

# بسم الله الرحمن الرحيم





شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم



شبكة المعلومات الجامعية

# جامعة عين شمس

التوثيق الالكتروني والميكروفيلم

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# بالرسالة صفحات لم ترد بالاصل

### Study of Fungal Infections in **Pediatric Oncology Patients**

Thesis Submitted for partial fulfillment of Master Degree In Pediatrics

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المالة العزاديم.

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سورة البقرة آية رقم :(٣٢)

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## LIST OF ABBREVIATION

ALL Acute lymphoblastic leukemia

**AMC** Absolute monocytic count

**AML** Acute myeloid leukemia

ANC Absolute neutrophil count

BAL Bronchoalveolar lavage

CD Cluster differentiation

CT Computed tomography

DNA Deoxyribose nucleic acid

**EBV** Epstein-Barr virus

ECG. Electrocardiogram

e.g. Exempli gratia (Latin= for example)

**ESR** Erythrocyte sedimentation rate

**FAB** French-American-British

FDA Food and drug Administration

FN Febrile neutropenia

**FUO** Fever of unknown origin

G-CSF Granulocyte-colony stimulating factor

**GM-CSF** Granulocyte-macrophage colony stimulating factor HD

Hodgkin disease

HIV Human immunodeficiency virus

**IgM** Immunoglobulin M

I.V. Intravenous

**NAC** Non-albicans Candida

NHL Non-Hodgkin's Lymphoma

PCR Polymerase chain reaction

**PMN** Polymorphonuclear neutrophil

**SNCCL** Small non-cleaved cell lymphoma

T.d.T. Terminal deoxynucleotide transferase

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#### INTRODUCTION

Infections are common and potentialy serious complications of cancer treatment. The most important risk factor for infection is an absolute neutrophil count less than 500 / mm<sup>3</sup>, and its duration (Oude Nijhuis et al., 2005).

In the early 1970s, the introduction of empirical use of intravenous broad-spectrum antibiotics for febrile neutropenic patients reduced infection-related mortality significantly. Until recently, routine hospitalization for administration of intravenous broad-spectrum antibiotics is still standard care and the paradigm in most hospitals (Oude Nijhuis et al., 2005).

Oncology patients overall had a decreased mortality rate over time, likely attributable to advances in medical therapy in treating a variety of neoplasms. However, patients who acquired fungal infections did not experience a similar improvement in mortality. Although fungal infections themselves were not usually fatal events, they were associated with a 60% mortality rate in cancer patients (*Rosen et al.*, 2005).

El-Mahallawy et al., (2002) studied fungal infections in children with cancer at the National Cancer Institute in Cairo. They found that 68% of infections were caused by Candida sp. and 18% by Aspergillus sp.

Fungal infections pose a serious risk to immunocompromised patients. Over the past decade, a significant rise in the number of new infections has been observed in this population (Groll et al., 1996). Although Candida remains the most common fungal pathogen in

oncology patients, the epidemiology has recently shifted toward non-albicans species (Moosa et al., 2002).

This change has been attributed to the widespread use of fluconazole prophylaxis *Bodey et al.*, (2002); however, Aspergillus and uncommon molds have also seen a recent rise as pathogens identified in immunocompromised individuals (*Perea et al.*, 2002).

The availability of newer therapeutic agents with improved efficacy, including immunomodulators, will be crucial in obtaining better control of infections due to fungal organisms (Anaissie et al., 1998; Boogaerts et al., 2001).