against *F. oxysporum* in tomato plants were tested under glasshouse conditions. *N. sphaerica* was more potent than *C. lunata* in suppressing 40% of *F. oxysporum* infection and had positive impact on tomato plant growth. Our study results highlight the potential use of *N. sphaerica* endophytic fungi as plant biofertilizers and bio-control agent under glasshouse conditions.

Keywords: Antifungal activity, *Curvularia lunata*, *F. oxysporum*, *Nigrospora sphaerica*, Plant growth promotion.

INTRODUCTION

Searching for new, safe, and eco-friendly alternatives of synthetic pesticides and chemical fertilizers is a major concern for clean environment and sustainable agriculture. To minimize environmental pollution and health problems, bio-control microorganisms and their metabolites could be a good alternative for different synthetic agrochemicals. Many microorganisms have the ability to promote plant growth and reduce disease and pest attack by different mechanisms. These modes of action include increasing nutrients availability by fixing, solubilizing, and mobilizing micro and macro elements and producing active secondary metabolites antagonistic to large scales of pest...
Endophytes are microorganisms that spend the whole or part of their lifecycle colonizing inter-and/or intracellular parts of their host plants tissues without causing any symptoms of disease (Azevedo and Araújo, 2007). Endophytes have a great impact on host plant growth and yield. It promotes plant growth, suppresses pathogens, increases tolerance to drought, and solubilizes nutrients. However, the endophyte-plant interaction relationship is not fully understood as it ranges from mutualism to latent pathogenesis depending on plants and microbes genotypes, environmental conditions, and the dynamic interactions between the plant microbiome (Redman et al., 2001; Tan and Zou, 2001). Therefore, the objectives of the present study were: (1) To isolate endophytic fungi from an important medicinal tree, namely, *Melia azedarach* L. (Meliaceae), which is known for its richness in limonoids and terpenoids compounds, and (2) To study the role of the isolated endophytic fungi in plant defense against phytopathogenic fungi in vitro and under greenhouse condition and their effects on plant growth and mineral nutrient uptake in phosphorus deficient soil.

**MATERIALS AND METHODS**

**Isolation of Endophytic Fungi**

Samples of healthy leaf of *M. azedarach* were collected from Faculty of Agriculture Garden, Alexandria (31° 12’ 56.3” N, 29° 57’ 18.97” E.) Egypt. Leaves were washed by tap water, followed by distilled water, then, surface sterilized by submerging the whole leaf in 90% ethanol for 1 minute, followed by 3.0% sodium hypochlorite for 3 minutes. Then, the leaves were rinsed in three changes of sterile/distilled water for 1 minute each. The sterilized samples were cut into 5 mm² pieces and placed in Petri plates containing Potato Dextrose Agar (PDA) media with 50 mg L⁻¹ of ampicillin to suppress the bacterial growth. Petri plates were sealed and incubated at 25±2°C and checked on alternate days until the endophytes started to emerge from the leaf samples. The growing endophytes were subcultured onto plates containing PDA (Sunitha et al., 2013).

**Identification of Fungal Isolates**

The endophytic fungi were stained with lactophenol cotton blue and examined in 40× light microscope and were identified on the basis of colony characteristics and microscopic characters of the spores using standard manual (Barnett and Hunter, 1998).

The isolated endophytic fungi were subjected to DNA extraction, using the Qiagen DNA extraction kit (Qiagen, Germany). These fungi were identified by amplification of ITS1-4 gene using universal primers according to Hafez and Elbestawy (2009). ITS 1 (Forward primer: 5’ TCC GTA GGT GAA CCT GCG G 3’) and ITS 4 (Reverse primer: 5’ TCC TCC GCT TAT TGA TAT GC 3’). 25 µL of PCR reaction mixture contained 5 µL master mix, 1 µL forward primer, 1 µL reverse primer, 1µL DNA Template and 17 µL distilled water. The PCR amplified products were analyzed by gel electrophoresis at approximately 600–700 bp. The Internal Transcribed Spacer (ITS) of ribosomal DNA were amplified, sequenced, and the nucleotide sequences were compared with those sequences already deposited in the data bank of the National Center for Biotechnology and Information (NCBI) using the nucleotide Basic Local Alignment Search Tool (BLAST) to find the most closely related sequences. The identification of the species was determined based on the best sequence alignment score. The nucleotide sequences were deposited in NCBI nucleotide sequence databases to get accession numbers.

**In Vitro Antagonistic Activity of Endophytic Fungi by Dual Culture**

The antagonism between the isolated endophytic fungi to each other and to two phytopathogenic fungi was carried out by dual culture according to Fokkema (1978).
Phytopathogenic fungi; *Alternaria solani* (EMCC 756), *Fusarium oxysporum* (EMCC 137) were obtained from Microbiological Resource Centre (Cairo MIRCEN), Faculty of Agriculture, Ain Shams University, Cairo, Egypt. One mycelial plug (5 mm) of each endophyte and pathogen were placed on the same dish 5 cm from each other with three replicates of each treatment and incubated at 27ºC. The Percent of Growth Inhibition (PGI) was calculated using the following formula:

\[ \text{PGI} (\%) = \left( \frac{\text{KR} - \text{R1}}{\text{KR}} \right) \times 100 \]

Where, KR is the average of three colony diameters (mm) from the point of inoculation to the colony margin on the control dishes, and R1 is the colony diameters from the point of inoculation to the colony margin in the direction of the antagonist (Korsten *et al*., 1995). The PGI was categorized on a Growth Inhibition Category (GIC) scale from 0 to 4, where 0= No growth inhibition; 1= 1-25% growth inhibition; 2= 26-50%; 3= 51-75%; and 4= 76-100%. Inhibition zone was recorded as the distance between the two fungal growths after seven days.

**Characterization of Endophytic Fungi for Their Plant Growth Promoting Capability**

The endophytic isolates were cultured in Potato Dextrose Broth (PDB) amended with 0.5 mg mL\(^{-1}\) tryptophan for 2 weeks at 28ºC in shaker incubator at 150 rpm. The ability of endophytes to produce IAA was determined by colorimetric method as preliminary test and then a more reliable HPLC method was performed to quantify the IAA produced (Glickmann and Dessaux, 1995; Szkop and Bielawski, 2013). For the colorimetric method, the development of pink or red color after adding Salkowski’s reagent to the previous supernatant and incubation for 30 minutes indicates IAA production, and concentration of IAA produced by the isolates were calculated using the calibration curve of pure IAA as a standard. For HPLC method, 10 mL of endophytic culture broth was centrifuged at 15,000xg at 4ºC for 30 minutes. The supernatant was analyzed on HPLC using UV-detector at 245 nm and C-18 column. Methanol: water (75:25) mixture was used as mobile phase with flow rate of 1 mL min\(^{-1}\). The growth hormone was identified on the basis of retention time of the standard IAA.

**Phosphate Solubilization and Acids Production by Endophytes**

Both fungal endophytes were examined for their ability to solubilize insoluble phosphate by using Illmer media according to Illmer *et al.* (1995) with a modification in CaHPO\(_4\) concentration (CaHPO\(_4\) 2.4 g). The presence of clearing zones around the fungal colonies was considered as indicator for positive solubilization activity. Acid production of the endophytic isolates was detected according to Louw and Webley (1958). The two endophytes were cultured in media containing bromocresol purple to improve the clarity and visibility of the yellow colored halo zone around the positive colonies.

**Ammonia Production**

Two mL from the PDB growth media were taken and distilled using Kjeldahl
apparatus by adding 0.2 g MgO. The resulting ammonia was collected in boric acid with mixed indicator, then, the distillate was titrated with 0.01 molar H$_2$SO$_4$ and the ammonia concentration was calculated as µg ammonia per 1 mL from the PDB media.

**Growth Promoting Activity by Endophytic Fungi and Mineral Analysis**

A split plot experimental design with three replicates was conducted to assess the efficacy of the two fungal endophytes (*N. sphaerica* and *C. lunata*) as bio-inoculants in enhancing tomato growth in a Phosphorus (P) deficient soil. Main plots were assigned to test three mineral fertilizers levels: Nitrogen (N), P, and potassium (K) at three rates (zero, 50, and 100%) of the recommended dose, with or without the two fungal endophytes spore suspensions as subplots. All the pots contained one kilogram of unsterile calcareous soil (EC= 3.3 dS m$^{-1}$, CaCO$_3$ = 10%, N= 10 mg kg$^{-1}$, P= 12 mg kg$^{-1}$ and K= 250 mg kg$^{-1}$). The endophytes’ spores were added to the soil around tomato seedlings two times at the rate of 10 mL pot$^{-1}$, each addition contained 10$^7$ fungal spores mL$^{-1}$ of *N. sphaerica* or *C. lunata*. The first inoculation for each of the endophytic fungi was at the beginning of cultivation and the second one was in the middle of the plant growing period. The plants were harvested after 45 days of growth, fresh weight was measured, and then the whole plants were dried at 70°C for 48 hours until constant weight. Then, 0.1 g of dried ground plant was digested by using a mixture of concentrated sulfuric acid and hydrogen peroxide and then the total N was determined according to Bremner and Mulvaney (1982). Total P was determined calorimetrically according to Olsen and Sommers (1982), and K content was determined by flame emission spectrophotometry using a flame photometer according to Horneck and Hanson (1998).

**Evaluation of Antagonistic Activities of Endophytes under Glasshouse Conditions**

Endophytes and phytopathogenic fungi (*F. oxysporum*) were grown in PDA plates at 28°C for 1 week. The cultures were flooded with sterile distilled water, scrapped with surface sterilized spatula, and filtered through cheesecloth. Spore suspension concentration was determined and adjusted to 10$^7$ and 10$^5$ spore mL$^{-1}$ for endophytes and pathogenic fungi, respectively. One-week old tomato seedling was planted in 9 cm pots containing 1 kg of unsterile soil in a greenhouse under natural sunlight at 30±2°C. At the beginning of the experiment, ten mL of endophytes suspension were applied to soil surface. Ten days later, the soil was re-inoculated by endophyte suspension before adding pathogenic fungi to encourage plant–endophyte association. Two types of control treatments were used. In the first control treatment, neither endophytes nor phytopathogenic fungi were added to the soil, while in the second control treatment, 10 mL of *F. oxysporum* spore suspension (10$^5$ spore mL$^{-1}$) was added. Pots were arranged in a randomized design in glasshouse at 30±2°C with 12 hours photoperiod (3.3 µmol m$^{-2}$ s$^{-1}$) and were irrigated with 100% nutrients as per required for 45 days. Plants were daily observed and disease symptoms were recorded. Then, the plants were uprooted to measure seedling growth parameters and to calculate disease severity index. Disease severity was evaluated for each plant on a zero to 4 rating scale according to percentage of shoot and roots affected by necrosis, wilt, and dark brown colors (Zhang et al., 2012): 0= Healthy plants, 1= 1 to 33%, 2= 34 to 66%, 3= 67 to 97%, 4= Dead plants.

Disease severity (%)= \{Σ (No. infected plants×Their infected degree)/(Total examined tested plants×Upper infected degree)\}×100.
Phytohormones Producing Fungal Endophytes

**Statistical Analysis**

All treatments were performed in triplicate and all the values were reported as average of triplicate determination. Data of the nutrition experiment was analyzed using split plot design. According to Naessens et al. (1986), the difference among the treatments at the 0.05 significant level was determined by Least Significant Difference (LSD) test using SAS procedures. Inhibition percent of phytopathogenic fungus by the endophytic fungus was subjected to one-way analysis of variance followed by Student–Newman–Keuls test (Cohort Software Inc. 1985) to determine significant differences between mean values at the probability level of 0.05.

**RESULTS**

**Identification of Endophytic Fungi**

Under the light microscope, fungal isolates (Figure 1) were identified to genus level as Curvularia sp. and Nigrospora sp., by their sporulation structures on PDA growing medium.

In the present study, PCR amplification of the ITS regions resulted in fragments ranged in length between 650 to 700 bp (Figure 2). The NCBI nucleotide database matching results of the sequences obtained in this study indicated that the two examined isolates belonged to two different genera. The observed high identity results (100%) to database sequences allowed the authentication of the isolates at the species level as C. lunata (GenBank accession no. MF113056) and N. sphaerica (GenBank accession no. MF113055).

**Antagonistic Activities of Endophytic Fungi by Dual Culture**

Dual culture bioassay showed antagonistic activities of both endophytic fungi to each other and to the phytopathogenic fungi. C. lunata antagonistic activity was characterized by the existence of inhibition zone (Figure 4) with A. solani, F. oxysporum and N. sphaerica. The antagonistic effects of C. lunata ranged from intermediate antagonism of both Alt. solani and F. oxysporum to high antagonism of N. sphaerica (Table 6). N. sphaerica showed medium antagonism of both F. oxysporum and A. solani with no inhibition zone (Figure 3).

**The Ability of Endophytic Isolates to Produce IAA and Ammonia and to Solubilize Phosphorus**

N. sphaerica showed significant production of IAA 40 µg mL⁻¹ and had the
ability to dissolve insoluble phosphate and produce acids. In contrast, *C. lunata* failed to solubilize phosphate or produce acid, and produced lower amount of IAA 3 µg mL\(^{-1}\) (Figure 3). Both fungal endophytes were able to produce ammonia at nearly the same rate.

**Effect of Mineral Fertilizer and Endophytes on Tomato Growth**

Analysis of variance presented in Table 2 shows the effect of mineral fertilizer levels and the tested fungal endophytes on both fresh and dry weight of shoot and root of tomato plant. The data revealed that shoot and root weights were highly significantly affected by the two main studied factors, except in case of root fresh weight. On the other hand, the interaction between the factors was not significant for all the tested parameters, except root dry weight. Mean values of shoot and root fresh weight and shoot dry weight were affected by mineral fertilizer levels and increased significantly by increasing NPK rates in soil, reaching 30.76, 7.85, and 4.99 g per plant, respectively (Table 4). Table 5 shows the mean values for shoot fresh and dry weight as affected by the two tested endophytes; the superiority of *N. sphaerica* compared to *C. lunata* was observed.

**Nutrients Content in Tomato Plants**

<table>
<thead>
<tr>
<th>Effect</th>
<th>df</th>
<th>Mean squares</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Fresh weight (g)</td>
<td>Dry weight (g)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Shoot</td>
<td>root</td>
<td>Shoot</td>
<td>root</td>
<td></td>
</tr>
<tr>
<td>Mineral fertilizer levels (A)</td>
<td>2</td>
<td>222.08 ***</td>
<td>4.90 *</td>
<td>8.52 **</td>
<td>.83 ***</td>
<td></td>
</tr>
<tr>
<td>Added endophytes (B)</td>
<td>2</td>
<td>60.48 ***</td>
<td>2.38 ns</td>
<td>4.26*</td>
<td>0.72 ns</td>
<td></td>
</tr>
<tr>
<td>AxB</td>
<td>4</td>
<td>12.31 ns</td>
<td>2.25 ns</td>
<td>0.81 ns</td>
<td>0.18 ns</td>
<td></td>
</tr>
</tbody>
</table>

* Significant at 0.05 level of probability; ** Significant at 0.01 level of probability; *** Significant at 0.001 level of probability; ns: Non-significant, df= Degree of freedom.

**Figure 3.** HPLC chromatogram of: (a) Standard Indole acetic acid, (b) *Nigrospora sphaerica* filtrate, (c) *Curvularia lunata* filtrate.

**Figure 4.** Antagonistic effect of endophytic fungi by dual culture: (a) *Curvularia lunata* and *Nigrospora sphaerica*; (b) *C. lunata* and *Fusarium oxysporum*; (c) *C. lunata* and *A. solani*.

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Table 2. Means squares and level of significance of fresh and dry weight (g) of tomato plant as affected by mineral fertilizers levels and endophytes treatments.
The effect of mineral fertilizer and the two tested endophytes on the concentration of N in the shoot and root of tomato plant is presented in Table 3. Analysis of variance revealed highly significant effect with using the increasing rates of NPK fertilizers. The use of endophytes also gave highly significant effect on N concentration. It is clear that *N. sphaerica* was able to increase the N content of the tomato plant when compared to the control (no endophytes) and this increase was clearly evident in plant that obtained the fully recommended mineral fertilization. The interaction between the effect of mineral fertilization and the endophytes inoculation produced positive results with tomato leaf and root N content. This is clearly shown in Figure 5, which indicates that the inoculation of *N. sphaerica* with 50% of the recommended mineral fertilization has led to a significant increase in N content of leaves and root of tomato plants.

Phosphorus concentration was significantly increased in leaves and roots of tomato plants, when inoculated with the *N. sphaerica*, and the same result as for N was obtained when we added the full dose of the recommended mineral phosphorus fertilization. Our results showed that *N.

**Table 3.** Means squares and level of significance of total nitrogen, phosphorus, and potassium concentration in tomato plant as affected by mineral fertilizers levels and endophytes treatments.

<table>
<thead>
<tr>
<th>Effect</th>
<th>df</th>
<th>Nitrogen (%)</th>
<th>Phosphorus (%)</th>
<th>Potassium (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Shoot</td>
<td>Root</td>
<td>Shoot</td>
</tr>
<tr>
<td>Mineral fertilizer levels (A)</td>
<td>2</td>
<td>8.65***</td>
<td>2.38***</td>
<td>0.30***</td>
</tr>
<tr>
<td>Added endophytes (B)</td>
<td>2</td>
<td>0.37***</td>
<td>0.47***</td>
<td>0.01*</td>
</tr>
<tr>
<td>A*B</td>
<td>4</td>
<td>0.05*</td>
<td>0.09*</td>
<td>0.003 ns</td>
</tr>
</tbody>
</table>

* Significant at 0.05 level of probability; ** Significant at 0.01 level of probability; *** Significant at 0.001 level of probability; ns: Non-significant. df= Degree of freedom.

**Table 4.** Means of fresh and dry shoot weights, phosphorus and potassium concentration in shoot of tomato plant as affected by mineral fertilizers levels.

<table>
<thead>
<tr>
<th></th>
<th>Shoot weight (g)</th>
<th>Phosphorus (%)</th>
<th>Potassium (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Shoot</td>
<td>Root</td>
<td>Shoot</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0 (%)</td>
<td>21.12 c</td>
<td>6.59 b</td>
<td>3.10 b</td>
</tr>
<tr>
<td>50 (%)</td>
<td>23.86 b</td>
<td>6.56 b</td>
<td>3.65 b</td>
</tr>
<tr>
<td>100 (%)</td>
<td>30.76 a</td>
<td>7.85 a</td>
<td>4.99 a</td>
</tr>
<tr>
<td>LSD (0.05)</td>
<td>2.4</td>
<td>1.11</td>
<td>1.09</td>
</tr>
</tbody>
</table>

* Means in the same column followed by the same letters are not significantly different at 0.05 level of probability. LSD$_{0.05}$= Least Significant Difference at 0.05 level probability.

**Table 5.** Means of shoot fresh and dry weight and phosphorus content of tomato plant as affected by endophytes treatments.

<table>
<thead>
<tr>
<th>Endophytes treatments</th>
<th>Shoot fresh weight (g)</th>
<th>Shoot dry weight (g)</th>
<th>Endophytes treatments</th>
<th>Phosphorus (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No endophytes</td>
<td>22.98 b</td>
<td>3.45 b</td>
<td>0.35 b</td>
<td>0.24 b endophytes</td>
</tr>
<tr>
<td>C. lunata</td>
<td>24.70 b</td>
<td>3.59 b</td>
<td>0.37 b</td>
<td>0.28 ab C. lunata</td>
</tr>
<tr>
<td>N. sphaerica</td>
<td>28.07 a</td>
<td>4.07 a</td>
<td>0.42 a</td>
<td>0.31 a N. sphaerica</td>
</tr>
<tr>
<td>LSD (0.05)</td>
<td>1.98</td>
<td>1.06</td>
<td>0.04</td>
<td>0.039 LSD (0.05)</td>
</tr>
</tbody>
</table>

* Means in the same column followed by the same letters are not significantly different at 0.05 level of probability. LSD$_{0.05}$= Least Significant Difference at 0.05 level probability.
**Figure 5.** Effect of three mineral fertilizers levels and the two tested endophytes on: (a) Nitrogen concentration in root, (b) N concentration in shoot. Bars marked with the same capital letter indicate no significant difference among the tested endophytes treatments under the same NPK level. Bars marked with the same small letter indicate no significant difference among the three mineral fertilizers levels for the same endophytes treatments.

`sphaerica` had the highest phosphate solubilization capability.

Concerning K, addition of the recommended complete mineral dose resulted in a significant increase in the concentration of the plant tissues; 37% higher than the plants fertilized with half the dose and three fold of the plants that were not fertilized by this element. In spite of the positive effect of the addition of K fertilization on the tomato plant, inoculation with both fungal endophytes did not have any positive or significant effect on its concentration in tomato leaves (Table 3).

**Table 6.** Antagonistic activities of endophytic fungi to each other and to phytopathogenic fungi by dual bioassay.

<table>
<thead>
<tr>
<th>Fungus</th>
<th><em>Curvularia lunata</em> (MF113056)</th>
<th><em>Nigrospora sphaerica</em> (MF113055)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Phytopathogenic fungi</td>
<td>Phytopathogenic fungi</td>
</tr>
<tr>
<td></td>
<td><em>C. lunata</em></td>
<td><em>N. sphaerica</em></td>
</tr>
<tr>
<td></td>
<td>PGI (%)</td>
<td>GIC</td>
</tr>
<tr>
<td><em>Alternaria solani</em></td>
<td>56</td>
<td>3</td>
</tr>
<tr>
<td>(EMCC 756)</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Fusarium oxysporum</em></td>
<td>50</td>
<td>2</td>
</tr>
<tr>
<td>(EMCC 137)</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Nigrospora sphaerica</em></td>
<td>70.2</td>
<td>3</td>
</tr>
<tr>
<td>(MF113055)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Antagonistic Activities of Endophytic Fungi against *F. oxysporum* (EMCC 137) under Greenhouse Condition**

Under pots condition, phytopathogenic fungi *F. oxysporum* caused significant reduction in shoot and root lengths, older leaves turned brown, and the plant frequently wilted. A cross section of the root showed that plants inoculated with *F. oxysporum* alone were infected with the pathogen and vascular tissue was brown and disease severity percent was 50%.
Interestingly, treatment of soil by *N. sphaerica* endophyte suppressed the infection and had 10% disease severity and increased root and shoot lengths of tomato plants (Table 7). *C. lunata* was less antagonistic to *F. oxysporum* under glasshouse conditions, causing disease severity of 30%, although it exhibited high antagonistic activity against *F. oxysporum* under *in vitro* dual culture bioassay.

**DISCUSSION**

To the best of our knowledge, this is the first isolation of *Curvularia lunata* as an endophytic fungus from *M. azedarach*. However, *C. lunata* was isolated before from *Azadirachta indica* by Verma et al. (2011) and from *Cymbopogon caesius* grass by Avinash et al. (2015). *N. sphaerica* was isolated from *Moringa oleifera* by Zhao et al. (2012), from *Melia azedarach* by Dos Santos and Rodrigues (2003) and from *Indigofera suffructiosa* by Santos et al. (2015).

This study showed the importance of using *N. sphaerica* and its ability to increase tomato plant biomass and its capability to reduce the infection rate of wilt disease caused by *F. oxysporum* in tomato plants under glasshouse conditions. The positive effect of *N. sphaerica* on fresh and dry weights of tomato might be attributed to its ability to produce plant growth promoters such as IAA, producing ammonia and acids, and solubilizing phosphorus, which can increase plant growth and minerals uptake. This result was supported by the findings of Glick et al., (2007), who found that increasing IAA and decreasing ethylene level increased root and shoot length as well as weight of the tested plant. In addition, Saad El-Din (2017) studied the effect of endophytes isolated from the medicinal plant (*Teucrium polium* L.) as bio-inocula for maize plant. The result showed higher fresh and dry weight for the inoculated plant and concluded that endophytes directly promote plant growth through the production of plant hormones, particularly IAA. Fungal endophytes can also help plants to access insoluble P through excretion of protons or enzymatic production that solubilize insoluble P and significantly improve plant growth (Oteino et al., 2015; Taktek et al., 2017). This may be due to the close linkage of endophytes inside plant tissues, which facilitates nutrients exchange and enzymes activity (Matsuoka et al., 2013; Murphy et al., 2014).

Many studies suggest that endophytes can mitigate disease symptoms of phytopathogenic fungi and improve plant growth by numerous modes (Shahzad et al., 2016; Egamberdieva et al., 2017). Although the mechanism is not fully understood, results of those studies suggest that pre-inoculation of plant root system by endophytes can interfere with the early infection processes and disease development by pathogenic fungi. Moreover, production of ammonia by fungal endophytes not only provides plants with N or decreases the cost of crop production but also intensifies the plant defense against phytopathogens colonization (Li et al., 2016). Endophytes can also increase P solubilization and nutrients uptake, which help the host plant to

**Table 7. Effects of antagonistic endophytes (*Curvularia lunata* and *Nigrospora sphaerica*) on seedling growth of tomato plant inoculated with *F. oxysporum* in greenhouse experiment.**

<table>
<thead>
<tr>
<th>Fungi treatments</th>
<th>Shoot length (cm)</th>
<th>Root length (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>45.0±1.00 a</td>
<td>20.0±0.00 a</td>
</tr>
<tr>
<td><em>F. oxysporum</em></td>
<td>41.5±1.50 b</td>
<td>10.5±2.50 c</td>
</tr>
<tr>
<td><em>C. lunata</em> + <em>F. oxysporum</em></td>
<td>41.5±1.50 b</td>
<td>13.5±2.50 c</td>
</tr>
<tr>
<td><em>N. sphaerica</em>+ <em>F. oxysporum</em></td>
<td>44.5±1.50 a</td>
<td>16.5±1.50 b</td>
</tr>
</tbody>
</table>

* Data are expressed as means±SE from experiments with three replicates. Means within a column sharing the same letter are not significantly different at the 0.05 probability level.
resist disease infection (Servin, 2004; Ongena and Jacques, 2008; Mei and Flinn, 2010; Zhao et al., 2014; Lecomte et al., 2016; Torres et al., 2016). This was the case with *N. sphaerica*, which had the ability to solubilize phosphorus, introducing acids and producing good amount of ammonia and IAA, so, all this may help tomato plant to resist *F. oxysporum* infection.

Although the laboratory tests proved that *C. lunata* had antagonistic activity against phytopathogenic fungi and it can produce ammonia and IAA hormone, we conclude that its poor results as a bio-control agent, or as plant growth promoter, when used in soil may be due to the lack of adaptation to high pH and lack of P availability in the calcareous soil used in this study. Concerning this issue, we propose testing *C. lunata* on different soil types to determine the optimum conditions for more efficient uses of this fungus. In contrast, *N. sphaerica* proved to be more adaptable and well suited as a bio-control agent and vital fertilizer for this type of soil. However, we cannot say by this study whether these endophytes isolated from *M. azedarach* colonized tomato plants and acted as endophytes or acted as rhizospheric organisms, though it enhanced tomato plant growth under these environmental conditions. As previous studies have found, some endophytes are able to colonize a wide range of host plants while others are specialized to one or a few hosts and some endophytes could colonize above- and below-ground plant tissues, i.e., the rhizosphere and aerial tissues (Zabalgozeazcoa, 2008; Rodriguez et al., 2009). This plant-endophyte interaction varies from mutualism to pathogenicity; it depends on a set of abiotic and biotic factors (Hardoim et al., 2015). Therefore, more studies are needed to focus on establishing the colonizing ability of these endophytes to understand the symbiotic relationship between endophytes and their host plant and other plant species under different environmental conditions. Further studies are required to isolate secondary metabolites, which may be responsible for the antifungal activities of these endophytic fungi, and to evaluate the potential of co-inoculation of the two endophytes and their effect on phytopathogenic microorganisms and plant growth under green house and field conditions.

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**REFERENCES**


قارچ های درون رست تولید کننده هورمون های گیاهی باعث بهبود وضعیت تغذیه و جلوگیری از عفونت قارچی گوجه فرنگی می‌شوند.

م. م. گ. سعد، و ه. بدري

چکیده

قارچ درون رست بیماری‌هایی در می‌آورد مثلا Curvularia lunata و Nigrospora sphaerica ساکن در درخت بیماری‌های گیاهی و بنابراین بهبود وضعیت تغذیه و جلوگیری از عفونت قارچی گوجه فرنگی می‌شوند. N. sphaerica از نگاه انتخابی آنتانگونیست‌شیک‌شان در داده‌های مایه DNA sequencing و PCR در حالت PCR مشخص شد. در تحقیق کلی، C. lunata با ترکیبی ۵۶٪ بهره‌برداری و Alternaria solani و Fusarium oxysporum با حذف ۴ تا ۶۳٪ از آسیب در حالت N. sphaerica و C. lunata به ترتیب تاثیر گذار می‌گردند. در ارتباط با وضعیت تغذیه گیاه گوجه فرنگی، در تیماری که ۵۰٪ کود معدنی توصیه شده مرکب بود، هم N. sphaerica و F. oxysporum در حالت N. sphaerica در درخت C. lunata با ترکیبی ۴۰٪ بهره‌برداری و اثرات مثبت می‌تواند به رشد گیاه گوجه فرنگی داشته. نتایج این‌گونه نشان دهنده استفاده از قارچ درون را به عنوان یک کود زیستی و کنترل کندنی‌هایی در شرایط گلخانه به خوبی نشان می‌دهد.