Dysfunctional Epstein-Barr virus (EBV)-specific CD8(+) T lymphocytes and increased EBV load in HIV-1 infected individuals


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Dysfunctional Epstein-Barr virus (EBV)--specific CD8\(^+\) T lymphocytes and increased EBV load in HIV-1 infected individuals progressing to AIDS-related non-Hodgkin lymphoma

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Introduction

Epstein-Barr virus (EBV) is a widespread human gamma herpesvirus. Primary infection with EBV usually occurs asymptptomatically,\(^1\) whereafter the virus persists for life in a latent form in B lymphocytes.\(^2\) The initial expansion and reactivation of these latently infected B lymphocytes is controlled by specific CD8\(^+\) major histocompatibility complex (MHC) class I–restricted cytotoxic T cells. Data on EBV-specific cellular immunity were correlated with EBV load. For comparison, individuals who progressed to AIDS with opportunistic infections (AIDS-OI) and long-term asymptomatics (LTAs) were studied. The number of virus-specific T cells was detected using tetrameric HLA–EBV-peptide complexes; function of these EBV-specific T cells was determined using the interferon-\(\gamma\) (IFN-\(\gamma\)) Elispot assay. It was observed that EBV-specific CD8\(^+\) T cells were present in normal numbers in human immunodeficiency virus (HIV)-infected individuals. However, their functional capacity was decreased compared with HIV\(^-\) individuals. In AIDS-NHL patients, EBV-specific T cells were not physiologically lost in the course of HIV-1 infection but showed progressive loss of their capability to produce IFN-\(\gamma\) in response to EBV peptides. This loss of function correlated with lower CD4\(^+\) T-cell numbers and was accompanied by increasing EBV load.

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these new methods allowed us to investigate at the peptide level whether EBV-specific CD8+ T cells become either deleted or dysfunctional in patients who have developed AIDS-NHL. In addition, using a real-time quantitative polymerase chain reaction (PCR) assay, EBV DNA in infected cells was quantitated. Presence and function of CD8+ T lymphocytes specific for peptides from both latent and lytic EBV antigens were studied, and EBV load was measured in the course of HIV-1 infection in AIDS-NHL patients. For comparison, HIV-1–infected individuals who progressed to AIDS with opportunistic infections (AIDS-OI) or who were long-term asymptomatic (AIDS-0I) were studied.

**Patients, materials, and methods**

**Study population**

This study was performed on participants of the Amsterdam Cohort Studies on AIDS and HIV-1 infection. From these individuals at risk of HIV-1 infection, we selected HIV+ homosexual males according to duration of follow-up, availability of samples, and HLA type. Blood samples from these individuals were collected every 3 months for HIV-1 serology and immunologic studies. In addition, at all times peripheral blood mononuclear cells (PBMCs) were cryopreserved.

We analyzed longitudinal PBMC samples from 5 HIV-1 infected individuals progressing to AIDS-related diffuse large cell NHL (NHL006, NHL0118, NHL6118, NHL0139, NHL0308), starting at or soon after HIV-1 seroconversion. For comparison, we studied PBMC samples from 3 HIV-1–infected individuals progressing to AIDS (classification of the Centers for Disease Control, 1993) with opportunistic infections (PROG0232, PROG0341, PROG0642) and 3 HIV-infected LTA (LTA0036, LTA0057 39 A2, 19; B27, 40 162 0.34 —

Intracellular IFN-γ staining after antigen-specific stimulation

A total of 2 × 10^6 PBMCs per milliliter were either stimulated with 1 μg EBV (RAK) peptide, used in the tetrameric complexes, or PMA/ionomycin (positive control) or not stimulated (medium alone as negative control) at 37°C for 4 hours in the presence of 3 μM monensin. This stimulation was sufficient to induce IFN-γ production in most potential cells. Stronger stimulation protocols (10 μg peptide per milliliter for 6 hours) did not substantially increase the number of IFN-γ–producing cells. After incubation, cells were washed and stained in PBS supplemented with 0.5% (vol/vol) BSA for 15 minutes with HLA-B8–RAK tetramers (APC) and

**Table 1. Characteristics of HIV-1–infected individuals**

<table>
<thead>
<tr>
<th>Participant</th>
<th>Age*</th>
<th>HLA class I</th>
<th>Follow-up†</th>
<th>CD4 counts‡</th>
<th>Diagnosis</th>
</tr>
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<tbody>
<tr>
<td>NHL0118</td>
<td>36</td>
<td>A2, 3; B51, 60</td>
<td>124</td>
<td>&lt; 0.19</td>
<td>NHL</td>
</tr>
<tr>
<td>NHL0139</td>
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<td>A2, B8, 51</td>
<td>87</td>
<td>&lt; 0.06</td>
<td>NHL</td>
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<tr>
<td>NHL0308</td>
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<td>A1, B8, 51</td>
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</tr>
<tr>
<td>NHL0606</td>
<td>41</td>
<td>A2, 32; B7, 35</td>
<td>76</td>
<td>0.1</td>
<td>NHL</td>
</tr>
<tr>
<td>NHL6118</td>
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<td>A2, 3; B8, 37</td>
<td>25</td>
<td>0.19</td>
<td>NHL</td>
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<tr>
<td>PROG0341</td>
<td>33</td>
<td>A24, 2B, B8, 13</td>
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<td>0.09</td>
<td>PCP</td>
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<tr>
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<td>PCP</td>
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<td>0.36</td>
<td>—</td>
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<tr>
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<td>A2, 19, B27, 40</td>
<td>162</td>
<td>0.34</td>
<td>—</td>
</tr>
<tr>
<td>LTA1160</td>
<td>35</td>
<td>A2, B8, 27</td>
<td>128</td>
<td>0.24</td>
<td>—</td>
</tr>
</tbody>
</table>

NHL indicates B-cell NHL; PCP, Pneumocystis carinii pneumonia; Mycobact, Mycobacterium tuberculosis extrapulmonary.

*Age in years at HIV-1 seroconversion or first seropositive visit.
†Follow-up visit, in months, from HIV-1 seroconversion or first seropositive visit until AIDS diagnosis or last time point studied.
‡CD4+ T-cell numbers x 1000/mL at AIDS diagnosis or last time point studied for LTAs.
PerCP-conjugated mAb CD8 (Becton Dickinson). After membrane staining, cells were washed with PBS/BSA and fixed with 4% paraformaldehyde, permeabilized (permeabilization kit, Becton Dickinson), and stained intracellularly with IFN-α-PE (Becton Dickinson) for 30 minutes at 4°C. At least 200,000 events in the lymphocyte gate were acquired using a FACSCalibur flow cytometer (Becton Dickinson).

**DNA extraction and real-time quantitative TaqMan assay**

PBMCs (1 × 10⁶) were lysed by addition of L6-lysis buffer.²⁰ Genomic DNA was extracted by precipitation with isopropanol, and DNA from 2 × 10⁵ cells was amplified using PCR primers selective for the EBV DNA genome encoding the nonglycosylated membrane protein BNRF1 p143.²¹,²² PCR amplification was performed as previously described²¹ using EBV/p143 forward and reverse primers resulting in a 74-base pair DNA product. In the PCR reaction, a fluorogenic EBV/p143-specific probe was added with a FAM reported molecule attached to the 5’ end and a TAMRA quencher linked at the 3’ end to detect amplified DNA. Amplification and detection were performed with an ABI Prism 7700 Sequence Detection System (PE Applied Biosystems, Foster City, CA). Real-time measurements were taken, and a threshold cycle value was calculated for each sample by determining the point at which the fluorescence exceeded a threshold limit of 0.04. Each run contained several negative controls (no template or EBV DNA), a positive control (a known amount of EBV copies), and a standard dilution of plasmid DNA containing the PCR product as insert, which was calibrated with an EBV-quantified standard (Advanced Biotechnologies, Epsom, United Kingdom). The analyzed sensitivity of the assay was between 50 and 5 × 10⁶ copies per milliliter. All reactions were performed in duplicate and only considered positive when both replications were above the threshold limit. The variation between duplicates was as low as 7.5%.

**Statistical analysis**

To compare EBV-specific tetramer+ T cells, IFN-γ-producing CD8 + T cells, and EBV load early and late in HIV infection, Wilcoxon tests were performed. To compare all variables early or late between groups, Mann-Whitney tests were performed using the software program SPSS 7.5 (SPSS, Chicago, IL). To test the relation between CD4 + T-cell numbers and EBV-specific functional CD8 + T cells, EBV-tetramer + T cells, or total CD8 + T cells, regression analyses (mixed linear model) were performed. Regression analysis and multivariate analysis to search for predictors of the number of functional T cells were performed after cube root transformation of all variables. To correct for a possible correlation between multiple observations from one person, compound symmetry was used as correlation structure using the Proc Mixed procedure of the software program SAS.

**Results**

**Direct visualization and functional analysis of EBV-specific CD8 + T lymphocytes**

To investigate the presence and function of EBV-specific CD8 + T cells, we studied CD8 + T cells specific for 2 epitopes derived from lytic antigens, the HLA-A2–restricted epitope GLCTLVAML (A2-GLC) and the HLA-B8–restricted epitope RAKFKQLL (B8-RAK), and for 1 epitope derived from a latent antigen, the HLA-B8–restricted epitope FLRGRAYGL (B8-FLR). To detect the presence of antigen-specific CD8 + T cells, we stained antigen-specific T cells using tetrameric HLA-peptide complexes. To assess the function of antigen-specific CD8 + T cells, we used the IFN-γ Elispot assay, which shows the number of IFN-γ-producing T cells after peptide stimulation. To determine the sensitivity of both assays, we applied them both to an HIV-specific T-cell clone that was selected in vitro to respond to one specific (HIV RT) peptide.²⁴ In Figure 1 (left panel), tetramer staining of this T-cell clone is shown, revealing that 87% of the T-cell clone was specific for the tetramer containing the specific peptide. IFN-γ Elispot revealed that virtually all tetramer + T cells from the T-cell clone produced IFN-γ upon stimulation with the specific peptide (Figure 1A, right panel), because almost every (93%) tetramer + T cell could be accounted for in the Elispot assay. In addition, IFN-γ production was similar in both Elispot assay and intracellular FACS staining (manuscript in preparation). Tetramer staining of PBMCs from a healthy EBV-carrying individual showed that approximately 3% of the CD8 + T cells were specific for an EBV-peptide (Figure 1B, left panel), half of which produced IFN-γ after peptide stimulation in a direct ex vivo assay using intracellular staining (Figure 1B, right panel) or Elispot assay (data not shown). Furthermore, both assays were reproducible and sensitive on both fresh and frozen material (data not shown).

In preliminary studies using IFN-γ Elispot assay, the selected EBV epitopes were shown to be immunodominant in that they were recognized by a high frequency of CD8 + T cells (100–400 peptide-specific T cells per 10⁶ PBMCs) in all HIV-infected individuals studied. HLA-A2–restricted latent antigen-specific T cells were either not present or present in a very low range (data not shown). Therefore, HLA-A2–restricted T cells against latent epitopes were not studied.

**Lower numbers of functional EBV-specific CD8 + T cells in HIV + versus HIV – individuals**

After immunodominance was established, the number and function of EBV peptide–specific CD8 + T lymphocytes was measured in HIV + and HIV – individuals in the course of HIV-1 infection. For
comparisons, both the total number of circulating (tetramer+ as percentage of CD8+ T cells as assessed by tetramer staining) and (B) proportion of IFN-γ-producing T cells of tetramer+ T cells (percentage of tetramer+ T cells determined after peptide stimulation) are shown for HIV− (n = 6, filled circles) and HIV+ individuals (n = 11) early in HIV infection (average of time points studied until CD4+ T cells drop below 200/μL or halfway through follow-up). HIV− individuals include individuals progressing to AIDS-NHL (n = 5, open triangles) or AIDS-OI (n = 3, open circles) and LTA HIV-infected individuals (n = 3, filled triangles).

In this study population, 0.2% to 5% of the CD8+ T cells were EBV-specific. (Figure 2A). No difference in the percentage of tetramer+ EBV-specific CD8+ T cells was observed between HIV− and HIV+ individuals early in HIV infection (P = .42, Mann-Whitney test) (Figure 2A). Interestingly, in HIV− individuals a higher proportion of these EBV-specific T cells produced IFN-γ in response to the selected dominant EBV peptides (median 22%, range 12%-28%) than in HIV+ individuals early in HIV infection (median 13%, range 6%-15%) (P = .014, Mann-Whitney test) (Figure 2B).

**Progressive loss of function of EBV-specific CD8+ T lymphocytes in AIDS-NHL patients**

To study the cause of the defective EBV-immune surveillance in AIDS-NHL patients, we investigated both the number and function of EBV-specific CD8+ T cells in the course of HIV-1 infection in AIDS-NHL patients. For comparison, progressors to AIDS-OI and LTA individuals were studied. Detailed results from 2 study participants (1 LTA, 1 AIDS-NHL) are shown in Figure 3. Most EBV-specific CD8+ T cells were directed against the lytic epitope B8-RAK (0.7%-3.6% in Figure 3C), whereas fewer T cells were directed against the latent epitope B8-FLR (0.1%-1.1% in Figure 3B). In HLA-A2+ individuals, 0.2% to 1.2% of the T cells were directed against the lytic epitope A2-GLC (0.6%-0.8% in Figure 3D).

![Figure 2. Percentage tetramer+ EBV-specific T cells and functionality of tetramer+ T cells in HIV− and HIV+ individuals.](image)

**Figure 2.** Percentage tetramer+ EBV-specific T cells and functionality of tetramer+ T cells in HIV− and HIV+ individuals. (A) The number of tetramer+ (percentage of CD8+ T cells as assessed by tetramer staining) and (B) proportion of IFN-γ-producing T cells of tetramer+ T cells (percentage of tetramer+ T cells determined after peptide stimulation) are shown for HIV− (n = 6, filled circles) and HIV+ individuals (n = 11) early in HIV infection (average of time points studied until CD4+ T cells drop below 200/μL or halfway through follow-up). HIV− individuals include individuals progressing to AIDS-NHL (n = 5, open triangles) or AIDS-OI (n = 3, open circles) and LTA HIV-infected individuals (n = 3, filled triangles).

![Figure 3. Presence and function of lytic and latent EBV antigen-specific CD8+ T lymphocytes in an AIDS-NHL patient (left) and an LTA individual (right).](image)

**Figure 3.** Presence and function of lytic and latent EBV antigen-specific CD8+ T lymphocytes in an AIDS-NHL patient (left) and an LTA individual (right). On the x-axis, follow-up is indicated in months after HIV-1 entry in the study. The arrows indicate the time of NHL-diagnosis (NHL) and start of therapy (DDI). The vertical dotted line indicates the time point at which CD4+ T-cell counts drop below 200/μL. (A) Longitudinal analysis of CD4+ and CD8− T-lymphocyte numbers. (B) Longitudinal analysis of B8-FLR–specific CD8+ T lymphocytes assessed by tetramer staining (percentage of CD8+ T cells, solid line) and IFN-γ Elispot assay (per 106 CD8+ T cells, dashed line); (C) IFN-γ. (D) Longitudinal analysis of A2-GLC–specific CD8+ T lymphocytes assessed by tetramer staining (solid line) and IFN-γ Elispot assay (dashed line), as described in “Patients, materials, and methods.” (A) indicates tetramer; (B), IFN-γ; (C), IFN-γ.
For all individuals, the total percentage of tetramer$^+$ and number of IFN-γ–producing (functional) EBV-specific CD8$^+$ T cells was calculated from the individual peptide-specific CD8$^+$ T cells. In Figure 4, average numbers of EBV-specific CD8$^+$ T cells early and late in HIV-1 infection are shown for the 3 groups studied. Figures 5 and 6 give detailed longitudinal information for several of the investigated participants.

In most individuals no change in the percentage of tetramer$^+$ T cells was observed in the course of HIV-1 infection (Figure 4A). In AIDS-NHL, patients, the function of these EBV-specific T cells significantly decreased in the course of HIV-1 infection ($P = .04$, Wilcoxon test) (Figure 4B). The decrease in IFN-γ–producing cells (6 IFN-γ) tended to be stronger than the decrease in tetramer$^+$ cells (6 tetramer$^+$) corrected for the first time point (−29% and −6%, respectively, $P = .07$, Mann-Whitney test), indicating a discrepancy in the kinetics of the number of EBV-specific T cells present in comparison with their functionality.

The decrease in IFN-γ–producing T cells was most pronounced in patients NHL6006, NHL0308, and NHL0118, in whom almost no IFN-γ–producing CD8$^+$ T cells were left after a few years of HIV-1 infection (NHL6006, left panel in Figure 5B,C; and B8-FLR–specific T cells in NHL0308, Figure 2B) or near lymphoma diagnosis (NHL0118, middle panel in Figure 5B). This loss of function was observed for both lytic and latent antigen-specific responses (patient NHL0308, Figure 2B,C).

Furthermore, the decline in IFN-γ–producing T cells, while the number of tetramer$^+$ T cells remained stable, leads to a loss of the proportion of tetramer$^+$ T cells that produced IFN-γ in individuals progressing to AIDS-NHL (Figure 4C). In contrast, in LTAs the proportion of tetramer$^+$ T cells that produced IFN-γ even increased in the course of HIV-1 infection. As a result, the decrease in IFN-γ (Δ IFN-γ) in AIDS-NHL patients tended to be stronger than the delta IFN-γ in LTAs corrected for the first time point (−29% and +3%, respectively, $P = .07$, Mann-Whitney test) (Figure 4C).

Progressors to AIDS-OI showed lower but stable percentages of tetramer$^+$ T cells (Figure 4A) and IFN-γ–producing T cells specific for EBV (Figure 4B), resulting in a stable proportion of IFN-γ$^+$ / tetramer$^+$ T cells (Figure 4C). Interestingly, PROG0232, a slow progressor to AIDS, showed an increase in the total number of IFN-γ–producing T cells just after the start of highly active antiretroviral therapy (HAART) (Figure 6B).

### Comparison of the number of IFN-γ–producing T cells determined by Elispot assay and intracellular staining

To confirm our findings with the Elispot assay, we also performed intracellular IFN-γ staining after peptide stimulation (B8-RAK) of PBMCs from an AIDS-NHL patient at an early time point (Figure 7, left panel) and PBMCs from an LTA individual at 2 time points (Figure 7, right panel). At an early time point in NHL0308, the percentage of RAK-specific IFN-γ–producing T cells as assessed by intracellular staining (0.19%, or 1900 per $10^6$ CD8) (Figure 7B) is of the same magnitude as the number of IFN-γ producers found by Elispot assay (~1900, Figure 7A, dotted line). Similar to the Elispot results, the number of CD8$^+$ T cells with intracellular IFN-γ staining increased in the course of infection for the LTA individual (from 0.16% to 0.3%) after stimulation, which correlated well with IFN-γ production in Elispot (Figure 7A). Moreover, the proportion of IFN-γ–producing tetramer$^+$ T cells for the AIDS-NHL patient early in infection was 9% (Figure 7C), which corresponded well with the percentages observed using tetramer staining in combination with IFN-γ Elispot assay (around 8% early in HIV-1 infection) (Figure 4C).

### EBV-specific CD8$^+$ T-lymphocyte function is dependent on CD4$^+$ T cells

Next, we investigated whether there was a possible relation between CD4$^+$ T-cell numbers and the number of EBV-specific CD8$^+$ T cells. In AIDS-NHL patients, CD4$^+$ T-cell numbers

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**Figure 4.** EBV-specific T cells and EBV load early and late in HIV-1 infection. For 3 groups of HIV-infected individuals, from the longitudinal data the average of early (period studied until CD4$^+$ T cells drop below 200/μL or halfway through follow-up) and late (average of time points from the drop in CD4$^+$ T cells below 200/μL or halfway through follow-up until AIDS diagnosis or last time point studied) time points in HIV-infection were calculated excluding time periods on therapy. For AIDS-NHL patients (left), LTA individuals (middle), and AIDS-OI patients (right), the figure depicts (A) the percentage of EBV tetramer$^+$ CD8$^+$ T cells, (B) the number of IFN-γ–producing EBV-specific T cells per $10^6$ CD8$^+$ T cells, (C) the proportion of IFN-γ$^+$ tetramer$^+$ T cells, and (D) EBV load per $10^6$ PBMCs.
Figure 5. Presence and function of EBV-specific CD8+ T cells and EBV load in 3 HIV-1 infected individuals with AIDS-NHL. On the x-axis follow-up is indicated in months after HIV-1 entry in the study. The arrows indicate the time of AIDS-NHL diagnosis (NHL) and start of antiretroviral therapy (DDI). The vertical dotted line indicates the time point at which CD4+ T cell counts drop below 200/μL. (A) Longitudinal analysis of CD4+ and CD8+ T-lymphocyte numbers. (●) indicates CD4; (○) CD8. (B) Longitudinal analysis of EBV-specific CD8+ T lymphocytes as assessed by tetramer staining as a composite of all tested tetramers (solid line) and IFN-γ Elispot assay (dashed line), as described in “Patients, materials, and methods.” (●) indicates tetramer; (○), IFN-γ. (C) Longitudinal analysis of EBV load, expressed as the number of virus copies per 10^6 PBMCs (solid line), in comparison with the number of functional EBV-specific CD8+ T lymphocytes per 10^6 CD8+ T cells (dashed line). (●) indicates EBV load; (○), IFN-γ.

Functional loss of EBV-specific CD8+ T lymphocytes is paralleled by an increase in EBV load

To investigate whether loss of functional T cells is associated with an increase in EBV load and whether this precedes the development of AIDS-NHL, EBV load was determined using a sensitive and specific real-time quantitative PCR. As shown in Figure 4, loss of EBV-specific CD8+ T-cell function (Figure 4B) was associated with an increase in EBV load in AIDS-NHL patients (P = 0.08, Wilcoxon test) (Figure 4D), although the absolute number of EBV copies was not different from absolute EBV load observed in LTAs and other progressors to AIDS (D.v.B. et al, manuscript submitted). Interestingly, in patient NHL0308 after the start of DDI treatment, EBV-specific CD8+ T cells slightly increased and a small reduction in EBV load was observed (Figure 5C).

In contrast, in LTAs no increase and, in progressors to AIDS-OI, even a decrease in EBV load was observed (Figure 4D, middle and right panels). The increase in load (∆ load) in AIDS-NHL patients was significantly different from the decrease in load (∆ load) in LTAs (P = 0.05) and progressors to AIDS-OI (P = 0.03, Mann-Whitney test) (Figure 4D). Interestingly, in LTAs occasional EBV peak loads (Figure 6C) were paralleled by an increase in the number of functional EBV-specific CD8+ T cells (Figure 6B) after which EBV load decreased, suggestive of EBV control. HAART, which led to an increase in the number of IFN-γ–producing CD8+ T cells (Figure 6B), resulted in a decrease of EBV load in progressor PROG0232 (Figure 6C).

To study whether EBV load is able to drive the number of IFN-γ–producing T cells, we performed multivariate analysis including the number of CD4+ T cells, EBV load, and the number of IFN-γ–producing T cells. Although multivariate analysis of all HIV-infected individuals (n = 11) showed that only the number of CD4+ T cells could predict the number of functional cells (β = 0.92, P = 0.0188), analysis of only AIDS-OI patients and
important role for CD4$^+$ T cells and EBV load in 2 HIV-1 infected LTA individuals and 1 HIV-1 infected individual progressing to AIDS-OI. The arrows indicate the time of AIDS diagnosis (O) and start of HAART. (A) Longitudinal analysis of CD4$^+$ and CD8$^+$ T lymphocyte numbers. (●) indicates CD4$^+$, (○) CD8$^+$. (B) Longitudinal analysis of EBV-specific CD8$^+$ T lymphocytes as assessed by tetramer staining as a composite of all tested tetramers (solid line) and IFN-γ Elispot assay (dashed line), as described in "Patients, materials, and methods." (●) indicates tetramer; (○), IFN-γ. (C) Longitudinal analysis of EBV load, expressed as the number of virus copies per 10$^6$ PBMCs (solid line), in comparison with the number of functional EBV-specific CD8$^+$ T lymphocytes per 10$^6$ CD8$^+$ T cells (dashed line). (●) indicates EBV load; (○), IFN-γ.

Discussion

To investigate the cause of decreasing numbers of CTLp in AIDS-NHL patients, which is believed to lead to defective EBV-immune control, we studied number (using MHC class I tetramers) and function (using IFN-γ Elispot assay) of EBV-specific CD8$^+$ T lymphocytes in the course of HIV infection in relation to EBV load. This is, to our knowledge, the first longitudinal study demonstrating a discrepancy between direct visualization using tetramer staining and functional assays enumerating IFN-γ-producing antigen-specific T cells. The major conclusions from our study are that (1) HIV$^+$ individuals have lower numbers of functional EBV-specific CD8$^+$ T cells than HIV$^-$ individuals; (2) in AIDS-NHL patients, EBV-specific CD8$^+$ T cells are lost preferentially at the functional level and are not physically lost; (3) loss of EBV-specific CD8$^+$ T-cell function is correlated with lower CD4$^+$ T-cell numbers; (4) increasing EBV load correlates with loss of EBV-specific immunity; and (5) the number of T cells directed against lytic antigens is higher than against latent antigens.

The observed correlation between loss of function of EBV-specific CD8$^+$ T cells and lower CD4$^+$ T-cell numbers indicates an important role for CD4$^+$ T cells in maintaining the functional capacity of CD8$^+$ T cells. Our data are in good agreement with studies on T-helper dependence of chronic lymphocytic choriomeningitis virus (LCMV)-specific CTL in mouse models.\textsuperscript{25-27} A critical role for CD4$^+$ T cells has also been shown during immunization,\textsuperscript{28} and progressive loss of CTL in the absence of adequate helper cell function has been demonstrated for several murine viral infections.\textsuperscript{29,30} Furthermore, CD4$^+$ T cells also appear to be essential for long-term persistence of adoptively transferred virus-specific CTL in humans.\textsuperscript{27,32,33}

In the natural course of HIV infection, it has been shown that progressors to AIDS lose CD8$^+$ CTLs when functional HIV-specific CD8$^+$ T cells disappear. In contrast, in nonprogressors, who have stable CD4$^+$ T-cell numbers, HIV-specific CTL responses can be sustained for long periods of time,\textsuperscript{34,37} indicating that sustained HIV-specific helper activity is required for maintenance of functional CD8$^+$ T-cell responses.\textsuperscript{34,36,39}

The fact that most EBV-specific CD8$^+$ T cells were directed against lytic epitopes suggests that these lytic antigen-specific T cells play a role not only during acute infection\textsuperscript{40} but also in controlling EBV reactivation by eliminating virus-producing cells at an early stage. Loss of functional lytic antigen-specific CD8$^+$ T cells could therefore lead to an increase in EBV DNA, as we indeed observed. As the pool of EBV-infected B cells grows, there is an increased risk of subsequent genetic hits resulting in malignant outgrowth of EBV-infected B cells. Because functional CD8$^+$ T cells specific for latent antigens appear to be lost as well, newly developed tumor cells will not be destroyed. Our data suggest that loss of both lytic and latent antigen-specific CD8$^+$ T cells may contribute to the risk for development of AIDS-NHL.

In HIV infection, 0.2% to 5% of the CD8$^+$ T cells were found to be EBV-specific, of which 13% were shown to produce IFN-γ in response to EBV peptides. This is lower than in healthy HIV$^+$ EBV carriers, where the percentage of IFN-γ-producing T cells was approximately 22% (our data), consistent with data from literature.\textsuperscript{31} In AIDS-NHL patients, the number of IFN-γ-producing T cells was found to decrease progressively in the course of HIV infection, resulting in a much lower percentage of IFN-γ-producing CD4$^+$ T cells. This suggests that EBV-specific CD8$^+$ T cells may be EBV-specific, of which 13% were shown to produce IFN-γ and function (using IFN-γ Elispot assay) of EBV-specific CD8$^+$ T lymphocytes as assessed by tetramer staining as a composite of all tested tetramers (solid line) and IFN-γ Elispot assay (dashed line), as described in "Patients, materials, and methods." (●) indicates tetramer; (○), IFN-γ. (C) Longitudinal analysis of EBV load, expressed as the number of virus copies per 10$^6$ PBMCs (solid line), in comparison with the number of functional EBV-specific CD8$^+$ T lymphocytes per 10$^6$ CD8$^+$ T cells (dashed line). (●) indicates EBV load; (○), IFN-γ.

Discussion

To investigate the cause of decreasing numbers of CTLp in AIDS-NHL patients, which is believed to lead to defective EBV-immune control, we studied number (using MHC class I tetramers) and function (using IFN-γ Elispot assay) of EBV-specific CD8$^+$ T lymphocytes in the course of HIV infection in relation to EBV load. This is, to our knowledge, the first longitudinal study demonstrating a discrepancy between direct visualization using tetramer staining and functional assays enumerating IFN-γ-producing antigen-specific T cells. The major conclusions from our study are that (1) HIV$^+$ individuals have lower numbers of functional EBV-specific CD8$^+$ T cells than HIV$^-$ individuals; (2) in AIDS-NHL patients, EBV-specific CD8$^+$ T cells are lost preferentially at the functional level and are not physically lost; (3) loss of EBV-specific CD8$^+$ T-cell function is correlated with lower CD4$^+$ T-cell numbers; (4) increasing EBV load correlates with loss of EBV-specific immunity; and (5) the number of T cells directed against lytic antigens is higher than against latent antigens.

The observed correlation between loss of function of EBV-specific CD8$^+$ T cells and lower CD4$^+$ T-cell numbers indicates an important role for CD4$^+$ T cells in maintaining the functional capacity of CD8$^+$ T cells. Our data are in good agreement with studies on T-helper dependence of chronic lymphocytic choriomeningitis virus (LCMV)-specific CTL in mouse models.\textsuperscript{25-27} A critical role for CD4$^+$ T cells has also been shown during immunization,\textsuperscript{28} and progressive loss of CTL in the absence of adequate helper cell function has been demonstrated for several murine viral infections.\textsuperscript{29,30} Furthermore, CD4$^+$ T cells also appear to be essential for long-term persistence of adoptively transferred virus-specific CTL in humans.\textsuperscript{27,32,33}

In the natural course of HIV infection, it has been shown that progressors to AIDS lose CD8$^+$ CTLs when functional HIV-specific CD8$^+$ T cells disappear. In contrast, in nonprogressors, who have stable CD4$^+$ T-cell numbers, HIV-specific CTL responses can be sustained for long periods of time,\textsuperscript{34,37} indicating that sustained HIV-specific helper activity is required for maintenance of functional CD8$^+$ T-cell responses.\textsuperscript{34,36,39}

The fact that most EBV-specific CD8$^+$ T cells were directed against lytic epitopes suggests that these lytic antigen-specific T cells play a role not only during acute infection\textsuperscript{40} but also in controlling EBV reactivation by eliminating virus-producing cells at an early stage. Loss of functional lytic antigen-specific CD8$^+$ T cells could therefore lead to an increase in EBV DNA, as we indeed observed. As the pool of EBV-infected B cells grows, there is an increased risk of subsequent genetic hits resulting in malignant outgrowth of EBV-infected B cells. Because functional CD8$^+$ T cells specific for latent antigens appear to be lost as well, newly developed tumor cells will not be destroyed. Our data suggest that loss of both lytic and latent antigen-specific CD8$^+$ T cells may contribute to the risk for development of AIDS-NHL.

In HIV infection, 0.2% to 5% of the CD8$^+$ T cells were found to be EBV-specific, of which 13% were shown to produce IFN-γ in response to EBV peptides. This is lower than in healthy HIV$^+$ EBV carriers, where the percentage of IFN-γ-producing T cells was approximately 22% (our data), consistent with data from literature.\textsuperscript{31} In AIDS-NHL patients, the number of IFN-γ-producing T cells was found to decrease progressively in the course of HIV infection, resulting in a much lower percentage of IFN-γ-producing CD4$^+$ T cells. This suggests that EBV-specific CD8$^+$ T cells may be EBV-specific, of which 13% were shown to produce IFN-γ and function (using IFN-γ Elispot assay) of EBV-specific CD8$^+$ T lymphocytes as assessed by tetramer staining as a composite of all tested tetramers (solid line) and IFN-γ Elispot assay (dashed line), as described in "Patients, materials, and methods." (●) indicates tetramer; (○), IFN-γ. (C) Longitudinal analysis of EBV load, expressed as the number of virus copies per 10$^6$ PBMCs (solid line), in comparison with the number of functional EBV-specific CD8$^+$ T lymphocytes per 10$^6$ CD8$^+$ T cells (dashed line). (●) indicates EBV load; (○), IFN-γ.
AIDS-NHL patient (NHL0308, left panel) and one LTA individual (LTA0036, right panel). (A) Number of tetramer **+** (percentage of CD8 **+** T cells, solid line) and IFN-γ **+** T cells (percent 10 **6** CD8 **+** T cells, dashed line), as assessed by Elispot assay, are shown in the course of HIV-1 infection for RAK-specific T cells of one AIDS-NHL patient (NHL0308, left panel) and one LTA individual (LTA0036, right panel). (B) Intracellular IFN-γ staining, as described in “Patients, materials, and methods,” after stimulation with RAK peptide is shown at an early time point in HIV-1 infection for the AIDS-NHL patient and an early and late time point for the LTA individual, revealing the percentage of IFN-γ **+** producing CD8 **+** T cells (upper right quadrant). (C) Combination of tetramer and intracellular IFN-γ staining after peptide stimulation is shown at an early time point for the AIDS-NHL patient, revealing the percentage tetramer **+** IFN-γ **+** (2.3%, upper left quadrant), IFN-γ **+** tetramer (0.21%, upper and lower right quadrant) T cells, and proportion of IFN-γ **+** tetramer **+** CD8 **+** T cells (9%, gray box).

Figure 7. Correlation of IFN-γ **+** producing T cells by Elispot and intracellular FACS-staining. (A) Number of tetramer **+** (percentage of CD8 **+** T cells, solid line) and IFN-γ **+** T cells (percent 10 **6** CD8 **+** T cells, dashed line), as assessed by Elispot assay, are shown in the course of HIV-1 infection for RAK-specific T cells of one AIDS-NHL patient (NHL0308, left panel) and one LTA individual (LTA0036, right panel). (B) Intracellular IFN-γ staining, as described in “Patients, materials, and methods,” after stimulation with RAK peptide is shown at an early time point in HIV-1 infection for the AIDS-NHL patient and an early and late time point for the LTA individual, revealing the percentage of IFN-γ **+** producing CD8 **+** T cells (upper right quadrant). (C) Combination of tetramer and intracellular IFN-γ staining after peptide stimulation is shown at an early time point for the AIDS-NHL patient, revealing the percentage tetramer **+** IFN-γ **+** (2.3%, upper left quadrant), IFN-γ **+** tetramer (0.21%, upper and lower right quadrant) T cells, and proportion of IFN-γ **+** tetramer **+** CD8 **+** T cells (9%, gray box).

The observed low percentage of IFN-γ **+** producing T cells is not likely to be a consequence of susceptibility to rapid activation-induced cell death in vitro, caused by preactivation in vivo. Because tetramer staining results before and after stimulation are virtually the same (data not shown), this indicates that antigen-specific cells are not lost after stimulation. In addition, when dead cells are included in the analysis, no increase in the percentage of tetramer **+** T cells was observed, suggesting that even if cells are lost, this is not occurring selectively in the EBV-specific CD8 **+** T cells. Furthermore, we studied expression of CD69, which has been shown to be highly expressed on nonresponsive T cells in LCMV-infected CD4 **+** T cells. However, expression of CD69 on EBV-specific CD8 **+** T cells was low, reaching no higher levels than 5% of the tetramer **+** T cells (data not shown). Indeed, it has been shown that, compared with healthy individuals, a higher percentage of PBMCs from HIV-infected individuals undergo apoptosis after overnight stimulation. However, death of T cells was not correlated with CD4 **+** T-cell numbers or T-cell function and was not confined to expression of T-cell activation markers. Moreover, it has been shown that virus-specific CD8 **+** T cells are able to rapidly reinstate cytokine production after recent stimulation. Thus, even if the EBV-specific T cells are recently activated or preactivated in vivo, this should result in a rapid production of IFN-γ. Overall, these observations make considerable in vitro death of EBV-specific T cells unlikely.

The low percentage of IFN-γ **+** producing T cells was also not due to a suboptimal assay condition, because the Elispot assay was shown to detect virtually all antigen-specific T cells when a T-cell line was analyzed. Furthermore, the percentage IFN-γ **+** producing T cells as assessed by intracellular staining was similar to the percentage of IFN-γ producers found by Elispot assay in both healthy controls (data not shown) and patients (Figure 7).

The phenomenon of antigen-specific CD8 **+** T-cell dysfunction has also recently been shown for hepatitis C virus, during a period of acute infection, for tumor-specific T cells in melanoma patients, and for HIV-specific T cells. This state of dysfunction has been shown to occur both at the level of IFN-γ production and cytolytic activity. In addition, in healthy individuals there is a correlation between IFN-γ **+** producing T cells and CTLp frequencies. Furthermore, our own observations indicate that the number of CTLp **+** and IFN-γ **+** producing T cells correlate, and both decreased in the course of HIV infection in AIDS-NHL patients.

In AIDS-NHL patients the observed loss of function of EBV-specific CD8 **+** T cells was accompanied by an increase in EBV load, although absolute EBV load did not differ between groups (data not shown). This increase in EBV load cannot be attributed to technical variation, because we have low variation in duplicate measurements (7.5%) and correlation for the quantity of input DNA did not change the observed patterns. In addition, increases in EBV load are not due to increases in the number of total B cells, because correction for the number of B cells in PBMC samples did not lead to altered patterns (data not shown).

Thus, finally, in AIDS-NHL patients immune control over EBV
seemed to be lost. Surprisingly, in LTAs enormous transient bursts of EBV load were observed. These peaks in viral load seemed to be paralleled by expansions of functional CD8+ T cells specific for EBV. Because EBV load subsequently decreased, these cells apparently were able to control EBV viremia. Indeed, multivariate analysis of AIDS-OI patients and LTAs showed that EBV load, besides the number of CD4+ T cells, could predict the number of functional T cells, indicating that EBV load is indeed able to drive EBV-specific T cells when sufficient CD4+ T cells are present. In progressing to AIDS-OI, EBV-specific CD8+ T cells on total were lower, suggestive of physical loss or lack of expansion of these T cells. In these patients, the relatively low number of EBV-specific T cells did, however, not lead to an increase in EBV load, suggesting that there is adequate control. Alternatively, it could be that these individuals eventually would have developed AIDS-NHL had they not developed AIDS-OI.

In conclusion, our data suggest that the development of AIDS-NHL is a multifactorial process involving at least virologic and immunologic parameters. Thus, both determinants are required to obtain a complete picture of the virus-host balance. We show that not so much the total number of circulating EBV-specific CD8+ T cells but, mainly, the number of functional EBV-specific CD8+ T cells is important in keeping EBV infection under control. When EBV-specific CD8+ T cells start to lose function, in most cases as a consequence of a decrease in CD4+ T cells, this is paralleled by an increase in EBV load. To be able to predict the occurrence of an AIDS-NHL, all these factors should be taken into account.

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References

11. Shibata D, Weiss LM, Hernandez AM, Nathwani RA, Pagano JS. Epstein-Barr virus replication in oropharyngeal epithelial cells is important in keeping EBV infection under control. When EBV-specific CD8+ T cells start to lose function, in most cases as a consequence of a decrease in CD4+ T cells, this is paralleled by an increase in EBV load. To be able to predict the occurrence of an AIDS-NHL, all these factors should be taken into account.

27.卡西乌斯·特拉库斯 (Cassius Gallicus) 是罗马帝国的将军和政治家。他在公元前60年被提比略·凯撒（Tiberius Caesar）任命为高卢总督，并在那里建立了沃尔肯·加德（Wolvens Gallicum），这是罗马在高卢地区的一个重要据点。卡西乌斯·特拉库斯在公元前37年被处决，结束了他在高卢地区的统治。卡西乌斯·特拉库斯是罗马历史上的一个重要人物，他对罗马帝国的扩张和巩固做出了重要的贡献。他在政治上的才能和军事上的勇气使他成为了一位杰出的领袖。


