Inhibitory Effect of Some Spice Essential Oils on *Penicillium digitatum* Causing Postharvest Green Rot in Citrus

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**Abstract:** The aim of this study was to evaluate the antifungal activity of two species collected in different locations of Morocco against *Penicillium digitatum*, the causal agent of citrus green mold. The in vitro antifungal activity of essential oil was determined using the method of agar diffusion assay. The results show that essential oil of *Lavandula hybrida* has an inhibitory capacity of mycelial growth and spore production compared to others essential oils. *Penicillium digitatum*; *Lavandula hybrida*; in vitro; antifungal activity; *Artemisia herba alba*

**Keywords:** *Penicillium digitatum*; *Lavandula hybrida*; in vitro; antifungal activity; *Artemisia herba alba*

1. Introduction

In Morocco citrus play a socio-economic role that is very important. On the economic map, citrus exports is an important source of foreign exchange in the order of 2.5 to 3 billion dirham’s a year.

In citrus as in any other agricultural production sector commercial character to the final volume of the production is the most important parameter that determines the interest of the culture of a variety thus the improvement and mastery of production techniques and protection prove necessary for a better profitability.

The most important citrus disease that cause commercially significant losses in Morocco (Elkhmass et al.,1994) and worldwide (Eckert and Ears;1989;Holmes and Eckert,1999;Zhu et al 2006) are green mold, caused by *Penicillium digitatum*. The chemical control remains by far the control method preferred against all rot mainly due to *Penicillium Spp*. Intensive and repeated use of fungicide protection in post citrus fruits favored the recolte development resistant strains worldwide (Harding,1972; Smoot et brown, 1974) even to the consumer tends to look more natural products, has prompted the research, development and application of new natural products with antimicrobial activities in order to use them as alternative to synthesis products in the field of food industrial.

2. Materials and Methods

**Plant Material**

Species used in experiment are listed in table 1. They were collected from different locations between Meknes and Azrou. These plants have been chosen for their medicinal properties and because it is a very important floristic local heritage and it is largely described as a botanical point of view, the aerial part of the plant was dried in the shade at room temperature and subjected to a hydro distillation 2h.

**Table 1: Species used for distilling essential oils**

<table>
<thead>
<tr>
<th>Common name</th>
<th>Botanical name</th>
<th>Family</th>
<th>Used part</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lavender</td>
<td><em>Lavandula hybrida</em></td>
<td>Lamiaceae</td>
<td>Leaves+flowers</td>
</tr>
<tr>
<td>Artemisia</td>
<td><em>Artemisia herba alba</em></td>
<td>Astéracées</td>
<td>Leaves+flowers</td>
</tr>
</tbody>
</table>

**Return Calculation**

The yield of essential oil is defined as the ratio between the mass of E.O obtained after extraction (M’) and the mass of the plant material used (M) (Bssaisi F., Gmira Meziane N and M. (2009)). The yield is expressed in percentage and is given by the following formula:

\[
\text{Yield (\%)} = \frac{M'}{M} \times 100
\]

**Pathogen culture**

In this study the fungus employed for assays of antifungal activity was isolated from decayed oranges. The oranges were obtained from conditioning station. The identification was performed based on the study of macroscopic and microscopic characters of isolates growing on PDA medium. After identifying *Penicillium digitatum*, it was cultured routinely on Potato Dextrose Agar and incubated at 25°C.

**Evaluation of antifungal activity of essential oils:**

This effect was tested using aromatogram method that described by Maychiew and Devahastin (2008) and Hussain and colaboratory (2010), 20 ml PDA cooled are cast in Petri dishes. After solidification of the culture medium, 200 µl of the fungal suspension under test (107 spores.ml) were plated on the surface until total desiccation.

Under aseptic conditions and using sterile forceps, Paper filter discs (Whatman Nr 3, 6 mm diameter) (1 disc / Petri Dish)
dish) are sterilized and then deposit on the dried agar inoculated beforehand with the fungal suspensions then these discs are loaded by volumes of essential oil (10, 20, 30, 50, 60, 80μl) using a micropipette for each volume, 3 repetitions were performed to minimize the risk of error.

The Petri dishes are kept at 4°C for 2 hours so that the essential oil may diffuse into the culture medium. For control Petri plate had no essential oil.

The Petri plates were incubated at 25°C for 5 to 7 days until the growth in the control petri dishes reaches the edges.

**Determination of minimum inhibitory concentration (MIC)**

The minimum inhibitory concentrations (MIC) of essential oils were determined according to the method reported by Remmal et al. (1993) and Satrani et al. (2001). Due to the immiscibility of essential oils to the water and thus in the middle of culture, emulsification was carried out through a solution d’agar 0.2%. It provides, in the middle, an even distribution of essential oils to maximize the Contact germ/compound. Dilutions are prepared 1/10th, 1/25th, 1/50th, 1/100th, 1/200th, 1/300th and 1/500th in this agar solution.

In test tubes each containing 13.5 ml of medium on PDA (Potato Dextrose Agar) for fungi, autoclaved for 20 min at 121°C and cooled to 45°C, aseptically added 1.5 ml of each dilution to give final concentrations of 1/100, 1/250, 1/500, 1/1 000 1/2 000 1/3 000 1/5 000 (v/v). Stir the tubes to disperse the essential oil in the culture medium before pouring in the Petri disches.

Petri dishes control, containing the culture medium and the agar solution at 0.2% alone are also prepared.

3. Results and Discussion

1-Essential oil yield

The extraction of E.O by hydro distillation has been used for three plants. The figure shows the yields obtained by E.O extraction plants.

![Figure 1: Yield of E.O of Lavandula hybrida et Artemisia herba Alba](image)

This figure shows that the yield of *Lavandula hybrid* is 1.7% which is higher than that of *Artemisia* (0.8 %) The yield of HE, which depends on many factors (growth stage, soil and climatic conditions, extraction technique, harvest etc.) is only 0.8% for Artemisia alba. It can be much higher in the same species, which is about 1.23% (M.Ghanmi et al, 2010). In this sense Ghanmi confirms that the essential oil content of white mugwort change depending on the date of harvest.

2- Testing the antifungal activity:

![Figure 2: development of the zone of inhibition caused by the essential oil of Lavandula hybrida and Artemisia herba alaba](image)

*P. digitatum* was sensitive to the majority of the tested concentrations of both EO The figure below expresses the change in the average diameters of the inhibition zones (mm) depending on the concentration for the two essential oil, changes in the development of *P. digitatum* is inversely proportional to the increase of the concentration. Also comparing to the positive control (treatment with imazalil 50%) *Artemisia herba alba* has proved significantly more effective than *Lavandula hybrid* especially for the 60μl volume which notes that the inhibition zone and the same diameter as that of the positive control the same for the 80μl dose with a difference of 10 μl.
3-Determination de la CMI:

Table 1: The CMI of E.O of Lavendula and Artemesia herba alba

<table>
<thead>
<tr>
<th>Dilution</th>
<th>1/100</th>
<th>1/250</th>
<th>1/500</th>
<th>1/1000</th>
<th>1/2000</th>
<th>1/3000</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Penicillium digitatum</em></td>
<td>-</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>-</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Lavendula hybrida</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Artemesia alba</em></td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The inhibitory power of essential oils overlooked a microbial strain is classified as: excellent inhibitory power for MIC < 50 µL/ml to interesting inhibitor for CMI 50 µl/ml and 250 µl/ml, low inhibitory power for MIC 250 µl/ml and 500 µl/ml and to poor or no inhibitor for MICs > 500 µl/ml (Koba et al., 2004). It can be inferred that both essential oils have an interesting inhibitory power on the fungal strains.

4. Conclusion

According to these results we can predict that the essential oils of *Lavendula hybrida* and *Artemesia herba alba* are very effective natural antifungals and can be a very important source of herbal constituents used to eradicate infections of fungal origin.

References


[6] Sensitivity of *Penicillium digitatum* and *P. italicum* to Postharvest Citrus Fungicides in California Gerald J. Holmes and Joseph W. Eckert first author: Department of Plant Pathology, North Carolina State University, Raleigh 27695; and second author: Department of Plant Pathology, University of California, Riverside 92521


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