Research Article

Eucalyptus Tree: A Potential Source of Cryptococcus neoformans in Egyptian Environment

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In Egypt, the River Red Gum (Eucalyptus camaldulensis) is a well-known tree and is highly appreciated by the rural and urban dwellers. The role of Eucalyptus trees in the ecology of Cryptococcus neoformans is documented worldwide. The aim of this survey was to show the prevalence of Cryptococcus neoformans during the flowering season of Eucalyptus at the Delta region in Egypt. Three hundred and eleven samples out of two hundred Eucalyptus trees, including leaves, flowers, and woody trunks, were collected from four governorates in the Delta region. Thirteen isolates of Cryptococcus neoformans were recovered from Eucalyptus tree samples (4.2%). Molecular identification of Cryptococcus neoformans was done by capsular gene specific primer CAP64 and serotype identification was done depending on LAC1 gene. This study represents an update on the ecology of Cryptococcus neoformans associated with Eucalyptus tree in Egyptian environment.

1. Introduction

The basidiomycetes yeast of genus Cryptococcus includes C. neoformans/C. gattii species complex, which is composed of two separate species, C. neoformans and C. deneoformans, and five species within C. gattii. The most clinically relevant complex species were recently revised based on phenotypic and genotypic diversity, supported by the presence of distinct and consistent lines, and it proposes to recognize C. neoformans var. grubii (represented by genotypes VNI and VNII) and C. neoformans (VNIII and VIV) as separate species, as well as five species of C. gattii (represented by genotypes VGI, VGII, VGIII, and VGIV) [1, 2].

C. neoformans has a worldwide distribution and has been recovered from pigeon droppings (Columba livia), urban environments, and soil. Many reports have shown the presence of C. neoformans in the hollows of different tree species, proposing that trees play a major role in Cryptococcus infection [3, 4]. The most common isolate responsible for this fungal infection is C. neoformans var. grubii serotype A [1, 5–7]. C. gattii has been proposed to have a specific ecological association with a number of Eucalyptus species [8].

E. camaldulensis is a well-known tree in Egypt since it was imported by Mohamed Ali, the Governor of Egypt (1805–1848), for fixing the River Nile banks in the 19th century. It is one of the most widely distributed trees in most of the arid and semiarid areas. This kind of tree exists on almost every roadside in Egypt, but there are no data about its role as potential carrier of fungal elements.

In Egypt, the incidence of Cryptococcus neoformans from Eucalyptus trees and pigeon droppings has been reported [9]. In this report, the author depended on the conventional differentiation methods to determine C. neoformans varieties. There is no recent noticeable information about the Eucalyptus tree role in the ecology of Cryptococcus neoformans in Egyptian environment. Therefore, the present study was aimed at determining the possible role of this tree as a potential dispersing source of Cryptococcus neoformans in Delta region’s environment.
Table 1: Eucalyptus tree collected samples in the Delta region.

| Governorate   | Tree number | Leaves | Flowers | Wood trunks | Total |
|---------------|-------------|--------|---------|-------------|-------|
| Giza          | 50          | 40     | 25      | 30          | 95    |
| Cairo         | 50          | 31     | 13      | 15          | 59    |
| Al-Sharqia    | 50          | 50     | 30      | 30          | 110   |
| Elmenofia     | 50          | 17     | 20      | 10          | 47    |
| Total         | 200         | 138    | 88      | 85          | 311   |

2. Materials and Methods

2.1. Study Area and Sampling Collection. The study area in Delta region covered 240 kilometers (150 mi) of Mediterranean coastline of Egypt. A total 311 samples out of 200 Eucalyptus trees, including leaves, flowers, and woody trunks, were collected from four different governorates in the Delta region (Cairo, Giza, Elmenofia, Al-Sharqia) by the rate of fifty samples for each region (Table 1). The samples were stored on ice in clean, sterile plastic bags and transferred to the Microbiology Department laboratory, Faculty of Veterinary Medicine. The samples were rinsed in sterile distilled water, then immersed in sterile saline solutions supplemented with chloramphenicol (10.0 mg/mL), and homogenized with ultrahomogenization for 4 min. The bottle was left for 30 min at room temperature to settle the sediment.

2.2. Isolation and Identification. From the supernatant fluid of each homogenized sample, a loopful was streaked onto plates of Sabouraud dextrose agar with chloramphenicol and incubated at 30°C for 48 hours. The colonies suspected to be C. neoformans were streaked on Eucalyptus leaves agar media [12]. The isolates were identified by classical mycological procedures of C. neoformans [13].

2.3. Molecular Characterization

2.3.1. DNA Extraction. The yeast cells from SDA slants were collected after 48-hour incubation with sterile PBS. The collected pellets were mixed into a microtube with 500 μL TES (100 mM Tris, pH 8.0, 10 mM EDTA, and 2% SDS); 50–100 μg Proteinase K from an appropriate stock solution was added and then incubated for 30 min (minimum) up to 1 h at 55°–60°C with occasional gentle mixing. The lysate mix salt concentration was adjusted to 1.4M with 5M NaCl (=140 μL); 1/10 vol (=65 μL) of 10% CTAB was added and incubated for 30 min at 65°C. The lysate was mixed gently and then incubated for 30 min at 0°C; finally, the mix was centrifuged for 10 min at 4°C, at 15000 rpm. The supernatant was transferred to a 1.5 mL tube followed by the addition of 0.55 vol isopropanol (=510 μL) to precipitate DNA followed by immediate centrifugation for 5 min, at 15000 rpm.

2.4. Molecular Identification by Capsular Gene. Detection of C. neoformans was done by using specific capsular gene primers CAP64. The primers for CAP64 were designed on the basis of DNA sequences (Table 2) [10].

2.5. Molecular Differentiation of Serotypes. This was applied by subjecting genomic DNA of identified strains by CAP64 gene to multiplex PCR amplification using a set of four primers of the laccase gene (LAC1) (Table 2) which were used for differentiating four major serotypes, A, D, B, and C, of C. neoformans [11].

3. Results

3.1. Recovery Rate of C. neoformans from Eucalyptus camaldulensis. In this study, E. camaldulensis trees acted as a potential refuge for C. neoformans. A total of 13 (4.2%) C. neoformans isolates out of 311 examined samples in Delta region were recovered during the flowering season of Eucalyptus tree. All the recovered Cryptococcus isolates were identified as C. neoformans strains based on all conventional and physiological characters of C. neoformans (Figure 1). Among these, 7 isolates (7.9%) were recovered from 88 Eucalyptus flowers, 5 isolates (3.6%) were recovered from 138 Eucalyptus tree leaves, and 1 isolate (1.1) was recovered from 85 woody trunks (Table 3).

3.2. Molecular Identification and Differentiation of C. neoformans. All tested isolates and reference strain were produced (400 bp) by CAP64 specific capsular gene primer (Figure 2). Molecular typing of C. neoformans isolates was done by four primers for LAC1 gene (Table 2) which were used for amplification; serotype A strains produced three DNA fragments with sizes of 0.88, 0.76, and 0.25 kb (Figure 3). All tested C. neoformans strains were identified as C. neoformans var. grubii and there are no other serotypes of C. neoformans detected.

4. Discussion

In Egypt, Eucalyptus trees are in abundance mainly as windbreaks and for afforestation of the drains and canals or other
Table 2: Primers used in this study.

| Primer | Primer sequence 3′-5′ | PCR product | Reference |
|--------|-----------------------|-------------|-----------|
| CAP64  | GCCAAGGGAGTCTTATATGG  | 400 bp      | [10]      |
|        | GCAAAGGTTCCACCAAATCG  |             |           |
| LAC1   | GGAACAGCAACCACACTACTG | 250 bp      | [11]      |
|        | CATATTGGGTGGCATCTTACTGAGGA | 760 bp |           |
|        | CCAGGGCAATGTGTGGAC     | 250 bp      |           |
|        | GTTGTGGAAGGCAAGAACAC   | 880 bp      |           |

Table 3: Recovery rate and distribution of C. neoformans in tested Eucalyptus trees sampled in the Delta region.

|          | Total samples | Giza | Cairo | Al-Sharqia | Elmenofia | Total | Recovery rate |
|----------|---------------|------|-------|------------|-----------|-------|---------------|
| Leaves   | 138           | 1    | 1     | 2          | 1         | 5     | 3.6           |
| Flowers  | 88            | 1    | 1     | 3          | 2         | 7     | 7.9           |
| Woody trunks | 85     | 1    | 0     | 0          | 0         | 1     | 1.1           |
| Total    | 311           | 3    | 2     | 5          | 3         | 13    | 4.2           |

Figure 2: Agarose gel electrophoresis of CAP64 gene specific PCR of all the examined C. neoformans isolates with production of amplicons of 400 bp, marker 100 bp DNA ladder (Jena Bioscience).

Figure 3: Agarose gel electrophoresis of LAC1 gene specific PCR of all the examined C. neoformans isolates with production of amplicons of 250, 760, and 880 bp, marker 100 bp DNA plus ladder (Jena Bioscience).

determine C. neoformans variety, which evoked a high need to investigate the environmental ecology of this fungus, depending on molecular techniques to determine the actual variety of C. neoformans in relation to E. camaldulensis in order to establish a real surveillance program and applying the preventive measures for this pathogen infection.

This study was applied on Eucalyptus trees during the flowering season, as most C. neoformans and C. gattii reported cases were associated with Eucalyptus showing strong seasonality in its occurrence, which coincides with the periods of flowering [17].

The results show that the isolation of C. neoformans from Eucalyptus flowers is more frequent than from leaves and woody trunk (Table 3). All examined isolates were identified as C. neoformans var. grubii with a recovery rate of 4.2% of the total examined samples. It is normally the high isolation rate of var. grubii as the global distributed isolate responsible for cryptococcal infection [1, 5–7]. Also, it is commonly the recovering of C. neoformans from pigeon droppings, soil, and decaying wood in hollow trees [3].

Ambitiously, the present study documents the first record for the isolation of var. grubii from E. camaldulensis leaves, flower, and woody trunks in Egypt. Most of the previous reports stated that C. grubii association with Eucalyptus trees or other types of trees is interpreted in one sentence: “C. neoformans presence might represent fecal contamination by birds inhabiting these trees” [9, 14, 16].

Globally, many reports are recorded for isolation of C. grubii from Eucalyptus tree parts or other types of trees. In India, more than one report states that C. grubii tree association and its distribution differ from each part of tree or season or time of the study. The prevalence of C. grubii (5.56%) and C. gattii (9.26%) from decayed wood inside trunk hollows of diverse tree species was reported [18].

Recently, C. grubii was isolated from the bark of Eucalyptus trees followed by flower, bud, fruit, and detritus [19]. The prevalence of C. grubii in this study (4.2%) is somewhat near to the rate of Nawange et al.’s (2006) [18] study (5.5%), while the recovery rate of C. grubii is the highest from flowers.
(7.9%), then leaves (3.6%), and finally woody trunks (1.1%) (Table 3).

In sunny countries, *C. neoformans* can escape from lethal effects of sunlight and drying by taking trees as a good natural habitat in the environment because these pathogens can live inside woody debris as well as trunk hollows. The result of the present study highlighted the potential role of tree parts of *E. camaldulensis* in environmental ecology of *C. grubii*. *Eucalyptus* flowers were the best natural habitat and a suitable transporting means for these pathogens infectious propagules in the surrounding environment. Flowering season of *Eucalyptus* tree is mainly in winter and spring from November to February. At this time of year in Egypt, the temperature is slightly low to temperate which gives a potential chance for isolation of this pathogen. The association between *C. grubii* and tree is controlled by many environmental factors including humidity, temperature, and solar radiation [13, 17].

In Egypt, the *Eucalyptus* tree exists almost along every roadside, especially in the Delta region around River Nile and its tributaries. These study results confirm the potential role of *Eucalyptus* trees as a major source for *C. grubii* in Egyptian environment which act as a high risk for immunocompromised patients.

Most of the reported cases of human cryptococcosis were registered as cryptococcal meningitis. Cryptococcal meningitis in Egypt is rarely diagnosed, but this may be due to inadequate investigation rather than absence of definite epidemiological data about the organism in Egypt. More attention should be considered for human cases of unexplained chronic meningitis that is not responding to conventional therapy as *C. neoformans* could be the main cause of such fatal meningitis [20–23].

The only survey for fungal meningitis was done in Egypt at NAMRU-3 during the period of 1998 to 2001 of 1000 CSF samples, where 10 *C. neoformans* were recovered at a rate of 0.01%. All isolates belonged to serotype A (*C. neoformans var. grubii*) [24]. Recently, *C. neoformans* serotype A is the most common variety in association of pet birds droppings in the Egyptian environment [25].

This study's findings come in the same direction with the previous surveillance of the main causes of cryptococcal infection in Egypt and it confirmed that *C. neoformans var. grubii* is the main etiological agent of cryptococcal infection in Egypt.

Conclusively, this is the first record describing isolation of *C. neoformans var. grubii* from *E. camaldulensis* in Africa and Egypt. The results highlighted the potential role and risk of *Eucalyptus* tree as a carrier reservoir of one of the high pathogenic fungal elements in Egypt.

**Conflict of Interests**

There is no conflict of interests.

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