Influence of Culture Media and Temperature on Growth and Sporulation of Lasiodiplodia theobromae, Pestalotiopsis microspora and Fusarium oxysporum Isolated from Ricinodendron heudelotii in Cameroon

Joseph Djeugap Fovo*, Daniel Dostaler and Louis Bernier

1Phytopathology Laboratory, Department of Plant Protection, Faculty of Agronomy and Agricultural Science, University of Dschang, Box 222 Dschang, Cameroon
2Plant Pathology Laboratory, Department of Phytology, Faculty of Agricultural Science and Food, Université Laval, G1V 0A6 Qc, Québec, Canada
3Forest Pathology Laboratory, Department of Wood science and Forest, Faculty of Forestry, Geography and Geomatic, Université Laval, G1V 0A6 Qc, Québec, Canada

*Corresponding author

A B S T R A C T

The effects of mycological media and temperature on mycelial growth and spore production of three fungal pathogens recently reported on Ricinodendron heudelotii were investigated. The pathogens were identified on the basis of the ITS sequences of their ribosomal DNA as Pestalotiopsis microspora (isolate PMHP_109L), Lasiodiplodia theobromae (isolate LTHP_110L) and Fusarium oxysporum (isolates FOBR_164S, FOBR_049L and FOHP_121L). The radial growth (mm/day) of the fungi and conidia concentrations (number of conidia/ml of suspension) were assessed on three culture media: potato dextrose agar (PDA), malt extract agar (MEA) and V8 juice agar (V8). All media were suitable for the growth of L. theobromae while V8 juice agar (V8) supported the fastest mycelial growth rate in P. microspora (13.4 mm/day). PDA and MEA media were appropriate for F. oxysporum. The growth rate and conidia concentration increased with temperature and attained their optimum at 23°C for P. microspora and L. theobromae and 28°C for Fusarium oxysporum strains. There was no growth of P. microspora and L. theobromae at 33°C while at this temperature, Fusarium oxysporum strains continued to grow and produce spores. The best temperature for spore production was 23°C for P. microspora and 28°C for F. oxysporum and 21, 23 and 28°C for L. theobromae. F. oxysporum isolates produced the highest concentration of conidia in all the culture media. These data contribute to the knowledge of the biology of these newly recognized parasitic fungi on R. heudelotii and show that species like F. oxysporum possesses a high level of phenotypic plasticity that allows it to survive and proliferate over a wide range of environmental conditions.

Keywords
Pestalotiopsis microspora, Lasiodiplodia theobromae, Fusarium oxysporum, Culture media, Temperature, Mycelial growth, Sporulation, Ricinodendron heudelotii.

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Introduction
Ricinodendron heudelotii Pierre ex Heckel belongs to family Euphorbiaceae and is endemic to Madagascar and the Congo Basin forests of Africa. Its roots, bark, leaves and seeds are used by local people for medicine and food (Yeboah et al., 2011). It was also established that the species has a high agroforestry potential (Djeugap et al., 2013)
and yearly income from the sale of its edible grains in Cameroon is estimated to 1,556,280 US dollars (Perez and Ndoye, 1999). In Cameroon, many fungi cause various diseases on Ricinodendron heudelotii in forests, agroforestry systems and nurseries. Disease symptoms commonly encountered include leaf spots, leaf and seed rot and shoot blight (Djeugap, 2013; Djeugap et al., 2016). Despite these food, medicinal and agroforestry properties, very little work has been devoted to the study of pathogens responsible to these diseases. It is known that fungi can grow and reproduce in or on diverse culture media requiring several specific nutrient elements. They are isolated on specific culture medium for cultivation, preservation, microscopic examination and biochemical and physiological characterization. Also, a wide range of media are used for isolation of different groups of fungi that influence the vegetative growth and colony morphology, pigmentation and sporulation depending upon the composition of specific culture medium, pH, temperature, light and water availability (Kuhn and Ghannoum, 2003; Kumara and Rawal, 2008). However, requirements for fungal growth are generally less restrictive than for sporulation. The aim of this work was to assess the growth and sporulation of three cosmopolitan and polyphagous fungi, namely Pestalotiopsis microspora, Lasiodiplodia theobromae and Fusarium oxysporum, recently isolated from R. heudelotii in Cameroon (Djeugap, 2013) on three culture media at five temperatures. Several studies have shown that P. microspora is responsible for leaf spots, dieback and fruit rots in various plant species around the world (Keith et al., 2006; Djeugap et al., 2009; Zhang et al., 2010). The pathogen L. theobromae infects more than 500 plant species on which it causes root and fruit rots, shoot blight, dieback and canker (Mohali et al., 2005; Gezahgne et al., 2014; Djeugap et al., 2016). It was established that F. oxysporum is responsible for diseases such as seed rots and wilt in many crop and woody plant species (Tantawi and Fernandez, 1993; Hussain et al., 2012). This study was conducted with a view to contribute to the knowledge of the biology of P. microspora, L. theobromae and F. oxysporum, three newly recorded fungi isolated from R. heudelotii.

Materials and Methods

Sample collection, isolation and identification of fungi

Infected leaves and stems of R. heudelotii seedlings were collected in natural forests and nurseries of the World Agroforestry Centre in Cameroon. Isolation of fungi was carried out on potato dextrose agar medium (PDA) supplemented with ampicillin at 250mg/ml at 22°C for 10 days (Djeugap, 2013). Then, the pure cultures were transferred on PDA plate with cellophane membrane for 10 days. The mycelia were harvested, frozen in liquid nitrogen and crushed to a fine powder. Genomic DNA was extracted using the DNA Mini Kit Plant protocol (Qiagen) (Griffin et al., 2002; Levy and Mavrodieva, 2004), resuspended in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH 8) and the concentration of DNA was determined with a spectrophotometer. Purified diluted DNA was used as template for the amplification of the internal transcribed spacer (ITS) of nuclear ribosomal RNA gene (rDNA) repeats with the universal ITS4 (TCCTCCGCTTATGATATGC) and ITS5 (GGAAGTAAAA GTCGTAACAAGG) primers (White et al., 1990). The total volume of the PCR reaction was 25µl made up of 1.2µl of each primer, 12.5µl of Premix ExTaq, 5µl of diluted DNA and 5.1µl of double sterile distilled water. Amplification was performed in a PTC-225 Thermal Cycler (MJ Research, MA, USA) as follows: DNA denaturation (94°C, 2min); annealing (70°C, 2min) and final extension
(72°C, 10min) with 35 cycles. Sequencing by the Sanger method was carried out in a Life Technology 3139 XL type sequencer with 16 capillaries. DNA sequences were compared to those of the Genbank collection by using the WU-BLAST (Washington University-Basic Local Alignment Search Tool) algorithm (Altschul et al., 1997).

**Morphological and cultural characteristics of mycelium**

Morphological characteristics of the fungi were evaluated on PDA, malt extract agar (MEA) and V8 juice agar (10%) medium. PDA and V8 juice media were prepared based on Rodrigues et al., (2010) protocol while Pradeep et al., (2013) protocol was used for preparation of MEA medium. A volume of 20 ml sterilized medium was poured in each Petri plate and allowed to solidify. Then, 5mm-diameter plugs were taken with the help of a cork borer from the margin of 7-days old isolates grown on PDA, and placed at the centre of each set of Petri plates containing different media. Petri dishes were incubated at 10, 23, 28 and 33°C. The diameter (in mm) of each isolate was recorded in two directions at right angles to each other and then average colony diameter was calculated.

Growth was measured daily until the full expansion of the culture using the formula: 
\[ DG = \frac{\sum_{n=1}^{\infty} (MDd_s - MDd_0) + (MDd_2 - MDd_1) + \ldots + (MDd_{n-1} - MDd_n)}{n} \]
where DG is the daily growth rate, MDd is the mean diameter growth of the nth measurement day, MDd = (d1 + d2)/2 where d1 is the first diameter measure on Petri dish and d2, the second. MDd = initial diameter of the mycelium disk which is 5 mm (Pandey et al., 1985; Singh et al., 1993). The appearance of colonies and pigmentation, the vegetative and reproductive structures were described after 10 days of incubation. The experiment was repeated three times for each culture medium and isolate.

**Microscopic characteristics of mycelium and conidia**

Microscopic observations of mycelium and conidia of each fungus on each culture medium and temperature was made using a Olympus BX-41 microscope (Carsen, Markham, ON, Canada) and photographs were taken with a digital camera (Media Cybernetics, Evolution model VF) connected to a computer. A 10 ml conidial suspension was prepared by pouring sterile distilled water on 10-days old culture in Petri dishes. The concentration of spores (number of conidia/ml of suspension) was assessed for each culture medium and temperature from a drop of conidia suspension deposited on the hemocytometer. This activity was repeated three times. The size (width and length) of spores was measured on 30 conidia of each species randomly selected using the graduated dial of the ocular of a light microscope and the number of cells per conidia was counted (Bakry et al., 2010). Morphological characteristics of spores were compared with those described in the literature (Pavlic et al., 2004; Hussain et al., 2012; Rahman et al., 2012).

**Statistical analysis**

Data obtained were subjected to one way analysis of variance (ANOVA) and means compared by Student Least Significant Difference (LSD) test at 5% using SAS software (version 9.2).

**Results and Discussion**

**Sequencing and BLAST**

Data obtained from sequencing are presented in table 1. Isolates used are from bimodal rain forest and high plateau agroecological zones collected either on infected leaves or stem of *R. heudelotii* in crop farm, ICRAF plantation,
The size of conidia varied among isolates of *F. oxysporum*. The width and length of microconidia and macroconidia for FOBR_049L isolate on PDA varied from 2.2-3.9 µm x 3-17.4 µm and 3.7-4.2 µm x 26-28.5 µm, respectively. Isolate FOHP_121L produced longer macroconidia than other isolates (Table 3). Microconidia in *F. oxysporum* are mono or multicellular (2-3 cells) while macro conidia are multicellular (4-10 cells). There was abundant production of chlamydospores in FOBR_049L isolate in V8 juice agar (Figure 3). The number of cells reached 10 macroconidia in FOHP_121L while the maximum was 7 in FOBR_049L and FOBR_164S (Figures 4, 5 and 6). Isolate FOHP_121L produced microconidia and macroconidia that were longer and wider than those of FOBR_049L and FOBR_164S. The width and length of microconidia of FOHP_121L ranged from 2.2-4.4 µm x 6-12.3 µm, 2.3-4.6 µm x 7-23.5 µm and 2-3.6 µm x 6.3-11 µm on PDA, MEA and V8 juice agar media, respectively. The size of macroconidia ranged from 3.1-5.3 µm x 26-42.2 µm, 4.2-6.5 µm x 25.6-43 µm and 2.9-4.9 µm x 26.8-39.3 µm on PDA, MEA and V8 juice agar media, respectively (Table 3). On PDA, conidia of *P. microspora* were hyaline, bi- to tetra cellular with a cylindrical carrot-like shape; the tapered lower end terminated in a filamentous appendage while the more rounded upper end was extended by 2, 3 or 4 filaments (Fig. 7A). Conidia of *L. theobromae* were larger and brown with a peanut pod-shaped (Fig. 7B). The width and length of the conidia of *L. theobromae* and *P. microspora* are relatively thicker and longer on MEA medium compared to other media. In fact, conidia size of *L. theobromae* varied from 10.4 to 15.7 µm x 25.3 to 30.2 µm, 11 to 17 µm x 22.1-35.3 µm and from 9.9-14.7 µm x 18-27.8 µm on PDA, MEA and V8 juice agar media.
respectively while in \( P. \text{microspora} \), they varied from 6.6 -7.8 µm x 18.7 -29 µm, 8 -12.9 µm x 22.1 - 38.6 µm and from 6.5 -8.8µm x 20.5 - 28 µmon PDA, MEA and \( V_8 \) juice agar media, respectively (Table 4).

**Effect of temperature on the concentration of fungal spores**

Temperature affected the concentration of fungal spores produced on all three culture media. In general, spore concentration increased with temperature until an optimum (28°C), and then decreased until no more spore production occurred at 33°C for \( P. \text{microspora} \) and \( L. \text{theobromae} \). However, \( F. \text{oxysporum} \) isolates continued to grow and produce spores at 33°C. Significant differences in spore concentration were observed between 28°C and the other temperatures (10, 23 and 33°C) for isolates FOBR_049L and FOHP_121L of \( F. \text{oxysporum} \). In contrast, no significant difference was observed in spore’s concentration of FOHP_164S at 23 and 28°C on the MEA and \( V_8 \) juice agar media. There was also no significant difference in spore concentration between 21, 23 and 28°C for \( P. \text{microspora} \) on PDA and for \( L. \text{theobromae} \) on PDA, MEA and \( V_8 \) juice agar media.

### Table.1 Ecological and molecular characteristics of fungal isolates used in the study

<table>
<thead>
<tr>
<th>Isolate code</th>
<th>Agro ecological zone</th>
<th>Locality</th>
<th>Ecological niche</th>
<th>Plant organ</th>
<th>ITS/BLAST(^a)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LTHP_110L</td>
<td>High Plateau</td>
<td>Bantoum III</td>
<td>Crop farm</td>
<td>Leaves</td>
<td>FJ904838.1</td>
</tr>
<tr>
<td>PMHP_109L</td>
<td>High Plateau</td>
<td>Bantoum III</td>
<td>Crop farm</td>
<td>Leaves</td>
<td>AF377292.1</td>
</tr>
<tr>
<td>FOBR_049L</td>
<td>Bimodal Rainforest</td>
<td>Yaoundé 7e</td>
<td>ICRAF plantation</td>
<td>Leaves</td>
<td>HM179532.1</td>
</tr>
<tr>
<td>FOBR_164S</td>
<td>Bimodal Rainforest</td>
<td>Yokadouma</td>
<td>Cocoa farm</td>
<td>Stem</td>
<td>HQ658965.1</td>
</tr>
<tr>
<td>FOHP_121L</td>
<td>High Plateau</td>
<td>Dschang</td>
<td>ARF, FASA</td>
<td>Leaves</td>
<td>GU724513.1</td>
</tr>
</tbody>
</table>

\(^a\) Sequences from public genbank (NCBI) with highest similarity with the sequence of fungal isolates from \( R. \text{heudelotti} \). L=isolated from leaves and S= isolated from stem (seedlings). AFR, FASA= Applied Research Farm of the Faculty of Agronomy and Agricultural Science.

### Table.2 Effect of culture media on the daily growth rate (mm/d) of fungal isolates incubated at 23°C

<table>
<thead>
<tr>
<th>Isolate code and fungal species</th>
<th>Culture media</th>
<th>PDA</th>
<th>MEA</th>
<th>( V_8 ) (10%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LTHP_110L ( (Lasiodiplodia theobromae) )</td>
<td>21.6 ± 2.3(^a)</td>
<td>20.6 ± 1.2(^a)</td>
<td>19.8 ± 0.9(^a)</td>
<td></td>
</tr>
<tr>
<td>PMHP_109L ( (Pestalotiopsis microspora) )</td>
<td>10.0 ± 1.1(^b)</td>
<td>9.7 ± 1.4(^b)</td>
<td>13.4 ± 1.7(^a)</td>
<td></td>
</tr>
<tr>
<td>FOBR_049L ( (Fusarium oxysporum) )</td>
<td>8.4 ± 1.2(^a)</td>
<td>8.1 ± 0.8(^a)</td>
<td>6.3 ± 0.8(^b)</td>
<td></td>
</tr>
<tr>
<td>FOBR_164S ( (Fusarium oxysporum) )</td>
<td>7.8 ± 0.9(^a)</td>
<td>8.6 ± 1.6(^a)</td>
<td>6.9 ± 0.6(^a)</td>
<td></td>
</tr>
<tr>
<td>FOHP_121L ( (Fusarium oxysporum) )</td>
<td>9.8 ± 1.9(^a)</td>
<td>9.1 ± 0.7(^a)</td>
<td>7.0 ± 1.2(^b)</td>
<td></td>
</tr>
</tbody>
</table>

\(^a,b\) Means follow by the same letter in the line are not significantly different based on Student’s LSD at 5%. LT: \( L. \text{theobromae} \), PM: \( P. \text{microspora} \) and FO: \( F. \text{oxysporum} \). HP: High Plateau, and BR: Bimodal Rainforest. L: isolate from leave and S: isolate from stem.
Table 3 Micro and macro conidia sizes (µm) of isolates of *F. oxysporum* on PDA milieu incubated at 21±1°C from 10-days old culture

<table>
<thead>
<tr>
<th>Isolates of <em>F. oxysporum</em></th>
<th>PDA</th>
<th>MEA</th>
<th><em>V₈</em></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Microconidia</td>
<td>Macroconidia</td>
<td>Microconidia</td>
</tr>
<tr>
<td></td>
<td>Width</td>
<td>Length</td>
<td>Width</td>
</tr>
<tr>
<td>FOBR-049L</td>
<td>2.2-3.9</td>
<td>3-17.4</td>
<td>3.7-4.2</td>
</tr>
<tr>
<td>FOBR-164S</td>
<td>1.7-3.1</td>
<td>4.6-19</td>
<td>3-4.4</td>
</tr>
<tr>
<td>FOHP-121L</td>
<td>2.2-4.4</td>
<td>6-12.3</td>
<td>3.1-5.3</td>
</tr>
</tbody>
</table>

Table 4 Conidia sizes (µm) of *Lasiodiplodia theobromae* and *Pestalotiopsis microspora* on PDA, MEA and *V₈* media incubated at 21±1°C from 10 days-old culture

<table>
<thead>
<tr>
<th>Isolates</th>
<th>PDA</th>
<th>MEA</th>
<th><em>V₈</em></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Width</td>
<td>Length</td>
<td>Width</td>
</tr>
<tr>
<td>LTHP-110 L <em>(Lasiodiplodia theobromae)</em></td>
<td>10.4-15.7</td>
<td>25.3-30.2</td>
<td>11-17</td>
</tr>
<tr>
<td>PMHP-109L <em>(Pestalotiopsis microspora)</em></td>
<td>6.6-7.8</td>
<td>18.7-29</td>
<td>8-12.9</td>
</tr>
</tbody>
</table>
**Figure 1** Variation of morphological characteristics of fungi on different culture media. A: FOBR_049L isolate (*F. oxysporum*), B: FO_BR164S isolate (*F. oxysporum*), C: FOHP_121L isolate (*F. oxysporum*), D: PMHP_109L isolate (*P. microspora*) and E: LTHP_110L isolate (*L. theobroma*). Fungi were incubated at 23°C during 6 days; arrows indicate fructifications.
Figure 2 Variation of morphological characteristics among *Fusarium oxysporum* isolates based on temperature of incubation. A: FOBR_049L, B: FOBR_164S and C: FOBR_121L. The culture medium in the 1st and last column is PDA while the medium in the 2nd and 3rd column is MEA.

Figure 3 Evolution of growth rate (mm/day) of fungi newly recorded from *Ricinodendron heudelotii* on PDA, MEA and V8 media at different incubation temperatures.
**Figure 4** Micro and macro conidia of *Fusarium oxysporum* (isolate FOBR_049L) on PDA and MEA media. Chlamydospores on V8_Agar medium from 14-day-old culture, 400X; gentian violet as mounting liquid.

**Figure 5** Micro and macro conidia of *Fusarium oxysporum* (isolate FOBR_164S), from infected stem of seedlings of *Ricinodendron heudelotii* incubated at 21°C on PDA, 400X; water as mounting liquid.

**Figure 6** Micro and macro conidia of *Fusarium oxysporum* FOHP_121L from infected leaves of seedlings of *Ricinodendron heudelotii* incubated at 10, 28 and 33°C on PDA, 400X; water as mounting liquid.

**Figure 7** Conidia of *Pestalotiopsis microspora* (A) and *Lasiodiplodia theobromae* (B) isolated from *Ricinodendron heudelotii* incubated at 23°C on PDA medium, 400X; water as mounting liquid.
Figure.8 Concentration of spores of fungi isolated from *Ricinodendron heudelotii* and incubated during 14 days on PDA, MEA and V₈ media. Means follow by the same letter in one fungal species are not significantly different based on Student’s test at 5%

However, spore concentration was significantly higher in these three temperatures and media compared to 10°C and 33°C for *L. theobromae* and *P. microspora* (Figure 8).

The study reveals marked differences in the mycelial growth and sporulation patterns for the three fungal species tested on three culture media that are routinely used in plant pathology laboratories. Culture medium and temperature affected the morphocultural...
characteristics of the fungi tested. Indeed, preferences of fungi vis-a-vis the culture medium and temperature varied among species and, within *Fusarium oxysporum*, from one isolate to another. Some fungi grow and sporulate better on organic media incubated at low or high temperature, others on more selective media. It is known that temperature affects the number of spores produced on the culture medium (Agrios, 2005; Rahman et al., 2012). Mycelial growth and sporulation were abundant on PDA, MEA and V₈ juice agar media. These media are organic substrates rich in carbohydrates which are source of energy for cell metabolism, growth and sporulation of fungi. Several workers have recognized the importance of various organic media for mycelial growth and spore production in fungi (Kim et al., 2005; Zhao et al., 2010; Pradeep et al., 2013). PDA is one of the most commonly used culture media because of its simple formulation and its ability to support mycelial growth of a wide range of fungi. Several researchers state that PDA is one of the best media for mycelial growth of many fungi followed by MEA (Xu et al., 1984; Maheshwari et al., 1999; Khadega and Al-Hussaini, 2015). V₈ juice agar was also reported as suitable for cultivation and sporulation of other fungal species like *Mycosphaerella fijiensis*, *M. musicola*, *Phomamacdon aldii* and *Phytophthora* spp (Medina and Platt, 1999; Drenth and Sendall, 2001; Abadie et al., 2005; Al Fadil, 2006; Tsompbeng et al., 2012). Morphological traits (mycelia appearance, conidium size and pigmentation of mycelium) observed in isolates of *F. oxysporum* from *R. heudelotii* are comparable at some extend to those reported in other *F. oxysporum* (Sharma et al., 2005; Hussain et al., 2012). Isolates of *F. oxysporum* from wilted guava grow and sporulate abundantly with production of chlamydospores on PDA medium at 20 and 25°C (Hussain et al., 2012) while corn meal agar medium was reported to be suitable for the growth of *Fusarium oxysporum* f. *psidii* on which length of microconidia and macroconidia reached 12.9 and 57.4 µm, respectively (Rahman et al., 2012).

The range of temperatures (from 10 to 33°C) at which isolates of *F. oxysporum* from *R. heudelotii* could grow and sporulate indicates that this pathogen possesses a high level of phenotypic plasticity that allows it to survive and proliferate over a wide range of environmental conditions. The optimal temperature of mycelial growth for *F. oxysporum* isolates was 28°C. This was also reported in *F. moniliforme* KUMB F 1201 isolate cultivated on PDA and potato dextrose broth media and isolated from paddy field soil in India (Pradeep et al., 2013) and in *Fusarium oxysporum* f. *sp. psidii* and *F. solani*, causal agents of wilt in guava cultivated on PDA (Gupta et al., 2010). However, it was established that *Fusarium* species like *F. oxysporum* f. *sp. narcissi* can growth at 45°C in liquid medium (Linfield, 1986).

Conidia sizes in *P. microspora* isolated from *R. heudelotii* are relatively similar to values reported for 16 isolates of the same fungus isolated from *Taxodium distichum* and incubated on PDA at 23°C (Li et al., 1996), and for isolates from *Psidium guajava* grown on PDA at 24°C (Keith et al., 2006). However, Keith et al. (2006) reported that all isolates of *P. Microspora* from *P. guajava* grew at temperatures ranging from 10 to 35°C and that the optimum growth temperature differed among isolates and varied between 22 and 28°C. In our study, *L. theobromae* and *P. microspora* failed to grow at 33°C probably due to the inactivation of enzymes by temperature increase with a resulting effect on metabolism which affects growth (Mishra et al., 2013; Pathak et al., 2014). The optimum growth temperature for
P. microspora isolate was 23°C on PDA, MEA and V₈juiceagar media. Morphocultural characters and sizes of conidia in L. theobromae are similar to those described by Pavlic et al., (2004) and Shah et al., (2010) for cultures isolated from infected cocoa (Theobromae cacao L.) and avocado (Perseaa mericana L.), respectively. Conidial sizes of L. theobromae ranged from 11-15 μm x20-35μmon MEA medium at 25°C (Pavlic et al., 2004).

For all the fungi, it was observed that as temperature increased, mycelial growth and spore production also increased but at 28°C they started to decline for F. oxysporum and 23°C for P. microspora and L. theobromae on PDA, MEA and V₈juice agar media. This could be attributed to increase in enzymatic activity of the fungi.

In conclusion, this study revealed that culture media differentially influence the growth, colony characteristics (physical aspect and pigmentation of the mycelium) and sporulation of P. microspora, L. theobromae and F. oxysporum recovered from R. heudelotii. Among the test media used in the present study, PDA and MEA were found to be most suitable for mycelial growth and sporulation of F. oxysporum while V₈juice agar was suitable for P. microspora and the three media were appropriate for L. theobromae. The optimum temperature for high production of spores was 28°C for F. oxysporum isolates and 23°C for P. microspora and L. theobromae. These data will be important for future large-scale studies involving cultural and morphological characterization of more isolates of these newly recorded fungi from R. heudelotii.

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