Aeromycological study at the intensive care unit of the “Dr. Manuel Gea Gonzalez” General Hospital

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A b s t r a c t
Introduction: An aeromycological study verifies the presence and quantifies the concentration of fungal propagules in the air. It is very important in the hospital setting because of the increasing numbers of immunosuppressed and severely ill patients. The objective of this study was to determine the concentration of fungi in the air of the intensive care unit (ICU) of “Dr. Manuel Gea González” General Hospital.

Methods: This is a descriptive, observational cross-sectional study. Air samples were obtained with a single stage Thermo-Andersen Viable Particle Sampler (Thermo Electron Corporation - Massachusetts, U.S.A.) in a Petri dish with potato dextrose agar for 15 minutes at two different times (morning and afternoon) and heights (1 and 1.5 meters). The Petri dishes were incubated for five to seven days at 27°C, the number of colonies was counted, and the total CFU/m³ was determined. The isolated fungal genera were identified by morphological features. Epi Info v. 3.4.3 © was used for statistical analysis.

Results: The mean concentration of fungi in the air of the ICU was 85.08 ± 29.19 CFU/m³; while in the outside air it was 84.3 ± 17.23 CFU/m³ (p = 0.96). The fungi isolated were: Cladosporium spp., Penicillium spp., Aspergillus spp. (non-fumigatus), Fusarium spp., Exophiala spp., Syncphalastrum spp., and Acremonium spp.

Discussion: Fungal spores were found in the air of the ICU and Cladosporium spp. was the most frequently isolated fungi. There was no difference according to sampling time or height.

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I n t r o d u c t i o n

Bioaerosols are aerial suspensions of particles from live organisms, microorganisms or other biological materials.1-3 Aeromycology is the branch of aerobiology that studies the dispersion of spores and other fungal elements in indoor and outdoor air, the changes in their concentrations, and the factors that affect those changes.3 Fungal spores enter hospitals...
through ventilation systems and fungi develop on multiple surfaces, releasing more spores.\textsuperscript{1,3} The number of spores in indoor air varies depending on climate, weather, air currents, humidity, temperature, time of the day, type and maintenance of ventilation systems, age of the buildings, movement of people, cleaning, and the presence of plants or food.\textsuperscript{1,3-7}

Immunosuppressed patients with severe neutropenia, chronic granulomatous disease, and acquired immunodeficiency syndrome (AIDS) have the highest risk of developing invasive fungal infection.\textsuperscript{8-10} The main fungal genera related with these diseases are \textit{Aspergillus}, \textit{Candida}, \textit{Fusarium}, \textit{Penicillium}, \textit{Mucor}, and \textit{Rhizopus}.\textsuperscript{11-17} Invasive aspergillosis (especially by \textit{A. fumigatus}), candidemia, disseminated fusariosis, infections by \textit{P. marneffei}, and zygomycosis have a mortality rate that can reach 100\%.\textsuperscript{10-17}

The single stage Thermo-Andersen Viable Particle Sampler (Thermo Electron Corporation – Massachusetts, USA) is the most frequently used air sampler.\textsuperscript{8,18,19} In the detection phase, the amount of colony forming units (CFU) per cubic meter of air is determined, and then the identification of the fungi is performed by culture or by molecular techniques.\textsuperscript{7,9,18-21}

Because the number of immunosuppressed and severely ill patients is increasing worldwide, especially in the hospital setting, this study was designed to determine the concentration of fungi in the air of the intensive care unit (ICU) of this hospital.

\section*{Materials and methods}

This was a descriptive, observational, cross-sectional study. The universe was the air of the ICU of “Dr. Manuel Gea Gonzalez” General Hospital. The ICU is divided in different sections with different air volumes: five single rooms (30 m\textsuperscript{3} each), one double room (120 m\textsuperscript{3}), one aisle (67.5 m\textsuperscript{3}), and one nurse module (37.5 m\textsuperscript{3}). For convenience, an air sample was taken from the area just outside the ICU (one meter away from the entrance door), from one single room (picked randomly), from the double room, from the aisle, and from the nurse module. The total air volume of the areas that were sampled inside the ICU was 255 m\textsuperscript{3}. The air volume sampled from the five selected areas (inside and outside the ICU) was 0.1415 m\textsuperscript{3}.\textsuperscript{14} The unit is ventilated by a central air conditioning unit without fans or open windows. The unit can accommodate up to seven patients and always has at least ten staff members working. The highest level of activity occurs during morning hours. Among the studied variables are: humidity, temperature, CFU per air volume, and fungal agents.

Air samples were obtained with a single stage Thermo-Andersen Viable Particle Sampler (Thermo Electron Corporation – Massachusetts, USA) in a Petri dish with potato dextrose agar. The sampler was placed in the center of each room, and each sample was collected with a vacuum flow of 28.3 liters/min for 15 minutes.\textsuperscript{14,22} In each sampling area, four samples were collected: one at each time period (morning and afternoon) and one at each height (one and 1.5 meters). Humidity and temperature were measured with a hygrothermograph (Control Company – Friendswood, Texas, USA). Nobody was allowed in or out of the room during sampling.

After sampling, the Petri dishes were incubated for five to seven days at 27\textdegree C.\textsuperscript{1,14} After that, the number of colonies was counted in every Petri dish. The total number of fungal colonies present in the ICU was determined by obtaining the mean of the colony count in the four areas sampled. This count was stratified by time and height of the sample. The concentration of fungal propagules in the air was expressed in CFU/m\textsuperscript{3}. For its calculation, a correction factor was used for the colony count, based on the probability that more than one viable propagule could have passed through the same hole and impacted the culture medium. The formula used was:

\[ \text{CFU} = N \ln[N/N-P] \]

Where CFU was the corrected colony count, \( N \) was the number of holes in the perforated plate of the sampler (400), and \( P \) was the number of colonies that grew in the medium.

The corrected count represents the real number of CFUs present in the sampled air. After that, the CFU/m\textsuperscript{3} of air was determined with the following formula:

\[ \text{CFU/m}^3 = \frac{\text{CFU}}{t} \times K \]

Where CFU was the corrected colony count, \( t \) was the total sampling time expressed in minutes, and \( K \) was a conversion factor from cubic feet to cubic meters (\( K = 35 \)).

The identification of fungi was performed by observation of the macroscopic characteristics of the colonies and the microscopic characteristics of the sporulating hyphae.\textsuperscript{5}

The statistical analysis was performed with the Epi Info v. 3.4.3© software, and the means and standard deviations of humidity, temperature, and colony forming units per air volume were determined according to sampling time and height. Independent Student’s \( t \)-test was used to determine whether there was a statistically significant difference between the means of CFU per air volume according to sampling area, time, and height. Differences were considered significant when the \( p \)-value was below 0.05.

All procedures performed were in accordance with and approved by the ethical standards of the “Dr. Manuel Gea Gonzalez” General Hospital Review Board and Ethics Committee. The principles of the Helsinki Declaration of 1975 with the modifications of 1983, and the Mexican General Health Law were followed.

\section*{Results}

There was no difference in the fungal spore concentration of the air inside the ICU when compared with the air outside the unit (Table 1). These concentrations were not affected by the time of sampling (\( p = 0.13 \) and \( p = 0.14 \), respectively) or the height of sampling (\( p = 0.99 \) and \( p = 0.92 \), respectively) (Table 1). Several colonies were cultured, and the fungal genera isolated were: \textit{Cladosporium} spp., \textit{Penicillium} spp., \textit{Aspergillus} spp. (non-\textit{fumigatus}), \textit{Fusarium} spp., \textit{Exophiala} spp., \textit{Syncephalastrum} spp. and \textit{Acremonium} spp. (Table 2). The mean humidity during sampling was 38.88 ± 3.20% and the mean temperature was 24.09 ± 0.82\textdegree C.
However, some of these fungal genera also represent a threat to immunocompetent hosts, such as the healthcare staff of the ICU. Fungal spores of several genera, such as Cladosporium spp., have been associated with several diseases such as rhinosinusitis, chronic cavitary pulmonary aspergillosis, aspergillosoma, allergic bronchopulmonary aspergillosis, and skin and wound infections, among others.10,12,13 However, some of these fungal genera also represent a threat to immunocompetent hosts, such as the healthcare staff of the ICU. Fungal spores of several genera, such as Cladosporium spp., have been associated with several diseases such as rhinosinusitis, chronic cavitary pulmonary aspergillosis, aspergillosoma, allergic bronchopulmonary aspergillosis, and skin and wound infections, among others.10,12,13

The present results underline the importance of establishing control measures to improve air quality inside the ICU, aiming to reduce fungal-related morbidity and mortality. Among these measures, the following can be mentioned: periodic measurements of fungal propagules in the air with aeromycological studies, rigorous cleaning with disinfectants and dust removal, routine equipment maintenance, humidity control, air filtration with high efficiency particle air filters, and use of laminar flow ventilation systems, among others.3,4,24

**Conclusion**

Fungal spores were found in the air of the ICU (85.08 ± 29.19 CFU/m³), and Cladosporium spp. was the most isolated fungi. There was no difference according to sampling time or height.

**Conflict of interest**

All authors declare to have no conflict of interest.

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REFERENCES


