Peripheral Blood Mononuclear Cells from Individuals Infected with Human T-Cell Lymphotropic Virus Type 1 Have a Reduced Capacity To Respond to Recall Antigens

Rita Elizabeth Mascarenhas,1,4 Cláudia Brodskyn,2 Geisa Barbosa,1,4 Jorge Clarêncio,2 Antônio Souza Andrade-Filho,3,4 Frederico Figueiroa,3,4 Bernardo Galvão-Castro,1,4 and Fernanda Grassi1,4∗

Advanced Laboratory of Public Health/Gonçalo Moniz Research Center (CPqGM), Oswaldo Cruz Foundation (FIOCRUZ), Salvador, Bahia, Brazil; Microbiology and Immunoregulation of Integrate Laboratory/Gonçalo Moniz Research Center (CPqGM), Oswaldo Cruz Foundation (FIOCRUZ), Salvador, Bahia, Brazil; Neurology and Neurosurgery Foundation, Salvador, Bahia, Brazil; and Bahiana School of Medicine and Public Health, Salvador, Bahia, Brazil

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Evidence indicates that human T-cell lymphotropic virus type 1 (HTLV-1) infection leads to chronic immunosuppression and a greater susceptibility to infectious diseases. Spontaneous in vitro proliferation of peripheral blood mononuclear cells (PBMC) is an important immunological feature of HTLV-1-infected individuals. However, the association between spontaneous proliferation and immunosuppression is not clear. In this study, we evaluated the cellular immune responses of PBMC from 58 asymptomatic HTLV-1-infected individuals with PBMC showing or not showing spontaneous proliferation. Individuals with PBMC that spontaneously proliferated had increased proportions of CD4 T cells expressing CD45RO and dramatically reduced responses to recall antigens. In addition, frequencies of positive responses to recall antigens were also decreased in HTLV-infected individuals without spontaneous proliferation of PBMC. There was a polyclonal expansion of multiple T-cell receptor Vβ families of CD4+ T lymphocytes in patients with spontaneous proliferation. We observed that HTLV-1 induced an immunosuppression characterized by a decrease in the stimulation index to a recall antigen, even in individuals who did not present spontaneous proliferation. On the other hand, only patients with PBMC presenting spontaneous proliferation showed polyclonal activation and increased proportion of CD4 T cells expressing CD45RO.

Human T-cell lymphotropic virus type 1 (HTLV-1) is a human retrovirus that has infected 10 to 20 million people worldwide (3). There are areas in Japan, the Caribbean, Melanesia, central and western Africa, and South America that are large endemic foci (17, 35). In Brazil, the overall prevalence of HTLV-1 infection in blood donors is 0.45%, and the city of Salvador (located in the northeast) has the highest prevalence (1.35%) in the country (4). Recently, a population-based cross-sectional study performed in Salvador, Brazil, found an HTLV-1 seroprevalence of 1.76% (2).

Two diseases are clearly associated with HTLV-1 infection: adult T-cell leukemia/lymphoma (6, 26, 34) and HTLV-1-associated myelopathy/tropical spastic paraparesis (5, 24). Although 95% of infected people remain asymptomatic (7), there is evidence that HTLV-1 has a much broader spectrum of disease manifestations, such as uveitis (20), arthritis (8, 12), polymyositis (22), lymphocytic interstitial pneumonia (31), and infective dermatitis in children (15). Severe immunosuppression is well documented in patients with adult T-cell leukemia/lymphoma (10, 19). However, a growing body of literature suggests that many HTLV-1-infected individuals show chronic immunosuppression, even in the absence of malignant disease. HTLV-1-infected individuals are more susceptible to several infectious diseases, such as strongyloidiasis (1, 30), Hansen’s disease (11), and tuberculosis (21, 25). In addition, they have reduced cutaneous delayed-type hypersensitivity responses to purified protein derivative (PPD) of Mycobacterium tuberculosis (14, 23, 33, 36).

The immunologic hallmark of HTLV-1-infected individuals is a spontaneous in vitro proliferation of their peripheral blood mononuclear cells (PBMC) (without any added antigen) observed in approximately fifty percent of patients (13, 27, 29). The memory CD4+ CD45RO+ subset is the principal T-cell population involved in this phenomenon (29). In this study, we evaluated the cellular immune responses of HTLV-1-infected individuals (with or without PBMC spontaneous proliferation) against candidin, cytomegalovirus (CMV), PPD, and tetanus toxoid (TT) recall antigens.

We observed that HTLV-1-infected individuals had reduced stimulation indexes to recall antigens, even when their PBMC did not spontaneously proliferate. On the other hand, only the individuals with spontaneous PBMC proliferation (SP+) had polyclonal T-cell activation (as shown by the study of T-cell receptor Vβ [TCR-Vβ] families) and an increase in the proportion of CD4 T cells expressing CD45RO.

MATERIALS AND METHODS

Patients. Fifty-eight asymptomatic HTLV-1-infected individuals studied at the HTLV-1 Reference Development of Science Foundation (Salvador, Bahia, Brazil) were included in the study. The group consisted of 47 women (81%) and 11 men (19%) and had a mean age of 43 years. Blood samples from 12 healthy
TABLE 1. Spontaneous proliferation of PBMC from HTLV-1-infected individuals\textsuperscript{a}

<table>
<thead>
<tr>
<th>PBMC</th>
<th>No. of HTLV-1-infected individuals</th>
<th>Mean cpm ± SD (range)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SP+</td>
<td>39</td>
<td>1,656 ± 956 (520-4,629)</td>
</tr>
<tr>
<td>SP-</td>
<td>19</td>
<td>217 ± 152 (18-476)</td>
</tr>
<tr>
<td>Total</td>
<td>58</td>
<td></td>
</tr>
</tbody>
</table>

\textsuperscript{a} The mean counts per minute of nonstimulated PBMC from uninfected controls was less than 100 cpm. For the HTLV-1-infected individuals, the mean cpm from nonstimulated cells was \(\leq 500\) cpm at day 6 for HTLV-1-infected individuals with SP+ PBMC, while it was \(\leq 500\) cpm at day 6 for HTLV-1-infected individuals with SP- PBMC.

TABLE 2. Percentages of T-cell subsets from HTLV-1-infected individuals and uninfected controls

<table>
<thead>
<tr>
<th>Group and parameter</th>
<th>CD4\textsuperscript{a}</th>
<th>CD4\textsuperscript{b}</th>
<th>CD4\textsuperscript{c}</th>
<th>CD4\textsuperscript{d}</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group A (SP+ HTLV-infected subjects [n = 19])</td>
<td>Mean 41</td>
<td>29</td>
<td>79</td>
<td>24</td>
</tr>
<tr>
<td>SD</td>
<td>14</td>
<td>12</td>
<td>10</td>
<td>6</td>
</tr>
<tr>
<td>Median</td>
<td>44</td>
<td>30</td>
<td>79</td>
<td>23</td>
</tr>
<tr>
<td>Range</td>
<td>10-61</td>
<td>10-53</td>
<td>55-92</td>
<td>12-37</td>
</tr>
<tr>
<td>Group B (SP- HTLV-infected subjects [n = 10])</td>
<td>Mean 43</td>
<td>32</td>
<td>67</td>
<td>21</td>
</tr>
<tr>
<td>SD</td>
<td>9</td>
<td>14</td>
<td>18</td>
<td>10</td>
</tr>
<tr>
<td>Median</td>
<td>45</td>
<td>31</td>
<td>68</td>
<td>19</td>
</tr>
<tr>
<td>Range</td>
<td>26-56</td>
<td>13-56</td>
<td>46-99</td>
<td>10-41</td>
</tr>
<tr>
<td>Group C (uninfected controls [n = 10])</td>
<td>Mean 42</td>
<td>37</td>
<td>67</td>
<td>21</td>
</tr>
<tr>
<td>SD</td>
<td>9</td>
<td>11</td>
<td>11</td>
<td>4</td>
</tr>
<tr>
<td>Median</td>
<td>40</td>
<td>37</td>
<td>64</td>
<td>22</td>
</tr>
<tr>
<td>Range</td>
<td>31-55</td>
<td>23-57</td>
<td>51-86</td>
<td>15-30</td>
</tr>
</tbody>
</table>

\textsuperscript{a} CD4\textsuperscript{a} cells expressing CD45RA.
\textsuperscript{b} CD4\textsuperscript{b} cells expressing CD45RO.
\textsuperscript{c} Differences between the values for SP+ and SP- HTLV-1-infected groups and uninfected controls were compared by the Kruskal-Wallis test (group A versus group B versus group C). Differences between two groups were analyzed by the Mann-Whitney U test (group A versus group B, group A versus group C, and group B versus group C). Differences between the values for SP+ and SP- HTLV-1-infected groups and uninfected controls that were statistically significant are shown in bold type.

RESULTS

PBMC spontaneous proliferation. As shown in Table 1, spontaneous PBMC proliferation was observed in 39 out of 58 (67%) HTLV-1-infected individuals (1,656 cpm ± 956 cpm; range, 520 to 4,629 cpm).

On the other hand, no spontaneous PBMC proliferation was observed in 19 out of 58 (33%) HTLV-1-infected individuals (217 ± 152 cpm; range, 18 to 476). In this case, the mean counts per minute for asymptomatic HTLV-1-infected individuals did not differ from the mean counts per minute for uninfected controls.

counts obtained in the presence of antigen to mean counts obtained without antigen. A stimulation index of \(\geq 3\) indicated a positive proliferative response. Therefore, in this study, a mean counts per minute of \(\geq 500\) for nonstimulated cells i.e., three times the mean counts per minute for uninfected non-stimulated cells, namely 159 ± 138 cpm; range, 6 to 462) was considered spontaneous proliferation of PBMC.

Statistical analyses. Data are expressed as means and standard deviations. We compared mean values of the percentages of CD4 and CD8 T-cell subsets and stimulation indexes of proliferative responses to recall antigen for the three groups (patients with spontaneous proliferation of PBMC, those without spontaneous proliferation, and controls) using the Kruskal-Wallis test. If a significant difference was found, a Mann-Whitney U test was performed. The statistical analysis of TCR-V\textsuperscript{b} was performed with the Wilcoxon signed-rank test. A P value of less than 0.05 denoted a statistically significant difference. BioEstat 3.0 software (Societ\‘d\‘ Academic Maimuru/MCT-CNPq) was used for all statistical analyses.
affected controls (159 ± 138 cpm; range, 6 to 462) (P = 0.32 by the Mann-Whitney U test).

**HTLV-1-infected individuals have an expansion of the memory CD4<sup>+</sup> CD45RO<sup>+</sup> T-cell subset.** As shown in Table 2, the percentage of circulating CD4<sup>+</sup> T lymphocytes in the HTLV-1-infected group with PBMC with spontaneous proliferation (41% ± 14%) was similar to those of HTLV-1-infected individuals without spontaneous PBMC proliferation (43% ± 9%) and of uninfected controls (42% ± 9%) (P = 0.84). However, the memory CD4<sup>+</sup> CD45RO<sup>+</sup> T-cell subset was expanded in HTLV-1-infected individuals with spontaneous PBMC proliferation (79% ± 10%) compared to that in HTLV-1-infected individuals without spontaneous proliferation (67% ± 18%) (P = 0.04) and uninfected controls (67% ± 11%) (P = 0.01). Additionally, the proportion of CD4 T cells expressing CD45RO was positively correlated to spontaneous proliferation of PBMC in culture (Spearman’s r = 0.37; P = 0.04) (data not shown).

There were no significant differences in the frequencies of naive CD4<sup>+</sup> CD45RA<sup>+</sup> and CD8<sup>+</sup> T-cell subsets in HTLV-infected individuals (with or without spontaneous PBMC proliferation) and in uninfected controls.

**PBMC from HTLV-1-infected individuals have a lower capacity to proliferate in response to recall antigens.** HTLV-1-infected individuals with spontaneous proliferation showed the lowest stimulation indexes to all antigens tested (candidin, PPD, tetanus toxoid, and cytomegalovirus antigens) compared to HTLV-1-infected individuals without spontaneous proliferation and uninfected controls. Only 5.3% (1 out of 19) responded to candidin, while 4.8% (1 out of 21) responded to CMV or TT (Fig. 1). Besides that, HTLV-1-infected SP− patients also showed decreases in stimulation indexes: 50% of them (4 out of 8) responded to candidin, 33% (4 out of 12) responded to PPD, 42% (5 out of 12) to cytomegalovirus, and 50% (6 out of 12) to tetanus toxoid antigen. Almost all uninfected controls had positive responses to recall antigens: 90% to candidin (9 out of 10), 83% to PPD (10 out of 12), 100% to tuberculin (11 out of 11), and 67% (8 out of 12) to CMV.

In addition, PBMC from HTLV-1-infected individuals re-
sponded to fewer recall antigens than uninfected controls: while 83% of uninfected controls had specific PBMC proliforative responses to three or more different antigens, PBMC from HTLV-1-infected SP+ individuals were unable to recognize more than one antigen \( (P < 0.0001) \). Moreover, only 33\% of HTLV-1-infected individuals without spontaneous PBMC proliferation responded to three or more antigens \( (P = 0.038) \).

TCR-Vβ analysis disclosed polyclonal expansion only in HTLV-1-infected individuals with spontaneous proliferation of PBMC. As shown in Fig. 2A, polyclonal expansion of CD4+ cells (TCR-Vβ families 2, 3.1, 6.1, 8, 11, 12, 16, 17, 21.3, and 22) was observed in patients with spontaneous PBMC proliferation. In contrast, HTLV-1-infected individuals without spontaneous PBMC proliferation had CD4+ TCR-Vβ repertoires similar to those of uninfected individuals (Fig. 2B).

**DISCUSSION**

Spontaneous proliferation is an immunological hallmark of PBMC from HTLV-1-infected individuals. This phenomenon is observed in up to 50\% of cultures from PBMC from HTLV-1 carriers (28). In this study, we evaluated the cellular immune responses from HTLV-1-infected individuals with PBMC showing and not showing spontaneous proliferation. Compared to uninfected controls, HTLV-1-infected individuals with PBMC presenting spontaneous proliferation had an increased proportion of CD4 T cells expressing CD45RO. Moreover, TCR-Vβ analysis showed a polyclonal expansion of almost all Vβ families studied concerning CD4 T-cell subsets. In addition, HTLV-1-infected individuals with spontaneous PBMC proliferation had markedly reduced stimulation indexes for candidin, PPD, tetanus toxoid, and cytomegalovirus.
Therefore, responses to these recall antigens could not be detected by cell proliferation.

The absence of detectable cellular immune responses to recall antigens in individuals with spontaneous PBMC proliferation was expected, since high levels of spontaneous proliferation would mask a specific response to the recall antigens evaluated by the stimulation index. However, previous findings support the idea that there could be different levels of T-cell anergy among HTLV-I-infected individuals.

One of the most important observations of this study was that cells from HTLV-I-infected individuals with PBMC without spontaneous proliferation showed significant decreases in the stimulation indexes to candidin, PPD, and tetanus toxoid. In addition, only 33% were able to recognize more than two antigens; in contrast, 83% of uninfected controls were able to recognize three or more antigens. In those individuals, detectable proliferation showed significant decreases in the stimulation indexes to candidin, PPD, and tetanus toxoid.

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