NOSOCOMIAL INFECTIONS IN HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTED AND AIDS PATIENTS: MAJOR MICROORGANISMS AND IMMUNOLOGICAL PROFILE

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ABSTRACT

Antiretroviral therapy advances have proportioned to AIDS patients a survival increase. At the same time, the permanence of the seropositive people in the nosocomial environment becomes common not only by the adverse reactions caused by this therapy, but also by several opportunistic diseases that take them into and out of hospital environment. During the hospital permanence, the patients expose their impaired immune system to the nosocomial virulent microorganisms, and acquire destructive nosocomial infections that sometimes can be lethal. Among several hospital syndromes described, little is known about infections in immunocompromised patients and how their immune system is able to determine the course of the infection. The objective of this study was to describe the major microorganisms involved in the nosocomial infections of HIV-1 seropositive patients associated with their immunological status. The survey was carried out with the Hospital Infection Control Service records, from University Hospital, Londrina, Paraná, Southern of Brazil, during the period from July 2003 to July 2004. From all the cases studied (n=969), 24 patients (2.5%) had AIDS diagnosis and a half of them was women with the mean of CD4+ T cells counts of 158/mm3. The main topography of the infection was pulmonary (50.0%) and the main isolated microorganisms were Staphylococcus aureus, Pseudomonas aeruginosa and Escherichia coli. A major incidence of infection was observed in patients with CD4+ T cells counts lower than 50/mm3. The study of the relationship between the impairment of the immune system and infectious agents could provide a better healthcare of people living with HIV/AIDS and advances into the nosocomial infection control systems.

Key-words: AIDS, CD4+ T cell counts, nosocomial infections, immunocompromised

INTRODUCTION

Nosocomial infections (NIs) are resulted from treatment in a hospital or hospital-like setting, but secondary to the patient’s primary condition and constitute a problem of world-wide public health, that promote greater nosocomial permanency and an increase in the developing countries rates of mortality and morbidity (4,8).

NIs are even more alarming in the 21st century as antibiotic resistance spreads. Reasons why these infections are common include hospitals house large number of people who are sick and whose immune system are often in a weakened state; medical staff move from patient to patient provoking a way for pathogens spreading; many medical procedures bypass the body’s natural protective barriers and the routine use of antimicrobial agents in hospital creates selection pressure for the emergence of resistant strains (8).

The most common NIs are of the urinary tract and various pneumonias (6,8,32) and few studies have been reported about their occurrence in immunocompromised individuals.

Currently, the immunossupression is a state that is very common because the advances of science to increase survival...
in the extremities of age and the increase of the incidence of diseases that impair the immune system, such as neoplasia and the Acquired Immunodeficiency Syndrome (AIDS) (3,23). AIDS is the advanced clinical phase of the infection caused by the human immunodeficiency virus types 1 (HIV-1) and 2 (HIV-2) (27). The progression of HIV-1 infection causes immunological dysfunctions that contribute either to the appearance of opportunist diseases or to the reactivation of latent infections.

The predominant immunological characteristic in the course of HIV-1 infection is the progressive depletion of CD4+ T helper type 1 subpopulation of lymphocytes (CD4+ Th1) that changes the normal rate between CD4+T/CD8+T lymphocytes and promotes the immunodeficiency (DIAS et al., 2003). The CD4+ T cell count has important role to the introduction of high active antiretroviral therapy (HAART) (13).

Barlett & Gallant (1) revised the correlation between the appearance of complications of HIV-1 infection and CD4+ T cell counts from AIDS patients. However, the available data reported the acquired opportunist infections that occurred in the community, and little is known about the correlation between the number of these cells and the appearance of NIs and its causal pathogens. Although various studies described the different infections developed by HIV/AIDS patients during the nosocomial period and their etiological agents, the correlation between the pathogens and the compromising immunological status is still little explored.

The aim of this study was to describe the immunological profile and to characterize the etiological agents of NIs in HIV/AIDS patients, in order to contribute for improvements in the nosocomial infection control and cares in these patients.

MATERIALS AND METHODS

A retrospective study was carried out involving all patients admitted at the University Hospital from Londrina, Paraná, Southern of Brazil, whose were registered NIs during the period from July 2003 through July 2004. Patients personal data, CD4+ T cell counts and the isolated microorganisms results were collected from notified data of the active researches of the Nosocomial Infections Control Center (CCIH) from this health service. This study was approved by the Research Ethics Committee of the Londrina State University. The variables investigated were age, gender, topography, CD4+ T cell counts, clinical evolution and microorganisms isolated from these patients.

RESULTS

The total number of registered NIs cases in the studied period was 969 (Table 1). Of them, 257 (26.52%) occurred in patients admitted to the Newborn Intensive Therapy Unit, 158 (16.03%) were patients admitted to the Intensive Treatment Unit I and 157 (16.20%) were patients admitted to the Female Nursing Unit. The distribution of the cases, according to the gender, is presented in Table 2. Most of the cases occurred in male patients representing 591 cases (61.0%).

Of the total 969 patients, 24 (2.5%) had the AIDS diagnosis. The distribution of the NIs number among these patients according to the sex was of 50.0% and the age ranged from 0 to 78 years old, with the mean of 40.2 years old (Table 3).

The mean time of permanence in the hospital was of 36.16 days and 13 patients dead during this period. The mean CD4+ T cell count was 158.4 cells/mm³ (Table 3). In relation to infection sites, the main topography was the lung (50.0%), followed by infections in multiple sites (25.0%), urinary tract infections (8.3%) and bloodstream (8.3%), as demonstrate the Table 3 and the

| Nosocomial Unit                              | Case Number (n) | %     |
|----------------------------------------------|-----------------|-------|
| Male Nursing Unit                            | 257             | 26.50 |
| Intensive Treatment Unit I                   | 158             | 16.30 |
| Female Nursing Unit                          | 157             | 16.21 |
| Newborn Intensive Treatment Unit             | 81              | 8.36  |
| Emergency                                    | 78              | 8.05  |
| Pediatric Unit                               | 68              | 7.02  |
| Intensive Treatment Unit II                  | 45              | 4.65  |
| Infectious Diseases Unit                     | 42              | 4.34  |
| Pediatric Intensive Treatment Unit           | 24              | 2.48  |
| Intensive Care Unit                          | 22              | 2.28  |
| Maternity Unit                               | 20              | 2.07  |
| Dialysis Unit                                | 11              | 1.14  |
| Tisiology Unity                              | 5               | 0.5   |
| Pediatric Nursing Unit                       | 1               | 0.1   |
| TOTAL                                        | 969             | 100.0 |

| Gender           | Number (n) | % |
|------------------|------------|---|
| Male             | 591        | 61.0  |
| Female           | 276        | 28.5  |
| Newborn *        | 102        | 10.5  |
| TOTAL            | 969        | 100.0 |

* The gender was not registered in these cases.
Table 3. Distribution of nosocomial infections registered cases in HIV/AIDS patients attended at University Hospital, Londrina, Paraná, from July 2003 to July 2004, according to gender, age, nosocomial permanence, topography, isolated microorganism, CD4+ T cell counts and clinical evolution.

| Case | Gender | Age (years) | Nosocomial permanence (days) | Topography of infection | Isolated Microorganism | CD4+ T cell counts (cell/mm³) | Clinical evolution to death |
|------|--------|-------------|-----------------------------|-------------------------|------------------------|----------------------------|-----------------------------|
| 1    | F      | 26          | 145                         | Meningis                | ——                     | 15                         | No                          |
| 2    | F      | 27          | 21                          | Urinary tract           | E. coli                | 38                         | No                          |
| 3    | F      | 65          | 22                          | Lung                    | E. coli                | 25                         | Yes                         |
| 4    | M      | 40          | 06                          | Lung                    | ——                     | ——                         | Yes                         |
| 5    | F      | 36          | 36                          | Urinary tract           | E. coli                | 13                         | Yes                         |
| 6    | F      | 38          | 82                          | Urinary tract           | P. aeruginosa          | 203                        | Yes                         |
| 7    | F      | 27          | 19                          | Lung                    | ——                     | 216                        | No                          |
| 8    | M      | 60          | 17                          | Lung                    | ——                     | ——                         | No                          |
| 9    | M      | 58          | 17                          | Urinary tract           | P. aeruginosa          | 32                         | Yes                         |
| 10   | F      | 35          | 66                          | Lung                    | S. aureus              | 23                         | Yes                         |
| 11   | M      | 35          | 08                          | Lung                    | A. baumanii            | 313                        | Yes                         |
| 12   | M      | 31          | 59                          | Urinary tract           | E. faecalis            | 91                         | Yes                         |
|      |        |             | Paranasal sinuses           |                         |                        |                            |                             |
| 13   | M      | 54          | 84                          | Lung                    | S. aureus              | ——                         | ——                          |
| 14   | M      | 48          | 27                          | Urinary tract           | P. aeruginosa          | 416                        | No                          |
| 15   | F      | 26          | 6                           | Lung                    | ——                     | 308                        | No                          |
| 16   | F      | 18          | 45                          | Perineum                | ——                     | 336                        | No                          |
| 17   | M      | 62          | 36                          | Lung                    | ——                     | 113                        | No                          |
| 18   | M      | 44          | 39                          | Lung                    | P. aeruginosa          | 216                        | No                          |
| 19   | M      | 54          | 42                          | Urinary tract           | P. aeruginosa          | 1                          | Yes                         |
|      |        |             | Catheter                   |                         |                        |                            |                             |
| 20   | M      | 40          | 18                          | Bloodstream             | K. pneumoniae          | 802                        | Yes                         |
| 21   | F      | 28          | 21                          | Lung                    | ——                     | 300                        | Yes                         |
| 22   | M      | 34          | 11                          | Lung                    | ——                     | 1                          | Yes                         |
| 23   | F      | 32          | 31                          | Bloodstream             | ——                     | 19                         | Yes                         |
| 24   | F      | 47          | 10                          | Lung                    | ——                     | 3                          | No                          |
| n=24 | F=12   | Mean        | Mean                        | 8.3% urinary tract      | ——                     | 158.4±40.3*                | 50.4%                       |
|      | M=12   | 40.2        | 36.2                        | 50.0% of infections had an isolated microorganism | ——                     | (n = 22) dead |                             |
| 157  |        |             |                             | 8.3% bloodstream        | ——                     |                            |                             |

**Notes:**
- a oxacilina resistant strain (ORSA);
- b betalactamase producing strain;
- c carbapenase producing strain (CESP);
- d carbapenase producing strain and resistant to the polimixina;
- e extended spectrum betalactamase producing strain (ESBL);
- * average ± standard deviation; n = total considered; M = male; F = female; ——: no data available.
Fig. 1. Microorganisms were isolated from culture in 50.0% of the cases. Of them, 66.7% showed antimicrobial drug resistance. The major incidence of microorganisms in AIDS patients and clinical evolution to death was observed when the CD4+ T cell counts were lower than 50 cells/mm³ (Tables 4 and 5, respectively). However, the differences were not significant (Fisher’s Exact Test, p ≅ 1.000 and p = 0.687, respectively). The most frequent isolated microorganisms were *Staphylococcus aureus* (*S. aureus*) in 36.8% of cases, *Pseudomonas aeruginosa* (*P. aeruginosa*) in 26.3% of cases and *Escherichia coli* (*E. coli*) in 15.8% of cases (Fig. 2).

**Table 4.** Distribution of HIV/AIDS patients attended at University Hospital, Londrina, Paraná, Southern Brazil, from July 2003 to July 2004, according to the isolation of microorganisms and CD4+ T cell counts.

| CD4+ T cell counts (cells / mm³) | Patients with isolated microorganism n (%) | Patients without isolated microorganisms n (%) | Total n (%) |
|----------------------------------|------------------------------------------|---------------------------------------------|-------------|
| ≥500                             | 1 (8.3)                                  | 0 (0.0)                                     | 1 (4.5)     |
| < 500 and ≥ 50                   | 5 (41.7)                                 | 5 (50.0)                                    | 10 (45.5)   |
| < 50                             | 6 (50.0)                                 | 5 (50.0)                                    | 11 (50.0)   |
| Total                            | 12 (54.5)                                | 10 (45.5)                                   | 22 (100.0)  |

n: number of patients; %: percentage, p ≅ 1.000 (Fisher’s Exact Test)

**Table 5.** Occurrence of death in HIV/AIDS patients attended at University Hospital, Londrina, Paraná, Southern Brazil, from July 2003 to July 2004, according to the isolation of microorganisms and CD4+ T cell counts.

| CD4+ T cell counts (cells / mm³) | Death cases in microorganism presence n (%) | Death cases in microorganisms absence n (%) | Total n (%) |
|----------------------------------|---------------------------------------------|--------------------------------------------|-------------|
| ≥500                             | 1 (11.1)                                   | 0 (0.0)                                    | 1 (7.7)     |
| < 500 and ≥ 50                   | 2 (22.2)                                   | 2 (50.0)                                   | 4 (30.8)    |
| < 50                             | 6 (66.7)                                   | 2 (50.0)                                   | 8 (61.5)    |
| Total                            | 9 (69.2)                                   | 4 (30.8)                                   | 13 (100.0)  |

n: number of patients; %: percentage, p = 0.687 (Fisher’s Exact Test)

**DISCUSSION**

NIs are pointed as the major cause of morbidity and mortality in hospitalized patients. According to Center of Diseases Control, the NIs occurrence costs approximately US$ 2,100, varying from US$ 680 in bladder infections until US$ 5,683 in respiratory tract infections.

The main described risk factors for HIV/AIDS patients to acquire NIs, beyond the immunossupression state, are the length of stay in the hospital, insertion of urinary catheter, bloodstream access, multiple admissions to the hospital, number and duration of invasive procedures, use of prophylactic antimicrobials and malnutrition (3,25). Colonization seems to play an important role in NIs among these immunocompromised patients (16,25).

Amongst the NIs cases in HIV/AIDS patients evaluated in this study, a similar gender distribution was observed. Few reports in literature showed the distribution of these infections in AIDS patients considering parameters as gender, age or length of hospitalization. The gender was pointed as a risk factor for nosocomial Gram-positives infections in male (16), what can be observed in 57.1% of HIV-infected patients of this study.
Studied carried out about NIs in HIV/AIDS patients (9,24,26) reported the bloodstream as the main topography. In the present study, however, a low frequency of this site of infection was demonstrated (8.3%) and can be related to the low sensitivity of the hemoculture, the major diagnosis method for bacteremia. The highest frequency in our study was the pulmonary topography, which is related as an important cause of treatable morbidity among HIV-1 patients (8). The infections related to the respiratory tract are pointed as the major NI syndrome (7), independently of the original disease and immunological status of the patients and can be related with the high number of invasive procedures in this topography, frequently used as the HIV-1 infection progresses (1). In a case-control study, Tumbarello and collaborators verified that the CD4+ T cell counting < 100/mm³ was a risk factor to the community and nosocomial acquired pneumonia. In our data, the distribution of this site of infection was 46.2% of the total number of patients with CD4+ T cell counting lower than 100/mm³, underscoring the fact that the nosocomial acquired pneumonia can occurs at all CD4+ T cell counting (8).

All the HIV-1 infected patients enrolled in this study had AIDS diagnosis, the most advanced phase of the HIV-1 infection. The CD4+ T cell counts in these patients showed results that ranged from 1 to 802 cells/mm³, with average of 158.4 cells/mm³, and the majority of these patients showed values bellow of 400 cells/mm³, as are expected in these patients in the phase of opportunistic diseases or reactivation of latent infections (1,3,9,25).

A reduction is observed in the evolution of CD4+ T cell counts throughout the years after the AIDS diagnosis and characterizes the last disease clinical phase with values bellow 200 cells/mm³; in this phase occurs the development of opportunistic disease, the reactivation of latent infections and other organic complications, as well it is considered an indicator of mortality (1). Many clinical and laboratory measures have been used to assos disease progression and disease progression in HIV-1 infection. Previous studies showed that the single best predictor of progression to AIDS was the percentage or absolute number of circulating CD4+ T cells; further studies showed that plasma viral load was a better indicator of prognosis that CD4+ T cell. Currently, the combined measurement of plasma viral load and CD4+ T cell counts is useful to predict the disease progression (21).

The CD4+ T cell gradually declines over several years with a more accelerated decline 1.5-2 years before an AIDS-defining diagnosis. Late-stage of the disease is characterized by a CD4+ T cell count < 200 cells/mm³ and the development of opportunistic infections, selected tumors, wasting and neurologic complications (BARLETT et al., 2000). Thus, the results obtained in this study are in agreement with the literature. Prophylactic treatment for opportunistic infections is recommended when CD4+ T cell counting is lower than 200/mm³ (27).

The major microorganisms isolated were from the patients with CD4+ T cell <500/mm³ which confirm the importance of this laboratory marker in the prognosis of HIV-1 infection. Considering that NIs also constitute in an opportunist disease for these patients, it is possible that exist a relation between CD4+ T cell counts and the incidence of NIs.

Major incidence of culture isolated microorganisms was observed in those patients that CD4+ T cell counts were bellow 500 cells/mm³. The main microorganisms isolated were S. aureus, P. aeruginosa and E. coli (Table 4 and Fig. 2), in accordance with previous reports in the literature (15,16, 29).

S. aureus is described as a NI agent in HIV-1 patients (16) and is distinguished for its capacity in producing resistance mechanisms that difficult the antimicrobial drug therapy. The meticillin-resistant S. aureus (MRSA) appeared as a nosocomial agent at 1960’s in countries of the Europe and 1976 in U.S.A. In Brazil, the meticillin is not used. Thus, the cited microorganisms as MRSA are tested against the oxacilina and so called oxacillin-resistant (ORS), since both antibiotics are equivalents.

Considering the microorganism strains isolated in this study, 85.0% presented oxacilin resistance, indicating the selective pressure exerted by the drug antimicrobial therapies on these agents. Gram – positive microorganisms are described as the main isolated agents in AIDS patients and showed drug-resistance mechanisms (19,24). HIV-infected patients frequently receive prophylactic antimicrobial therapy that can be a possible cause of drug-resistance in the described cases.

The transmission of the MRSA occurs mainly through the contaminated hands of the health professionals, since these microorganisms are able to survive for hours in the environment. The surfaces of the environment also can be a source of transmission (15,16). The main risk factors for infection with MRSA include age, previous long time hospitalization, long hospital permanence, admission into intensive care units, chronic illness, previous antimicrobials drug therapy, manipulation for colonized professionals, and accomplishment of invasive procedures (15) that frequently occur in HIV-patients. MRSA NI in these patients is mainly related with the prolonged time of hospitalization, in agreement with the data showed in the present study (7).

Amongst the isolated Gram-negative microorganisms in this study, P. aeruginosa and E. coli presented the highest frequency. This result is in agreement with the previous reports which affirmed that P. aeruginosa is common in NIs and is pointed as an important agent of morbidity and mortality among HIV-1 patients because it can be fatal in the advanced stages of HIV-1 disease, specially when the CD4+ T cell counting is low (11,22,31). Pseudomonas spp. are isolated from the studied patients and 80.0% of the isolated microorganisms showed some drug resistance mechanism, in accordance with previous studies which point multiresistant microorganisms as emergent problems for the NIs control, specially in HIV-1 infected patients (31).
A microorganism is considered multiresistant when presents resistance at least three groups of antibiotics (16). The multiresistant microorganisms are more common in intensive treatment units than in other hospital units, due to the number of risk factors and intensive antimicrobial drugs use. The P. aeruginosa infections are considered of difficult treatment, and the most serious cases are described in weakness situations, compromised people or with reduced resistance (20). According with Table 3, 60.0% of patients infected with Pseudomonas spp. had clinical evolution to death, that confirmed the severity of this infection in the studied cases.

Sorvillo and co-workers (31) described P. aeruginosa as one of the main etiologic agent of urinary tract infections in AIDS patients, as showed in 80.0% of the cases in the present study. This find can be due to the excess of invasive procedures in these patients and to its virulence factors, which are also associated with the infections in other sites in these patients (20,22,28).

Another agent isolated with high frequency in this study was E. coli. This gram-negative microorganism integrates the group of the enterobacteria, common infectious agents acquired by the community as in the hospital environment and that are able to cause infections in the most diverse topographies. E. coli is characterized by presenting an ample drug resistance pattern due its capacity in degrading these drugs through the extended spectrum beta-lactamas (ESBL). The ESBL are enzymes that confer resistance to the antibiotic treatment with cefalosporines, constituting a sufficiently common hospital problem, since E. coli and Klebsiella spp. are common nosocomial pathogens (8). In this work, E. coli was isolated only from patients with CD4+ T cells counts lower than 50/mm³ and these microorganisms didn’t show any antibiotic resistance, what can be related with the favoring environment created by the immune system weakness.

Fungal agents as Histoplasma capsulatum (14), Cryptococcus neoformans (35), Candida spp. (18,33) and mycobacteria (2,5,10,12,17) are also important causes of infection in HIV/AIDS patients. All these microorganisms are related both with multiple drug-resistance and the clinical evolution and can be potentially lethal (30). In this study, however, these microorganisms were not isolated in the cultures, a result that can be in part related to the difficult grown of these agents in the usual culture media which determines the low sensitivity for laboratorial diagnosis.

High mortality rate was observed in the HIV-1 infected patients of this study and the disease progression to death was observed in 45.8% of them. The association of the mortality with the NIs and with the CD4+ T cell was not observed, probably, because of the limited number of HIV-infected patients included in this study. The mortality rate observed could not be associated only to the NIs, since these patients were at risk for other common non-nosocomial diseases such as the reactivation of latent infections caused by the deterioration of the immune system and HIV disease progression.

CONCLUSION

The HIV infection has a complex and heterogeneous clinical course and the ability to control the infection and the disease progression depends of the balance between host and viral factors. With the deterioration of the immune system and HIV disease progression, revealed by the CD4+ T cell counts, the AIDS patients are at an increased risk for NIs that contribute to the morbidity, to the significant increase of the length of hospitalization, and to the mortality of them. In this way, the results of nosocomial pathogens characterization and its relation with the immune status revealed by the CD4+ T cell counts in HIV/AIDS patients could contribute to advances in the hospital cares, to the improvement of the life quality of these patients and costs reduction for the health institutions.

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RESUMO

Infecções hospitalares em pacientes infectados com HIV-1 e com AIDS: principais microrganismos e perfil imunológico

Avanços na terapia anti-retroviral têm proporcionado aos pacientes com AIDS um aumento na sobrevida. Ao mesmo tempo, a permanência de pacientes soropositivos no ambiente nosocomial torna-se comum não só pelos efeitos colaterais desta terapia, mas também pelas diversas doenças oportunistas que acometem estes indivíduos dentro e fora do ambiente hospitalar. Durante o período de internação, a fragilidade do sistema imunológico é exposta à virulência da microbiota nosocomial, adquirindo infecções hospitalares graves e muitas vezes fatais. Dentre as diversas síndromes de infecções hospitalares descritas, pouco se sabe sobre estas infecções em pacientes imunocomprometidos e sobre como o estado imunológico é capaz de determinar o curso destas infecções. Este trabalho teve como objetivo determinar os principais microrganismos envolvidos nas infecções hospitalares de pacientes soropositivos para a infecção pelo HIV-1 e descrever a associação com seu perfil imunológico. Realizou-se análise de dados de notificações do Serviço de Controle de Infecção Hospitalar do Hospital Universitário, Londrina, Paraná, na região sul do Brasil, no período de julho de 2003 a julho de 2004. Do total de casos estudados (n=969), 24 pacientes (2,5%) tinham o diagnóstico de AIDS, sendo metade do gênero feminino, com contagem média...
de células T CD4+ de 158,4/mm³. A principal topografia foi o sítio pulmonar (50,0%), sendo Staphylococcus aureus, Pseudomonas aeruginosa e Escherichia coli os principais microrganismos isolados. Observou-se maior incidência de infecção em pacientes com contagem de células T CD4+ menor que 50/mm³. O estudo da relação entre sistema imunológico e microrganismos causadores de infecções poderá contribuir para melhorias nos cuidados de pacientes com AIDS e avanços nos sistemas de controle de infecção hospitalar.

**Palavras-chave:** AIDS, contagem de células TCD4+, infecções nosocomiais, imunocomprometidos.

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