

RESEARCH ARTICLE

Open Access



Prevalence and risk factors for Human T-Lymphotropic Virus Type 1 (HTLV-1) among maintenance hemodialysis patients

Rilma F. S. Santos^{1,2*}, Gildásio C. Conceição³, Márcia S. Martins⁴, Angiolina Kraychete⁴, Maria A. C. Penalva⁵, Edgar M. Carvalho^{6,7}, Antonio Alberto Lopes^{4,7} and Paulo Novis Rocha^{2,7}

Abstract

Background: Infection with the human T-cell lymphotropic virus type 1 (HTLV-1), although asymptomatic in most cases, can lead to potentially grave consequences, such as adult T-cell leukemia-lymphoma and HTLV-1-associated myelopathy / tropical spastic paraparesis. Its prevalence varies widely across different populations and geographic regions. A population-based study in the city of Salvador, located in the Northeast region of Brazil, showed an overall prevalence of HTLV-1 seropositivity of 1.7%. Blood borne virus infections are recognized as important hazards for patients and staff in maintenance hemodialysis (MHD) units but most studies focus on hepatitis B, hepatitis C and human immunodeficiency viruses. There are scarce data about HTLV-1 infection in the MHD population. We aimed to determine the prevalence and risk factors for HTLV-1 infection among MHD patients in the city of Salvador-Bahia, Brazil.

Methods: We conducted a multi-center, cross-sectional study nested in a prospective cohort of MHD patients enrolled from four outpatient clinics. HTLV-1 screening was performed with ELISA and positive cases were confirmed by Western Blot. Factors associated with HTLV-1 seropositivity were identified by multivariable logistic regression.

Results: 605 patients were included in the study. The overall prevalence of HTLV-1 infection was 2.48% (15/605), which was similar to that of hepatitis B [1.98% (12/605)] and C [3.14% (19/605)] viruses in our sample. HTLV-1 seropositivity was positively associated with age [prevalence odds ratio (POR) 1.04; 95% confidence interval (CI) 1.01–1.08], unmarried status (POR 3.65; 95% CI 1.13–11.65), and history of blood transfusion (POR 3.35; 95% CI 1.01–11.13).

Conclusions: The overall prevalence of HTLV-1 infection in a sample of MHD patients was similar to that of other viral infections, such as hepatitis B and C. Our data revealed that MHD patients who are older, unmarried or who have received blood transfusions are at higher risk for HTLV-1 infection.

Keywords: Blood Transfusion, Chronic Kidney Disease, Hemodialysis, Hepatitis B, Hepatitis C, Human T-Lymphotropic Virus Type 1

Background

The human T-cell lymphotropic virus type 1 (HTLV-1), the first human retrovirus identified [1], is the etiologic agent of adult T-cell leukemia-lymphoma (ATLL) and HTLV-1-associated myelopathy / tropical spastic paraparesis (HAM/TSP) [2–5]. HAM/TSP is the most severe form of neurologic manifestation of HTLV-1 infection, but a large percentage of infected subjects present with

milder forms of neurologic involvement, such as erectile dysfunction or neurogenic bladder [6, 7]. Patients with HTLV-1-related neurogenic bladder may present with urinary symptoms that mimic urinary tract infections [8] and ultimately evolve to obstructive kidney failure [9]. More recently, HTLV-1 infection has been linked to other manifestations, such as uveitis, Sicca syndrome, chronic periodontitis, infective dermatitis, polymyositis and polyneuropathy [10–12], although most seropositive patients remain asymptomatic for prolonged periods.

* Correspondence: rilma_santos@yahoo.com.br

¹Division of Nephrology, Hospital Geral Roberto Santos, Salvador, Brazil

²Postgraduate Program in Health Sciences, Federal University of Bahia, Salvador, Brazil

Full list of author information is available at the end of the article



HTLV-1 infection has become a worldwide epidemic, but studies focusing on population prevalence are scarce and stem mainly from endemic areas [13]. The distribution of the virus is not homogeneous and its prevalence varies according to the geographic area and characteristics of the population, with a spatial variation within each region [13–15]. Data obtained mainly from non-population-based studies indicate that the areas with the highest prevalence of HTLV-1 infection are Southern Japan, Caribbean islands, west and central regions of Africa, Malanesia islands, some regions of the Middle East and some regions of South America [16–19].

In Brazil, most of the prevalence data have been obtained from serological screening of asymptomatic blood donors. These data show a great variation in prevalence within the country, being highest in the northeast region [20–23]. In northeast Brazil, the highest prevalence of HTLV-1 infection is observed in the city of Salvador, capital of the state of Bahia. In a population-based study, the overall prevalence of HTLV-1 seropositivity in Salvador was 1.7% [95% CI 1.1–2.5%] [24].

Blood borne virus infections are recognized as important hazards for patients and staff in maintenance hemodialysis (MHD) units but most studies focus on hepatitis B (HBV), hepatitis C (HCV) and human immunodeficiency virus (HIV). Few studies were specifically designed to estimate the prevalence of HTLV-1 infection in MHD patients [25–30]. These studies, conducted in the last century, revealed that, similarly to what is observed in the general population, the prevalence of HTLV-1 infection among MHD patients varies according to the geographic area. In a study of patients undergoing MHD in several regions of Japan, the average prevalence of HTLV-1 infection was 2.7% but there was a wide variation across regions, reaching up to 21.2% in Okinawa, a highly endemic area of HLTV-1 among asymptomatic blood donors [26]. A more recent study, conducted in the Mazandaran province in north Iran, a non-endemic region of HLTV-1, found a prevalence of HTLV-1 of 0.6% [31]. We found no published data about the prevalence of HTLV-1 in MHD patients treated in Brazil.

Given the high prevalence of HTLV-1 infection in the general population in Salvador-Bahia and the scarcity of data about this infection in the MHD population, we conducted a cross-sectional study nested in a prospective, multicenter cohort of MHD patients, to identify the prevalence and factors associated with HTLV-1 seropositivity in this population.

Methods

Subjects and sites

We screened MHD patients enrolled in the Prospective Study of the Prognosis of Chronic Hemodialysis Patients (PROHEMO) cohort. The current phase of the PROHEMO

study cohort began in 2010; it includes MHD patients from four (out of a total of twelve) outpatient dialysis units in the city of Salvador, the capital of Bahia, located in the northeast region of Brazil [32, 33]. We do not have an epidemiologic profile of the remaining eight clinics, but we believe the four clinics studied to be representative of all clinics in Salvador. We included all adult (18 or older) members of the PROHEMO cohort who were alive and undergoing MHD at the time of the collection of blood samples for HTLV-1 serology screening (between August and October 2011). At this time, the cohort contained 695 patients; we excluded 90 patients due to lack of essential clinical data. Therefore, our analyses were conducted on 605 patients, corresponding to 87% (605/695) of the entire cohort.

Design

Cross-sectional study nested in a prospective cohort study of MHD patients. Information on socio-demographic, laboratorial and clinical variables were obtained directly from the PROHEMO database, which contains data prospectively collected from patients and medical records. Information on HTLV-1 status was obtained by screening stored blood samples. Between August and October 2011, at the time of blood collection for monthly dialysis labs, additional 10 ml samples of blood were drawn and stored at -20°C for future analyses. These samples were screened for antibodies against HTLV 1/2, using an enzyme immunoassay (EIA) - Gold ELISA HTLV-1/2 (REM Indústria e Comércio LTDA) and the positive cases were confirmed by a Western Blot (WB) immunoassay (HTLV Blot 2.4, MP Biomedicals Asia Pacific Pte Ltd. Singapore).

The economic class of the patients (A, B, C, D and E) was determined using the classification system of the Brazilian Institute of Market Research (Abipeme) that considers consumer goods, such as refrigerators, televisions and telephones. Patients in classes D and E were categorized as poor or very poor [34].

A variable “history of blood transfusion” was available in the dataset. Participants of the cohort were asked the following question “Did you receive blood transfusion over the past three months?” There were no data on transfusions past this period or on the number of units transfused.

Ethical issues

All participants signed an informed consent form agreeing with the use of their clinical information and collection and testing of blood samples for HTVL serology for research purposes. The medical director of each participating dialysis unit signed a consent letter approving the study protocol. The Institutional Review Board of the Medical School of Bahia of the Federal University of Bahia approved the study protocol.

Sample size calculation

In December 2011, the population of MHD patients of the city of Salvador was estimated to be around 2000 patients (unpublished data from the State's Health Secretariat). Using the software Openepi (www.openepi.com), we estimated that 548 patients would be needed to detect a prevalence of HTLV-1 seropositivity in MHD patients in Salvador of 2%, with 95% confidence limits extending 1% on either side.

Statistical analyses

Continuous data were summarized using mean \pm standard deviation (SD) or median and interquartile range (IQR); categorical variables were summarized using relative frequencies. Comparisons of continuous variables between seropositive and seronegative groups were performed using the Student's *t*-test for independent samples; categorical variables were compared using the chi square (with continuity correction) or Fischer's exact test. Variables that showed an association with HTLV-1 seropositivity characterized by a *p* value ≤ 0.25 in univariate analyses were selected for a multivariable logistic regression model to identify independent predictors of HTLV-1 seropositivity. We used an automated multivariable logistic regression method called forward stepwise. Briefly, in this method the computer tests the variables one by one for entry into the model, based on the significance level of the score statistic. The variable with the smallest *p* value is entered into the model (entry criterion: *p* < 0.05). After each entry, variables that are already in the model are tested for possible removal. The variable with the largest *p* value is removed (removal criterion: *p* > 0.10), and the model is re-estimated. Variables in the model are then evaluated again for removal. When no more variables satisfy the removal criterion, covariates that are not in the model are evaluated for entry. Model building stops when no more variables meet entry or removal criteria or when the current model is the same as a previous model. A *p* value ≤ 0.05 in final analyses was considered statistically significant. The analyses were performed using SPSS for Windows version 21.0 and the 95% confidence interval of the HTLV-1 prevalence was calculated using the bootstrap (50 replications) of the STATA, version 12.0.

Results

Sixteen patients were HTLV-1 seropositive by the ELISA technique but one was considered a false positive due to a negative Western Blot. Therefore, 15 out of 605 MHD patients studied had confirmed HTLV-1 infection, for an overall prevalence of 2.48% (95% CI 1.22–3.74%). There were no cases of HTLV-2 seropositivity in our sample.

In two out of these fifteen cases (13.3%), the ESRD was attributed to urological complications of HTLV-1 infection, specifically neurogenic bladder leading to obstructive kidney failure. The first case was a 70 year-old female,

wheelchair-bound due to HAM/TSP secondary to HTLV-1. She had a well-documented case of neurogenic bladder and was followed by a team of Neurologists and Urologists at an ambulatory clinic specialized in HTLV-1 at the University Hospital of the Federal University of Bahia. She did not have diabetes or hypertension and other causes of ESRD were excluded on clinical grounds. The second case was a 47 year-old man who initially presented to the emergency department of a public general hospital with uremia. He was found to have a distended bladder and bilateral hydronephrosis on ultrasound, with a normal-sized prostate gland. A Foley catheter was placed, but renal failure did not improve and he remained dialysis-dependent. He had no prior history of hypertension or diabetes.

Table 1 shows the socio-demographic, clinical and laboratorial characteristics of our sample, stratified by HTLV-1 serology. In this univariate analysis, unmarried status and a history of blood transfusion were the only variables significantly associated with a positive HTLV-1 serology. The mean age tended to be greater and time on dialysis tended to be lower in the seropositive group. Overall, there were 12/604 patients (1.99%) with HBV and 19/605 (3.14%) with HCV; the frequency of positive HBV or HCV serology tended to be higher in the HTLV-1 group. Seropositive patients also had higher serum creatinine, higher parathyroid hormone (PTH) and lower leucocyte count whereas hemoglobin levels did not differ among groups (data not shown).

Figure 1 shows the prevalence and prevalence ratio of HTLV-1 seropositivity in subgroups of MHD patients. The highest prevalence ratios were encountered in those who received blood transfusions, unmarried and with HBV or HCV. The prevalence of HTLV-1 infection also increased with age, going from 1.6 to 2.2% and 2.9% among those 18–30, 31–50 and > 50 years old, respectively.

Table 2 shows the multivariable logistic regression to identify variables independently associated with HTLV-1 seropositivity. In this analysis, the variables that showed statistically significant associations with HTLV-1 seropositivity were older age, unmarried status and a history of blood transfusion.

Discussion

Our initial hypothesis was that the prevalence of HTLV-1 in the MHD population would be greater than that observed in the general population. This hypothesis was based on older literature data from other countries [27–29, 35] and on the following postulates: a) ESRD could be a risk factor for HTLV-1, mediated by anemia and greater frequency of blood transfusion or due to the risk of acquiring the virus during the hemodialysis procedure, as it is known to happen with HBV and HCV [36, 37]; b) HTLV-1 and end stage renal disease (ESRD) could share risk factors, such as poverty and health illiteracy [38, 39]; c) HTLV-1

Table 1 Demographic, clinical and laboratorial variables of 605 maintenance hemodialysis patients from four outpatient dialysis units in Salvador, Bahia, Brazil, stratified by HTLV-1 serology

Independent variables	HTLV (+) N = 15	HTLV (-) N = 590	POR (95% CI)	<i>p</i>
Demographic				
Age, years	52.5 (46.9–71.0)	47.8 (37.7–59.0)	1.03 (1.00–1.07)	0.080
Female (vs. male)	8/15 (53.3)	224/590 (38.0)	1.87 (0.67–5.22)	0.234
Non-white (vs. white)	13/15 (86.7)	513/590 (86.9)	1.03 (0.23–4.63)	0.974
Poor or very poor (vs. classes A thru C)	10/15 (66.7)	265/580 (45.7)	2.38 (0.80–7.04)	0.118
Less than high school (vs. high school or greater)	12/15 (80.0)	359/589 (61.0)	2.56 (0.72–9.18)	0.148
Unmarried (vs. married)	11/15 (73.3)	261/588 (44.4)	3.45 (1.09–10.95)	0.036
Clinical				
Dialysis by catheter (vs. AV fistula)	4/15 (26.7)	76/588 (12.9)	1.12 (0.31–4.02)	0.867
Time on MHD, months	20.4 (4.5–58.4)	47.8 (19.6–97.4)	0.99 (0.97–1.00)	0.064
EPO, IU ($\times 10^3$)/week	12 (6–12)	10 (6–12)	1.00 (1.00–1.00)	0.870
Diabetes mellitus (vs. no)	5/15 (33.3)	100/590 (16.9)	2.45 (0.82–7.32)	0.109
Systemic hypertension (vs. no)	13/15 (86.7)	533/590 (90.3)	0.70 (0.15–3.16)	0.638
Blood transfusion (vs. no)	4/15 (26.7)	55/582 (9.2)	3.48 (1.07–11.31)	0.038
Hepatitis B (vs. no)	1/15 (6.7)	11/589 (1.9)	3.75 (0.45–31.10)	0.220
Hepatitis C (vs. no)	1/15 (6.7)	18/590 (3.1)	2.27 (0.28–18.21)	0.440
Hepatitis B or C (vs. no)	2/15 (13.3)	28/589 (4.8)	3.08 (0.66–14.33)	0.151

Continuous data (Age, Time on MHD and EPO dose) are expressed as median (P25–P75%); categorical data are expressed as n/total (%); POR Prevalence Odds Ratio, EPO Erythropoietin, MHD maintenance hemodialysis. For continuous data, POR reflects the odds of a positive HTLV serology with each increase in one unit of the continuous variable (one year for Age, one month for Time on MHD, and 10^3 IU/week for EPO); for categorical data, POR reflects the odds of a positive HTLV serology in the variable of interest compared to its complement. History of blood transfusion was limited to available data that reported transfusion only for the previous three months. *P* values in bold highlight the variables that were selected for the multivariable logistic regression model. Although a positive serology for Hepatitis B was associated with a $p \leq 0.25$, it was not included in the multivariable model because we preferred to use the variable Hepatitis B or C.

could be a risk factor for ESRD as the urological manifestations of the virus, such as neurogenic bladder, can lead to post-renal kidney failure [9]. Nevertheless, the overall prevalence of HTLV-1 seropositivity in our sample of 605 MHD patients was 2.48% (95% CI 1.22–3.74%), which is only marginally higher than that observed by Dourado et al. in a population-based study in the city of Salvador (1.7; 95% CI 1.1–2.5%) [24].

In their study, Dourado et al. studied individuals from 1 to 89 years. The prevalence of HTLV-1 infection increased with age, going from 0.3 to 1.1 to 1.1 and 8.4% among those 0–15,

16–30, 31–50 and > 50 years old, respectively. In the present study of MHD patients, the prevalence of HTLV-1 infection also increased with age, going from 1.6% in those 18–30 to 2.9% among those > 50 years old; for every increase in one year of age, we found a 4% increase in the odds of having a positive HTLV-1 serology. A potential explanation for this association is that increasing age provides a greater length of exposure to events that might result in acquiring the virus, such as sexual activity, intravenous drug use or blood transfusion. Since there is no cure for HTLV-1, once acquired, the infection is carried forward over time (cohort effect).

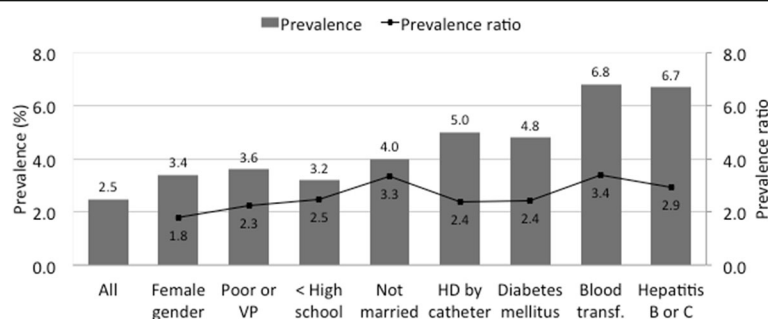


Fig. 1 Prevalence and prevalence ratio of HTLV-1 seropositivity in subgroups of maintenance hemodialysis patients from four satellite clinics in Salvador, Bahia, Brazil

Table 2 Multivariable logistic regression to identify variables independently associated with HTLV-1 seropositivity in 605 maintenance hemodialysis patients from four satellite clinics in Salvador, Bahia, Brazil

Independent variables	POR	95% CI		p
		Lower	Upper	
Age, years	1.04	1.01	1.08	0.041
Unmarried (vs. married)	3.65	1.13	11.75	0.030
Blood transfusion (vs. no)	3.35	1.01	11.13	0.048

Nine variables were selected to enter the automated, forward stepwise multivariable logistic regression model based on a p value ≤ 0.25 on univariate analyses: age (continuous), gender (female as reference), marital status (unmarried as reference), time on dialysis, history of blood transfusion (yes/no), seropositivity for hepatitis B or C virus (yes/no), socioeconomic status ("poor or very poor" as reference), level of schooling ("less than high school" as reference), and diabetes (yes / no). History of blood transfusion was limited to available data that reported transfusion only for the previous three months

The association between HTLV-1 infection and blood transfusion is well established and has been confirmed in several studies using blood donors in different countries [39–41], including Brazil [38]. Blood transfusions among MHD patients were much more common before the advent of recombinant human erythropoietin (rhEPO) [42, 43]; indeed, this might help explain the high prevalence of HTLV-1 in the MHD population of some Japanese studies published before the advent of rhEPO [25, 26]. Currently, the routine use of rhEPO markedly reduced the need for blood transfusion, particularly among those patients who are already established on MHD programs for more than three months [44].

In Brazil, however, there is still a considerable need for blood transfusions among patients who are recent on MHD, since many of them start dialysis emergently, without the benefit of pre-dialysis medical care. This was illustrated in a study conducted at the largest public hospital in Salvador [45], which is a referral service for the entire state of Bahia. The authors studied all patients who initiated MHD at that institution after being admitted through the emergency department and found that 57% of them first learned about their kidney disease at the time of diagnosis of ESRD. Moreover, 71% of them had not been evaluated by a nephrologist before arriving at the emergency department. Their mean hemoglobin at admission was 7.7 mg/dL; no patient had received rhEPO previously and 50% of them required blood transfusion during the hospitalization [45]. In the present study, we found that a history of blood transfusion tripled the odds of having a positive HTLV-1 serology. Spread of HTLV-1 during the hemodialysis procedure seems less likely, based on the biology of the virus. Some studies indicate that transmission of HTLV-1 requires the contact between living cells and not simply cell-free body fluids [46, 47].

In their population-based study, Dourado et. al showed that the prevalence of HTLV-1 was greater among females, those with lower income, worse living conditions

and less education [24]. In the present study of the MHD population, we also found a numerically higher prevalence of HTLV-1 infection among females, those who were poor or very poor and those who had less than high school education, although these associations did not reach statistical significance. Unmarried status, however, more than tripled the odds of having a positive HTLV-1 serology. We posit that unmarried status might have functioned as a proxy for sexual promiscuity and/or unprotected sex, which are known risk factors for HTLV-1 infection [38, 39].

In Brazil, the main etiologies of ESRD are hypertension (35%), diabetes mellitus (30%), chronic glomerulonephritis (12%) and polycystic kidney disease (4%) [48]. Herein, urological complications of HTLV-1 infection were felt to be the cause of ESRD in only 0.3% of patients (2/605). We are not aware of other studies that evaluated the contribution of HTLV-1 to ESRD.

Our study has several limitations. An inherent limitation of cross sectional studies is that associations do not imply causality. We utilized the database of an ongoing multicenter cohort study of MHD in our area and all of our analyses were limited to the variables that were present in their database. We did not have information on sexual practices, living conditions, intravenous drug use or neurologic symptoms. Moreover, our data on blood transfusion was categorical (yes/no) and restricted to the previous three months; we had no information on lifetime exposure to blood products. Finally, our study provides no information about HIV co-infection as none of the four MHD units studied admit HIV positive patients.

Conclusion

The prevalence of HTLV-1 infection in MHD patients from four hemodialysis units in northeast Brazil was not higher than that encountered in a population-based study of the same geographic area. HTLV-1 seropositivity in MHD patients was similar to that of other viral infections, such as HBV and HCV. Older age, unmarried status and blood transfusions were independently associated with HTLV-1 infection.

Abbreviations

ATLL: Adult T-cell leukemia-lymphoma; Blood Transfusion: Chronic Kidney Disease, Hemodialysis, Hepatitis B, Hepatitis C, Human T-Lymphotropic Virus Type 1; ESRD: End stage renal disease; HAM / TSP: HTLV-1-Associated Myelopathy / Tropical Spastic Paraparesis; HBV: Hepatitis B virus; HCV: Hepatitis C virus; HIV: Human immunodeficiency virus; HTLV-1: Human T-lymphotropic virus type 1; MHC: Maintenance hemodialysis; PROHEMO: Prospective study of the prognosis of chronic hemodialysis patients; PTH: Parathyroid hormone; RhEPO: Recombinant human erythropoietin

Acknowledgements

We are grateful for the support received from the medical directors and staff of the following dialysis units: *Clínica Nephron-Barris, Clínica Nephron-Itapuã, Instituto de Nefrologia e Diálise and Clini Rim.*

Funding

The study was supported by research grants from “The Brazilian National Council for Scientific and Technological Development (CNPq)”, grants #480292/2009-4 and #309468/2009-4. CNPq had no role in the design of the study and collection, analysis, and interpretation of data and in writing the manuscript.

Availability of data and materials

The datasets generated and analysed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

RFSS conceived and designed the study; acquired data; analyzed and interpreted the data; drafted the initial version of the manuscript; wrote the final version of the manuscript, incorporating suggestions from the reviewers and coauthors. GCC performed the laboratorial analyses and contributed importantly with the manuscript drafting and with its corrections. MSM acquired data and contributed importantly with the manuscript drafting and with its corrections. AK acquired data and contributed importantly with the manuscript drafting and with its corrections. MACP acquired data and contributed importantly with the manuscript drafting and with its corrections. EMC acquired data and contributed importantly with the manuscript drafting and with its corrections. AAL conceived and designed the study; acquired data; analyzed and interpreted the data; critically reviewed the manuscript for important intellectual content. PNR conceived and designed the study; acquired data; analyzed and interpreted the data; provided statistical support; critically reviewed the manuscript for important intellectual content. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

Not applicable.

Ethics approval and consent to participate

All participants signed an informed consent form agreeing with the use of their clinical information and collection and testing of blood samples for HTLV serology for research purposes. The medical director of each participating dialysis unit signed a consent letter approving the study protocol. The Institutional Review Board of the Medical School of Bahia of the Federal University of Bahia approved the study protocol.

Author details

¹Division of Nephrology, Hospital Geral Roberto Santos, Salvador, Brazil. ²Postgraduate Program in Health Sciences, Federal University of Bahia, Salvador, Brazil. ³Department of Biochemistry, APAE, Salvador, Brazil. ⁴Unit of Clinical Epidemiology and Evidence Based Medicine of the Professor Edgard Santos University Hospital, Federal University of Bahia, Salvador, Brazil. ⁵Diretoria Médica da Clínica Nefron-Itapua, Salvador, Brazil. ⁶Centro de Pesquisas Gonçalo Moniz (Fiocruz-Ba), Salvador, Brazil. ⁷Department of Internal Medicine and Diagnostic Support, Medical School of Bahia, Federal University of Bahia, Salvador, Brazil.

Received: 5 October 2016 Accepted: 9 February 2017

Published online: 15 February 2017

References

- Poiesz BJ, Ruscetti FW, Gazdar AF, Bunn PA, Minna JD, Gallo RC. Detection and isolation of type C retrovirus particles from fresh and cultured lymphocytes of a patient with cutaneous T-cell lymphoma. *Proc Natl Acad Sci U S A*. 1980;77:7415–9.
- Gessain A. Antibodies to human T-lymphotropic virus type-I in patients with tropical spastic paraparesis. *Lancet*. 1985;326:407–10.
- Kaplan JE, Osame M, Kubota H, Igata A, Nishitani H, Maeda Y, Khabbaz RF, Janssen RS. The risk of development of HTLV-I-associated myelopathy/tropical spastic paraparesis among persons infected with HTLV-I. *J Acquir Immune Defic Syndr*. 1990;3:1096–101.
- Tajima K, Kuroishi T. Estimation of rate of incidence of ATL among ATL (HTLV-I) carriers in Kyushu, Japan. *Jpn J Clin Oncol*. 1985;15:423–30.
- Takatsuki K, Yamaguchi K, Kawano F, Hattori T, Nishimura H, Tsuda H, Sanada I, Nakada K, Itai Y. Clinical diversity in adult T-cell leukemia-lymphoma. *Cancer Res*. 1985;45(9 Suppl):4644s–5.
- Souza A, Tanajura D, Toledo-Cornell C, Santos S, de Carvalho EM. Immunopathogenesis and neurological manifestations associated to HTLV-1 infection. *Rev Soc Bras Med Trop*. 2012;45:545–52.
- Tanajura D, Castro N, Oliveira P, Neto A, Muniz A, Carvalho NB, Orge G, Santos S, Glesby MJ, Carvalho EM. Neurological manifestations in Human T-cell Lymphotropic Virus Type 1 (HTLV-1)-infected individuals without HTLV-1-associated myelopathy/tropical spastic paraparesis: a longitudinal cohort study. *Clin Infect Dis*. 2015;61:49–56.
- Rocha PN, Rehem AP, Santana JF, Castro N, Muniz AL, Salgado K, Rocha H, Carvalho EM. The cause of urinary symptoms among Human T Lymphotropic Virus Type I (HTLV-I) infected patients: a cross sectional study. *BMC Infect Dis*. 2007;7:15.
- Araujo AQ-C. Update on neurological manifestations of HTLV-1 infection. *Curr Infect Dis Rep*. 2015;17:459.
- Ferraz-Chaoui AK, Atta AM, Atta MLS, Galvão-Castro B, Santiago MB. Study of autoantibodies in patients with keratoconjunctivitis sicca infected by the human T cell lymphotropic virus type 1. *Rheumatol Int*. 2010;30:775–8.
- Garlet GP, Giozza SP, Silveira EM, Claudino M, Santos SB, Avila-Campos MJ, Martins W, Cardoso CR, Trombone APF, Campanelli AP, Carvalho EM, Silva JS. Association of human T lymphotropic virus 1 amplification of periodontitis severity with altered cytokine expression in response to a standard periodontopathogen infection. *Clin Infect Dis*. 2010;50:e11–8.
- Höllsberg P, Hafler DA. Seminars in medicine of the Beth Israel Hospital, Boston. Pathogenesis of diseases induced by human lymphotropic virus type I infection. *N Engl J Med*. 1993;328:1173–82.
- Hlela C, Shepperd S, Khumalo NP, Taylor GP. The prevalence of human T-cell lymphotropic virus type 1 in the general population is unknown. *AIDS Rev*. 2009;11:205–14.
- Gessain A, Cassar O. Epidemiological aspects and world distribution of HTLV-1 infection. *Front Microbiol*. 2012;3:1–23.
- Proietti FA, Carneiro-Proietti ABF, Catalan-Soares BC, Murphy EL. Global epidemiology of HTLV-I infection and associated diseases. *Oncogene*. 2005;24:6058–68.
- Hedayati-Moghaddam MR, Fathimoghdam F, Eftekharzadeh Mashhadi I, Soghandi L, Bidkhori HR. Epidemiology of HTLV-1 in Neyshabour, Northeast of Iran. *Iran Red Crescent Med J*. 2011;13:424–7.
- Minamoto GY, Gold JWM, Scheinberg DA, Hardy WD, Chein N, Zuckerman E, Reich L, Dietz K, Gee T, Hoffer J, Mayer K, Gabrilove J, Clarkson B, Armstrong D. Infection with human T-cell leukemia virus type I in patients with leukemia. *N Engl J Med*. 1988;319:243–4.
- Mueller N. The epidemiology of HTLV-I infection. *Cancer Causes Control*. 1991;2:37–52.
- Verdonck K, González E, Van Dooren S, Vandamme A-M, Vanham G, Gotuzzo E. Human T-lymphotropic virus 1: recent knowledge about an ancient infection. *Lancet Infect Dis*. 2007;7:266–81.
- Carneiro-Proietti ABF, Ribas JGR, Catalan-Soares BC, Martins ML, Brito-Melo GEA, Martins-Filho OA, Pinheiro SR, de Araújo A Q-C, Galvão-Castro B, de Oliveira MSP, Guedes AC, Proietti FA. Infection and disease caused by the human T cell lymphotropic viruses type I and II in Brazil. *Rev Soc Bras Med Trop*. 2002;35:499–508.
- Catalan-Soares B, Carneiro-Proietti ABF, Proietti FA. Heterogeneous geographic distribution of human T-cell lymphotropic viruses I and II (HTLV-I / II): serological screening prevalence rates in blood donors from large urban areas in Brazil. *Cad Saude Publica*. 2005;21:926–31.
- Ferreira Júnior OC, Vaz RS, Carvalho MB, Guerra C, Fabron AL, Rosembit J, Hamerschlak N. Human T-lymphotropic virus type I and type II infections and correlation with risk factors in blood donors from São Paulo, Brazil. *Transfusion*. 1995;35:258–63.
- Galvão-Castro B, Loures L, Rodrigues LG, Sereno A, Ferreira Júnior OC, Franco LG, Muller M, Sampaio DA, Santana A, Passos LM, Proietti F. Distribution of human T-lymphotropic virus type I among blood donors: a nationwide Brazilian study. *Transfusion*. 1997;37:242–3.
- Dourado I, Alcantara LCJ, Barreto ML, da Gloria TM, Galvão-Castro B. HTLV-I in the general population of Salvador, Brazil: a city with African ethnic and sociodemographic characteristics. *J Acquir Immune Defic Syndr*. 2003;34:527–31.
- Lee SY, Mastushita K, Machida J, Tajiri M, Yamaguchi K, Takatsuki K. Human T-cell leukemia virus type I infection in hemodialysis patients. *Cancer*. 1987;60:1474–8.
- Morikawa K, Kuroda M, Tofuku Y, Uehara H, Akizawa T, Kitaoka T, Koshikawa S, Sugimoto H, Hashimoto K. Prevalence of HTLV-1 antibodies in hemodialysis patients in Japan. *Am J Kidney Dis*. 1988;12:185–93.

27. De Rossi A, Vertolli U, Romagnoli G, Bertoli M, Dalla Gassa O, Chieco-Bianchi L. LAV/HTLV-III and HTLV-I antibodies in hemodialysis patients. *Nephron*. 1986;44:377–8.
28. Méry JP, Fessi H, Dosquet P, Dazza MC, Simon F, Elias A. HTLV-1 antibodies in hemodialysis patients. *Nephron*. 1992;61:124.
29. Devito C, Pampuro S, Del Pino N, Martinez Peralta L, Libonatti O. HTLV-I/II survey on hemodialysis patients in Buenos Aires. *J Acquir Immune Defic Syndr Hum Retrovirol*. 1996;12:525–6.
30. Davies J, Jabbar Z, Gagan F, Baird RW. Blood-borne viruses in the haemodialysis-dependent population attending Top End Northern Territory facilities 2000–2009. *Nephrology (Carlton)*. 2012;17:501–7.
31. Ghaffari J, Ebrahimi M, Makhloogh A, Mohammadjafari H, Nazari Z. Seroepidemiology of human T-cell lymphotropic virus 1 infection in hemodialysis patients: should we be concerned about it? *Iran J Kidney Dis*. 2013;7:187–90.
32. Lopes GB, Matos CM, Leite EB, Martins MTS, Martins MS, Silva LF, Robinson BM, Port FK, James SA, Lopes AA. Depression as a potential explanation for gender differences in health-related quality of life among patients on maintenance hemodialysis. *Nephron Clin Pract*. 2010;115:c35–40.
33. Lopes GB, Nogueira FCP, de Souza MR, Penalva MA, de Amorim JL, Pisoni RL, Robinson BM, Lopes AA. Assessment of the psychological burden associated with pruritus in hemodialysis patients using the kidney disease quality of life short form. *Qual Life Res*. 2012;21:603–12.
34. Critério ABIPEME. <ftp://ftp.est.ufmg.br/pub/lupericio/ABIPEME.doc>.
35. Perez G, Ortiz-Interian C, Lee H, de Medina M, Cerney M, Allain JP, Schiff E, Parks E, Parks W, Bourgoignie JJ. Human immunodeficiency virus and human T-cell leukemia virus type I in patients undergoing maintenance hemodialysis in Miami. *Am J Kidney Dis*. 1989;14:39–43.
36. Fabrizi F, Messa P. Transmission of hepatitis C virus in dialysis units: a systematic review of reports on outbreaks. *Int J Artif Organs*. 2015;38:471–80.
37. Sowole L, Labbett W, Patel M, O'Riordan A, Cross J, Davenport A, Haque T. The prevalence of occult hepatitis B virus (HBV) infection in a large multi-ethnic haemodialysis cohort. *BMC Nephrol*. 2015;16:12.
38. Soares BC, de Proietti Ab FC, Proietti FA. HTLV-I/II and blood donors: determinants associated with seropositivity in a low risk population. *Rev Saude Publica*. 2003;37:470–6.
39. Hedayati-Moghaddam MR, Tehrani F, Bayati M. Human T-Lymphotropic Virus Type I (HTLV-1) infection among Iranian blood donors: first case-control study on the risk factors. *Viruses*. 2015;7:5736–45.
40. Schreiber GB, Murphy EL, Horton JA, Wright DJ, Garfein R, Chien HC, Nass CC. Risk factors for human T-cell lymphotropic virus types I and II (HTLV-I and -II) in blood donors: the Retrovirus Epidemiology Donor Study. *NHLBI Retrovirus Epidemiology Donor Study*. *J Acquir Immune Defic Syndr Hum Retrovirol*. 1997;14:263–71.
41. O'Brien SF, Goldman M, Scalia V, Yi Q-L, Fan W, Xi G, Dines IR, Fearon MA. The epidemiology of human T-cell lymphotropic virus types I and II in Canadian blood donors. *Transfus Med*. 2013;23:358–66.
42. Crowley JP, Nealey TA, Metzger J, Pono L, Chazan JA. Transfusion and long-term hemodialysis. *Arch Intern Med*. 1987;147:1925–8.
43. Schiller GJ, Berkman SA. Hematologic aspects of renal insufficiency. *Blood Rev*. 1989;3:141–6.
44. MacDougall IC, Obrador GT. How important is transfusion avoidance in 2013? *Nephrol Dial Transplant*. 2013;28:1092–9.
45. Godinho TM, Lyra TG, Braga PS, de Queiroz RA, Alves JA, Kraychete AC, Gusmão ENA, Lopes AA, Rocha PN. Profile of patients initiating chronic hemodialysis at a public hospital in Brazil. *J Bras Nefrol*. 2006;28:96–103.
46. Igakura T, Stinchcombe JC, Goon PKC, Taylor GP, Weber JN, Griffiths GM, Tanaka Y, Osame M, Bangham CRM. Spread of HTLV-I between lymphocytes by virus-induced polarization of the cytoskeleton. *Science*. 2003;299:1713–6.
47. Bangham CRM. The immune control and cell-to-cell spread of human T-lymphotropic virus type 1. *J Gen Virol*. 2003;84(Pt 12):3177–89.
48. Sesso RC, Lopes AA, Thomé FS, Lugon JR, Dos Santos DR. Brazilian Chronic Dialysis Survey 2013 - trend analysis between 2011 and 2013. *J Bras Nefrol*. 2014;36:476–81.

Submit your next manuscript to BioMed Central and we will help you at every step:

- We accept pre-submission inquiries
- Our selector tool helps you to find the most relevant journal
- We provide round the clock customer support
- Convenient online submission
- Thorough peer review
- Inclusion in PubMed and all major indexing services
- Maximum visibility for your research

Submit your manuscript at
www.biomedcentral.com/submit

