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Extracellular phospholipase activity of Malassezia strains isolated from individuals with and without dermatological disease

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ABSTRACT

Background: The Malassezia genus includes mainly lipophilic yeasts belonging to the cutaneous microbiota of man and other mammals. Some Malassezia species have been associated with various dermatological diseases. The factors permitting the transformation of yeasts of the Malassezia genus from a commensal organism to a pathogenic agent are still little known but the production of various enzymes such as lipase, phospholipase and lipooxygenase could contribute to the pathogenic activity of these yeasts.

Aims: Here we have determined and compared the extracellular phospholipase activity of sixty human isolates of Malassezia so as to relate this feature to the species of Malassezia and to the origin (from dermatological diseases or not) of the strains examined.

Methods: Phospholipase production was determined using the semi-quantitative egg-yolk plate method.

Results and conclusions: Malassezia obtusa, Malassezia slooffiae, Malassezia globosa, Malassezia restricta had difficulty developing in the chosen culture medium so that it was not possible to measure phospholipasic activity. Malassezia pachydermatis showed the highest phospholipase activity. Twenty-nine Malassezia sympodialis strains produced phospholipase; the isolates from patients with pityriasis versicolor had significantly higher phospholipasic activity than those isolated from healthy individuals. This observation suggests that the phospholipasic activity of Malassezia may play a role in the onset of skin lesions, especially in the case of pityriasis versicolor.

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The *Malassezia* genus includes mainly lipophilic yeasts belonging to the cutaneous microbiota of man and other mammals. These yeasts were first described by Eichstedt in 1846 as being associated with lesions of pityriasis versicolor.

The taxonomy and nomenclature of the *Malassezia* genus has, in recent years, been revised several times. In fact, up until 1995 only three species had been identified: *Malassezia furfur*, *Malassezia pachydermatis* and *Malassezia sympodialis*. The species *Malassezia globoza*, *Malassezia restricta*, *Malassezia obtusa*, and *Malassezia slooffiae* were described in 1995 on the basis of their morphology, ultrastructura, physiology and genomic features. In the last few years, the new lipid-dependent species *Malassezia dermatitis*, *Malassezia japonica*, and *Malassezia yamatoensis* were isolated from human skin. Other species were found on animals: *Malassezia nana* on cats and dogs, *Malassezia equina*, *Malassezia caprae* and *Malassezia cuniculi* found mainly on horses, goats and rabbits respectively.

Yeasts belonging to the *Malassezia* genus have been associated with various dermatological diseases: pityriasis versicolor (PV), dandruff, seborrheic dermatitis (SD), atopic dermatitis, folliculitis, psoriasis, onychomycosis, and blepharitis. The pathogenic role of *Malassezia* in PV has now been universally accepted although authors disagree about which species is most widely associated with the disease. Some of them suggest that *M. globoza* is the causal agent of PV, and *M. furfur* is the causal agent of SD.

As well as these diseases, cases of fungemia caused by *M. pachydermatis* and *M. furfur* have been reported in premature newborns and immunocompromised patients artificially fed with lipid emulsions.

Most *Malassezia* species are lipid-dependent; as a result these yeasts probably require lipolytic enzymes such as lipases and phospholipases to use the environmental lipids essential for their growth. In a recent work, Xu described the genome and secretory proteome of two species of *Malassezia*, suggesting that the lipid-dependency of these species is associated with the apparent absence of the gene for fatty acid synthase. Further studies have demonstrated extracellular production of phospholipase and lipase.

Phospholipases are a heterogeneous group of enzymes which hydrolyse one or more ester bonds of glycerol-phospholipids. Phospholipids and proteins are the main chemical constituents of the cellular membranes of the host so that the phospholipases and proteases are involved in the destruction processes of such cellular membranes. Many fungal species, such as *Candida albicans*, *Cryptococcus neoformans* and *Aspergillus fumigatus* can produce enzymes belonging to the phospholipase group. Researchers have associated some extracellular phospholipases with the virulence of *C. albicans*.

The factors permitting the transformation of yeasts of the *Malassezia* genus from a commensal organism to a pathogenic agent are still little known, but the production of various enzymes such as lipase, phospholipase and lipoxygenase could contribute to the pathogenic activity of these yeasts too.

Little information is available in literature regarding the phospholipasic activity of species of the *Malassezia* genus. Using the Price method (egg yolk medium), extracellular phospholipasic activity was demonstrated in vitro on strains of *M. furfur* isolated from skin lesions in humans. These works were prior to the classification introduced by Guého et al. so, bearing in mind the current classification, some strains included in these studies could be reclassified as different species.

The phospholipasic activity of *M. pachydermatis* has also been associated with skin lesions in dogs, but with infections by *M. sympodialis*.

In this work we have analysed and compared the extracellular phospholipasics of *Malassezia* isolates belonging to different species, identified according to the current classification so as to verify whether the various species of *Malassezia* have different phospholipasic activity and to relate this feature to the origin (from dermatological diseases or not) of the strains examined.

### Materials and methods

The strains included in this study were isolated from patients with PV or SD, both from skin lesions and from healthy skin, and from individuals without any dermatological disease. Fourteen reference strains from the Pasteur Institute (IP) and Central Bureau voor Schimmelcultures (CBS) (Table 1) were also examined.

<table>
<thead>
<tr>
<th>Species</th>
<th>Number of examined strains</th>
<th>PV patients skin with lesions</th>
<th>PV patients skin without lesions</th>
<th>SD patients skin with lesions</th>
<th>SD patients skin without lesions</th>
<th>Healthy individuals</th>
<th>IP or CBS</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>M. furfur</em></td>
<td>5</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. pachydermatis</em></td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. obtusa</em></td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. slooffiae</em></td>
<td>4</td>
<td>4</td>
<td>1</td>
<td>4</td>
<td>1</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. globosa</em></td>
<td>17</td>
<td>6</td>
<td>4</td>
<td>4</td>
<td>1</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. restricta</em></td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. sympodialis</em></td>
<td>31</td>
<td>4</td>
<td>3</td>
<td>2</td>
<td>4</td>
<td>15</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>60</td>
<td>10</td>
<td>3</td>
<td>8</td>
<td>5</td>
<td>20</td>
<td>14</td>
</tr>
</tbody>
</table>

CBS: Central Bureau voor Schimmelcultures; IP: Pasteur Institute; PV: pityriasis versicolor; SD: seborrheic dermatitis.
In addition, higher phospholipasic activity was shown in some strains. Similarly, it has been shown that the phospholipasic activity of the Malassezia species expressed as a Pz value mean after 15 and 20 days of culture is significantly higher in strains isolated from patients with PV and SD, from individuals with and without obvious skin lesions. Measuring of the precipitation zone taken at the day 15 and at day 20 from incubation were therefore deemed significant.

Table 2: Phospholipasic activity of the Malassezia species expressed as a Pz value mean after 15 and 20 days of culture.

<table>
<thead>
<tr>
<th>Species</th>
<th>Strains examined</th>
<th>Strains with positive development</th>
<th>Strains with positive phospholipase</th>
<th>Pz 15 days</th>
<th>Pz 20 days</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>M. furfur</em></td>
<td>5</td>
<td>5</td>
<td>1</td>
<td>1</td>
<td>0.68</td>
</tr>
<tr>
<td><em>M. pachydermatis</em></td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0.65</td>
<td>0.53</td>
</tr>
<tr>
<td><em>M. obtusa</em></td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td><em>M. slooffiae</em></td>
<td>4</td>
<td>3*</td>
<td>0</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. globosa</em></td>
<td>17</td>
<td>8*</td>
<td>0</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. restricta</em></td>
<td>1</td>
<td>0</td>
<td>0</td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>M. sympodialis</em></td>
<td>31</td>
<td>31</td>
<td>29</td>
<td>0.68</td>
<td>0.62</td>
</tr>
</tbody>
</table>

Pz: ratio of colony diameter to total diameter of the colonies plus the precipitation zone.

* Weak development.

The phospholipasic activity determined by the Price method is a simple technique applicable to a large number of strains. However, the measurement of the precipitation zone was not easy; in fact, the colony was already visible after 48 hours of incubation, while the precipitation zone around the colony was not observed until 8-10 days later, appearing extremely dense but clearly distinguishable from the colony. Up until day 15 of incubation both the diameter of the colony and that of the precipitation zone increased, whereas from day 15 to day 20 only the precipitation zone increased. Measurements of the precipitation zone taken at the day 15 and at day 20 from incubation were therefore deemed significant.

The ability of the species belonging to the Malassezia genus to produce phospholipase is shown in Table 2.

* *M. furfur* grew well on this medium but only for one strain it was possible to distinguish a late (day 20) precipitation zone around the colony. *M. pachydermatis* grew well and showed phospholipasic activity. *M. sympodialis* grew well and showed phospholipasic activity with the exception of two strains, one isolated from healthy subjects, the other supplied by IP, ex human normal skin.

*M. obtusa*, *M. slooffiae*, *M. globosa*, *M. restricta*, showed poor growth and did not produce a precipitate.

The greatest phospholipasic activity (Pz value) was shown by the strain *M. pachydermatis* after 20 days of incubation.

For *M. sympodialis* we compared the phospholipasic activity expressed as a Pz value mean with the dermatological disease presented by the patient.

Table 3 shows the Pz mean value of the strains isolated from patients with PV or SD, from individuals without any dermatological disease and from the reference strains of IP and CBS.

The Student t test highlighted significant differences between the Pz values of strains isolated from patients with PV and those of strains isolated from healthy individuals, both for readings on day 15 and day 20 subsequent to incubation (P = 0.04 and 0.05 respectively), while such difference was not significant for strains isolated from patients with SD (P = 0.08 and 0.4). The CBS and IP reference strains were isolated from individuals without any dermatological disease and showed a mean Pz value the same as that of the healthy individuals included in our research. Table 3 shows the phospholipasic activity of *M. sympodialis* strains isolated from patients with PV or SD, from sites with and without evident skin lesions.

After 15 days of incubation, the strains isolated from disease (PV and SD patients), both from healthy skin and lesioned skin, had the same Pz value, while, after 20 days, the strains isolated from the lesioned skin of patients with SD showed higher phospholipasic activity than the strains isolated from healthy skin (but not significantly P = 0.1) (Table 4).

Table 4: Phospholipasic activity of *M. sympodialis* with reference to strains origin.

<table>
<thead>
<tr>
<th>Strains origin</th>
<th>Pz 15 days</th>
<th>Pz 20 days</th>
</tr>
</thead>
<tbody>
<tr>
<td>PV patients lesioned skin</td>
<td>0.63</td>
<td>0.58</td>
</tr>
<tr>
<td>PV patients healthy skin</td>
<td>0.64</td>
<td>0.59</td>
</tr>
<tr>
<td>SD patients lesioned skin</td>
<td>0.69</td>
<td>0.60*</td>
</tr>
<tr>
<td>SD patients healthy skin</td>
<td>0.69</td>
<td>0.68</td>
</tr>
</tbody>
</table>

Pz: ratio of colony diameter to total diameter of the colonies plus the precipitation zone.

* Not statistically significant difference (P = 1).

Discussion

Extracellular phospholipases are considered virulence factors for many pathogenic bacteria and protozoa such as *Clostridium* species, *Listeria monocytogenes*, *Pseudomonas* species, *Staphylococcus aureus*, *Mycobacterium tuberculosis*, *Toxoplasma gondii* and *Entamoeba histolytica* and for some fungi such as *Candida albicans*, *Aspergillus fumigatus* or *Cryptococcus neoformans*.

Barret-Bee et al. in particular, were the first to correlate the production of phospholipase by *C. albicans* with its pathogenic nature, demonstrating that isolates with a high pathogenic potential (high level of adhesion to oral epithelial cells and greater pathogenicity for the mouse) had higher phospholipasic activity than yeasts with a low pathogenic potential. In addition, *C. albicans* blood isolates have shown greater in vitro phospholipasic activity than oral isolates from healthy patients. Similarly, it has been shown that isolates of *C. neoformans* taken from infections in AIDS patients show a higher level of phospholipasic activity than isolates taken from non AIDS patients or from bird droppings.

For the Malassezia genus some authors have observed the production of different extracellular enzymes such as lipase and phospholipase.

Using the Price method, extra-cellular phospholipasic activity was shown in vitro on strains of *M. furfur* isolated from human skin lesions. In addition, higher phospholipasic activity was observed in isolates of *M. pachydermatis* from skin lesions in dogs than in those from healthy skin.

The method used in this study to highlight the production of phospholipase by yeasts belonging to the Malassezia genus proved simple to apply and produced valid results for the species *M. furfur*.
M. pachydermatis and M. sympodialis, while the strains belonging to the species M. obtusa, M. slooffiae, M. globosa, M. restricta, had difficulty developing in the chosen medium so that it was not possible to measure phospholipasic activity. To measure the phospholipasic activity of Malassezia, ten days at least of incubation at 32 °C were needed, unlike the two or six days needed for C. albicans and C. neoformans, respectively.26,34 These results confirm those of other authors7 and indicate that the yeasts of the Malassezia genus produce phospholipase more slowly than C. albicans and C. neoformans.

For M. sympodialis it was possible to compare the production of phospholipase (Pz value) with the origin of the strains. Isolates of M. sympodialis taken from patients with PV showed significantly higher phospholipasic activity (P = 0.04) than those isolated from healthy individuals. Strains isolated from patients with SD and from healthy individuals did not show significant differences (P = 0.08) in the production of phospholipase. Strains isolated from the same patient, from healthy skin and from lesions, had the same Pz on average, both in the case of pityriasis versicolor and of seborrhoeic dermatitis. It is probable that the same strain colonises the entire individual but producing lesions is only able in some parts of the body, probably in relation to specific local conditions connected with the production of cutaneous lipids and/or alterations of the skin “ecosystem”.

These results are only partially comparable with previous works since only a limited amount of data in the literature refers to new species of Malassezia.

However, because egg-yolk contains substrates for both phospholipases (phospholipids) and lipases (triglycerides), the egg-yolk-based assay is not entirely specific (although the Pz value correlates with hydrolysis of phosphatidylcholine26) and should be considered as an initial screening which requires further confirmation using more specific enquiry such as radiometric or colorimetric methods.

This study suggests however that the phospholipasic activity of Malassezia may play a role in the onset of skin lesions, especially in the case of PV, even though phospholipases should be considered as only one of the many factors involved in the complex interaction between the yeast and its host leading to the development of skin lesions. More in-depth studies will be needed to understand the pathogenic role played by the enzymes secreted by Malassezia spp.

Conflict of interest

The authors report no conflict of interest.

References