Associations of Simian Immunodeficiency Virus (SIV)-Specific Follicular CD8+ T Cells with Other Follicular T Cells Suggest Complex Contributions to SIV Viremia Control

Mohammad Arif Rahman, Katherine M. McKinnon, Tatiana S. Karpova, David A. Ball, David J. Venzon, Wenjin Fan, Guobin Kang, Qingsheng Li and Marjorie Robert-Guroff

*J Immunol* 2018; 200:2714-2726; Prepublished online 5 March 2018;
doi: 10.4049/jimmunol.1701403
http://www.jimmunol.org/content/200/8/2714

References  This article cites 62 articles, 26 of which you can access for free at:
http://www.jimmunol.org/content/200/8/2714.full#ref-list-1

**Why *The JI*?** Submit online.
- **Rapid Reviews! 30 days** from submission to initial decision
- **No Triage!** Every submission reviewed by practicing scientists
- **Fast Publication!** 4 weeks from acceptance to publication

*average

**Subscription**  Information about subscribing to *The Journal of Immunology* is online at:
http://jimmunol.org/subscription

**Permissions**  Submit copyright permission requests at:
http://www.aai.org/About/Publications/JI/copyright.html

**Email Alerts**  Receive free email-alerts when new articles cite this article. Sign up at:
http://jimmunol.org/alerts
Associations of Simian Immunodeficiency Virus (SIV)-Specific Follicular CD8+ T Cells with Other Follicular T Cells Suggest Complex Contributions to SIV Viremia Control

Mohammad Arif Rahman,* Katherine M. McKinnon,* Tatiana S. Karpova,† David A. Ball,‡ David J. Venzon,§ Wenjin Fan,§ Guobin Kang,§ Qingsheng Li,§ and Marjorie Robert-Guroff*  

Follicular CD8+ T (fCD8) cells reside within B cell follicles and are thought to be immune-privileged sites of HIV/SIV infection. We have observed comparable levels of fCD8 cells between chronically SIV-infected rhesus macaques with low viral loads (LVL) and high viral loads (HVL), raising the question concerning their contribution to viremia control. In this study, we sought to clarify the role of SIV-specific fCD8 cells in lymph nodes during the course of SIV infection in rhesus macaques. We observed that fCD8 cells, T follicular helper (Tfh) cells, and T follicular regulatory cells (Tfreg) were all elevated in chronic SIV infection. fCD8 cells of LVL animals tended to express more Gag-specific granzyme B and exhibited significantly greater killing than did HVL animals, and their cell frequencies were negatively correlated with viremia, suggesting a role in viremia control. Env- and Gag-specific IL-21+ Tfreg positively correlated with fCD8 cells in LVL animals and negatively correlated with viremia, suggesting a potential benefit of Tfreg via suppression of chronic inflammation. In contrast, in HVL macaques, Tfreg and fCD8 cell frequencies tended to be negatively correlated, and a positive correlation was seen between Tfreg number and viremia, suggesting possible dysfunction and suppression of an effective fCD8 cell immune response. Our data suggest that control of virus-infected cells in B cell follicles not only depends on fCD8 cell cytotoxicity but also on complex fCD8 cell associations with Tfh cells and Tfreg. The Journal of Immunology, 2018, 200: 2714–2726.

Among the earliest manifestations of the epidemic disease that came to be known as AIDS was persistent, generalized lymphadenopathy, first seen in homosexual males (1, 2). Lymph node (LN) follicles were subsequently identified as important sites of replication and trapping of the etiologic agent of the disease, HIV (3, 4), as well as of SIV in the rhesus macaque model (5). Subsequently, CD4+ Tfh cells in LN follicles, now known as T follicular helper (Tfh) cells (6, 7), were identified as key targets of both HIV (8–10) and SIV infection (11, 12) in secondary lymphoid tissue. During HIV infection, Tfh cells in B cell follicles produce HIV and are responsible for persistent virus transcription in long-term aviremic individuals treated with antiretroviral therapy (ART) (13). Significantly higher concentrations of SIV-producing cells have been reported to occur in B cell follicles compared with extrafollicular regions of the spleen, LN, and gut-associated lymphoid tissue of SIV-infected macaques during chronic asymptomatic infection (14). Furthermore, residual SIV infection has been localized in B cell follicles of rhesus macaques undergoing fully suppressive ART (15). Such observations have suggested that germinal center (GC) Tfh cells comprise an immune-privileged site for HIV/SIV replication (14, 16, 17), which may not be readily accessible to ART or to antiviral CD8+ T cells that lack expression of the follicular homing molecule, CXCRI5. Thus, the production of HIV/SIV in GC Tfh cells represents a major obstacle to obtaining a functional cure for HIV/SIV infection.

In HIV infection, CD8+ T cells, especially Gag-specific CTL (18, 19), play a role in control of viral load. Early studies showed that depletion of CD8+ T cells in SIV-infected animals impaired viremia control (20, 21). Furthermore, cytotoxic CD8+ T cells were detected in LN GC of HIV-infected individuals (22, 23), as well as in LNs of SIV-infected nonhuman primates (24, 25). However, LNs, among other tissues, have come to be considered sanctuaries where reservoirs of Tfh cells infected with HIV or SIV can persist (15, 26). The observation that tetramer+CD8+ T cells, although present in extrafollicular areas of LNs of HIV-infected subjects were mostly absent in follicles, provided a rationale for the persistence of HIV/SIV in lymphoid Tfh cells (16). The growing focus of the field on obtaining an HIV cure, requiring elimination of viral reservoirs, has stimulated new studies on quantitation and functional capability of CD8+ T cells in lymphoid follicles.

In healthy humans, a subset of CD8+ T cells was reported to use CXCR5 to enter B cell follicles (27). CXCR5+CD8+ T cells,
termed follicular cytotoxic T cells, were subsequently identified in the lymphoctic choriomeningitis virus (LCMV) mouse model and shown to enter B cell follicles and eradicate LCMV-infected Tfh cells and B cells infected with the herpesvirus, MuHV-4 (28). CXCR5+CD8+ T cells were also reported to express lower levels of inhibitory receptors than the CXCR5+ subset and exhibited more potent cytotoxicity against chronic LCMV infection (29). With regard to HIV/SIV infection, CD8+ T cells with follicular cytotoxic T cell characteristics, including potential cytotoxic function, have been identified in lymphoid follicles. CXCR5+ CD8+ T cells in individuals infected with HIV were shown to correlate inversely with viral load (29). Granzyme B+ CM-9-tetramer+ (SIV Gag-specific) CD8+ T cells in lymphoid follicles of mucosal tissue were enriched in chronically SIV-infected macaques that controlled infection compared with noncontrollers, and correlated inversely with lymphoid Tfh cells (30). Similarly, CXCR5+CD8+ T cells (termed follicular CD8+ T [fCD8] cells) were increased in LN follicles and GC of HIV-infected individuals, and they exhibited expression of both granzyme B and perforin. In the presence of anti-HIV/anti-CD3 bispecific Ab these cells exhibited better killing in vitro compared with non-fCD8 cells (31). Overall, these reports substantiate that fCD8 cells are located in B cell follicles of secondary lymphoid tissue, the site of active HIV/SIV replication, and that they might be key contributors to viremia control.

In this study, we sought to clarify the role of viral-specific fCD8 cells during the course of SIV infection in rhesus macaques. Although previous studies have suggested that fCD8 cells might contribute to viremia control in chronic viral infections, they are observed in chronically SIV-infected macaques exhibiting high viral loads (HVL). This raises the question of whether fCD8 cells in HVL animals are functional. Furthermore, the interactions of other cell populations in B cell follicles and their effect on fCD8 cells have not been fully explored. In this study, we show that in low viral load (LVL) animals, fCD8 cells are associated with viremia control whereas associations of Tfh cells with fCD8 cells suggest provision of help through secretion of IL-21. In contrast, in HVL animals Tfh cells apparently fail to provide adequate help, T follicular regulatory cells (Tfreg) are associated with diminished iCD8 cells and increased viremia, and fCD8 cells overall are not correlated with a protective outcome.

Materials and Methods
Study animals and sample collection
Rhesus macaques (27 females, 24 males) were maintained at Advanced Bioscience Laboratories (Rockville, MD) and at the National Cancer Institute animal facility (Bethesda, MD) under the guidelines of the Association for the Assessment and Accreditation of Laboratory Animal Care and according to the recommendations of the Guide for the Care and Use of Laboratory Animals. Protocols and procedures were approved by the Institutional Animal Care and Use Committee of the respective facility. Inguinal LN biopsy specimens were collected from: 1) 12 naive macaques (including preimmunization specimens from 4 of the acute group and 1 each from the LVL and HVL groups with no apparent clustering in the paired values); 2) 15 acutely infected macaques (samples obtained 24–40 wk after SIVmac239 infection); 3) 24 chronically infected macaques (samples obtained 40–50 wk after SIVmac239 infection). These were processed and stained with antibodies specific for SIV gp120 or gp160 as described (32), or with either ALVAC–SIV recombinant plus SIV gp120 protein or SIV DNA plasmid plus gp120 protein (T.A. Musich, S. Helmold, Z. Mushtaq, L. Miller-Novak, M.A. Rahman, T. Demberg, A. Valentin, D.A. Vargas-Inchaustegui, D.J. Venzon, G. Franchini, B.K. Felber, G.N. Pavlakis, and M. Robert-Guroff, manuscript in preparation). They were divided into LVL (11 macaques) and HVL (13 macaques) groups. One macaque each in the LVL and HVL groups had received empty Ad vector and adjuvant only. Samples from 11 of each group were used for flow cytometry, and 5 from each group were used in killing and ELISPOT assays. Median plasma viral loads during week 12 until necropsy for LVL and HVL animals were <3 × 106 and >7 × 106 SIV RNA copies per milliliter of plasma, respectively. Geometric means of the median values for the HVL and LVL macaques were 1.20 × 103 and 1.86 × 103, respectively. CD4 counts were not performed as routinely as the viral loads; however, median CD4 counts from the week 12 set point onward ranged from 641 to 923 for LVL macaques and from 248 to 552 for the HVL macaques. The means of the median CD4 counts for this time period were 805 and 384 for the LVL and HVL macaques, respectively. For LN viral loads, LN cells were lysed using QIAzol lysis reagent (Qiagen) and stored at −70°C until RNA extraction and viral load determination by droplet digital PCR as previously described (33). FACS analysis of the viral load (Fig. 1A–C), CD4 counts (Fig. 1D–F), and LN viral loads (Fig. 1G) of macaques in the LVL and HVL groups. LN viral loads were directly correlated with both the median plasma viral loads during chronic infection (Fig. 1H) as well as with plasma viral loads at the time of LN collection (Fig. 1I). This is consistent with a previous report that significantly correlated plasma viral loads with SIV follicular cells assessed by in situ hybridization (14). None of the macaques was devided with clinical signs of disease.

LN biopsies were minced and passed through a 40-μm cell strainer after lysis of RBCs. Cells were washed and resuspended in R10 (RPMI 1640 containing 10% FBS, 2 mM L-glutamine, 1% nonessential amino acids, 1% sodium pyruvate, and antibiotics). Cells were viable frozen in FBS plus 10% DMSO and thawed for all assays. Additional LN biopsies from three naive, five acute, four LVL, and five HVL animals were collected in OCT (Sakura Finetek, Torrance, CA) and frozen in liquid nitrogen for immunohistochemical analysis.

Flow cytometric detection of fCD8 cells, Tfh cells, and Tfreg
LN cells were stained with Aqua Live/Dead viability dye (Invitrogen, Carlsbad, CA) at room temperature (RT) for 15 min in PBS, washed with FACS wash, and surface stained with the following anti-human fluorochrome-conjugated mAbs cross-reactive with rhesus macaque Abs: Alexa Fluor 700 anti-CD4 (MQ1–17H12) (BD Biosciences), Brilliant Violet 711 anti-CD4 (L200), PE–Cy5 anti-CD95 (DX2) and allophycocyanin–eFluor 780 anti-CCR7 (3D12) from eBioscience (San Diego, CA); Brilliant Violet 786 anti-CD25 (BC96) and Brilliant Violet 605 anti–PD-1 (EH12.2H7) from BioLegend (San Diego, CA); and PE anti–CXCR5 (7D082.1) from the National Institutes of Health Nonhuman Primate Resource Center (Baton Rouge, LA). For detection of Tfreg, after surface staining cells were permeabilized with Foxp3/3transcription factor staining buffer (eBioscience) and stained intracellularly with PerCP-Cy5.5 anti–Foxp3 (236A/E7) (BD Biosciences). At least 500,000 singlet events were acquired on a SORP LSR II (BD Biosciences) and analyzed using FlowJo software (FlowJo, Ashland, OR). For all samples, gating was established using a combination of isotype and fluorescence-minus-one controls.

Cytokine staining of SIV-specific Tfh and fCD8 cells
SIV-specific production of granzyme B, perforin, and IL-21 by LN resident fCD8 and Tfh cells was assayed by stimulating LN cells in the presence of 2 μg/ml anti-CD49d and PE-Cy7 anti-CD28 (BD Biosciences, BD GolgiPlug, BD GolgiStop, and allophycocyanin–eFluor 780 anti–CCR7 (3D12) at the manufacturer’s recommended concentrations. One million cells were stimulated with pooled peptides spanning the SIVmac239 Gag (125 peptides) or Env (216 peptides) protein. Peptides were 15-mer peptides comprised by 9–11 residues, with an overlapping peptide of 14 residues each. Oligonucleotides were purchased from Health AIDS Research and Reference Reagent Program). Following incubation for 6 h at 37°C in 5% CO2, cells were stained with Live/Dead dye followed by surface staining and permeabilization with Fixp3 transcription factor staining buffer. Intracellular staining was performed with Alexa Fluor 647 anti–IL-21 (3A3–N2.1) (BD Biosciences), FITC anti–perforin (P9–344) (Mabtech, Cincinnati, OH), and PE–Texas Red anti–granzyme B (GB11) (Invitrogen/Life Technologies). Acquisition and analysis were as described above. SIVmac239 Gag- or Env–specific granzyme B, perforin, and IL–21 responses were calculated by subtracting the percentage stimulated response from the percentage peptide-stimulated response. Specific mean fluorescence intensity (MFI) was calculated as the peptide-stimulated response minus the unstimulated response.

Cell sorting
LN cells from acutely and chronically infected animals were stained with Brilliant Violet 711 anti-CD4 (L200), FITC anti–CD8 (RPA-T8) (BD Biosciences), Brilliant Violet 780 anti–CCR7 (3D12), and PE–Cy5 anti–CD95 (DX2) and allophycocyanin–eFluor 780 anti–CCR7 (3D12) at the manufacturer’s specified concentrations. One million cells were stimulated with pooled peptides spanning the SIVmac239 Gag (125 peptides) or Env (216 peptides) protein. Peptides were 15-mer oligonucleotides comprised by 9–11 residues, with an overlapping peptide of 14 residues each. Oligonucleotides were purchased from Health AIDS Research and Reference Reagent Program). Following incubation for 6 h at 37°C in 5% CO2, cells were stained with Live/Dead dye followed by surface staining and permeabilization with Fixp3 transcription factor staining buffer. Intracellular staining was performed with Alexa Fluor 647 anti–IL-21 (3A3–N2.1) (BD Biosciences), FITC anti–perforin (P9–344) (Mabtech, Cincinnati, OH), and PE–Texas Red anti–granzyme B (GB11) (Invitrogen/Life Technologies). Acquisition and analysis were as described above. SIVmac239 Gag- or Env–specific granzyme B, perforin, and IL–21 responses were calculated by subtracting the percentage stimulated response from the percentage peptide-stimulated response. Specific mean fluorescence intensity (MFI) was calculated as the peptide-stimulated response minus the unstimulated response.
**ELISPOT assay**

The ELISPOT assay was conducted using monkey IFN-γ ELISPOT® (HRP) kits. The 96-well polyvinylidene plates were precoated with anti-IFN-γ mAb (MT126). After washing and blocking the plates, CD8+ CXCR5+ or CD8+CXCR5* sorted LN cells were serially diluted and plated in duplicate at 1 × 10^5, 0.5 × 10^5, and 0.25 × 10^5 cells per well. Cells were stimulated by adding either SIVmac239 Env or Gag pooled peptides (1 μg/ml each peptide) and incubating for 20 h at 37°C. Subsequently, plates were washed with PBS before the addition of biotinylated anti-IFN-γ mAb (7-B6-1) at 1 μg/ml. Following incubation at RT for 120 min, plates were washed with PBS and incubated with the manufacturer’s recommended dilution of streptavidin-conjugated HRP for 60 min at RT. Following washes with PBS, spots resulting from individual IFN-γ-producing cells were visualized after a 15-min reaction with tetramethylbenzidine ready-to-use substrate solution, counted with an Eliphosto counter (CTL analyzer), and recorded per 10^6 CD8+ T cells. As negative controls, cells were plated without peptide. SIV-specific spots were calculated by subtracting spots resulting from unstimulated cells.

**Killing assay**

Cytotoxic activity of CD8+ T cells was assayed using a modified flow cytometry–based killing assay (34). Sorted autologous CD4+ T cells, used as targets, were fluorescently labeled according to the manufacturer’s recommendation with the CellTrace Violet dye kit (Thermo Fisher Scientific/Life Technologies, Grand Island, NY). The labeled CD4+ cells were washed and pulsed with SIVmac239 Gag pooled peptides (1 μg/ml each peptide) for an hour. A portion of the cells was not peptide pulsed and served as control. Target cells (with/without peptide pulsing) were washed twice and 10,000 cells per well were plated in R10 in U-bottom 96-well plates. Effector CD8+CXCR5+ or CD8+CXCR5* sorted LN cells were placed in different wells at 25:1 E:T ratios to a final volume of 200 μl. Plates were incubated at 37°C for 5 h. After incubation, cells were labeled by adding 0.25 μl of yellow Live/Dead viability dye (Invitrogen) in 100 μl of PBS per well. Plates were washed twice with PBS and cells were resuspended in 200 μl of PBS–parafomaldehyde solution, transferred to acquisition tubes, and read on an LSR II. Killing was measured by incorporation of the yellow Live/Dead viability dye in CellTrace Violet+ target cells. Specific killing was defined as percentage killing of peptide-pulsed targets minus percentage killing of targets without peptide pulsing.

**Immunohistochemical detection of GCs and GC-resident fCD8 and Tfh cells**

Adjacent LN sections were stained for Tfh and fCD8 cells, respectively. Briefly, 6-μm sections were cut, adhered to silanized slides, and pretreated in 1 mM EDTA (pH 8) in a Presto pressure cooker (National Presto Industries, Eau Claire, WI) at 121°C for 35 s to unmask Ags. Sections were blocked with normal horse serum and incubated overnight at 4°C with the following primary Abs: rabbit monoclonal anti-human CD4 (clone EPR6855, catalog no. ab133616, 1:200; Abcam) or mouse monoclonal anti-human CD8 (clone RPA-T8, catalog no. 557084, 1:200; BD Biosciences), goat anti-human PD-1 (AF1086, 1:40; R&D Systems), and mouse monoclonal anti-Ki67 (clone MIB1, catalog no. VP-K40; Vector Laboratories) for CD4 and PD-1 staining combination or rabbit monoclonal anti-human Ki67 (clone EPR3610, catalog no. ab92742, 1:200; Abcam) for CD4 and Ki67 staining combination. After washing in PBS, slides were incubated at RT for 2 h with the following secondary Abs for CD4 staining: AlexaFluor 594 donkey anti-rabbit IgG, AlexaFluor 488 donkey anti-goat IgG, and AlexaFluor 647 donkey anti-mouse IgG and counterstained with DAPI. For CD8 staining secondary Abs included AlexaFluor 647 donkey anti-mouse IgG, AlexaFluor 488 donkey anti-goat IgG and AlexaFluor 594 donkey anti-rabbit IgG with DAPI counterstaining. The secondary Abs (diluted 1:200) were from Life Technologies. After washing in PBS, slides were coverslipped and examined. Images were collected on an LSM 780 confocal microscope with Zen software (Carl Zeiss, Thornwood, NY) using Plan-apochromat X40, 1:40 numerical aperture objective and gallium arsenide red laser line. Panels of single focal plane images were acquired with pixel size of 415 or 519 nm at 12-bit image depth with averaging (2) and pixel dwell time of 1.58 μs in the following channels: magenta (excitation, 633 nm; emission, 650–758), red (excitation, 561 nm; emission, 570–694 nm), green (excitation, 488 nm; emission, 490–553 nm), and blue (excitation, 405 nm; emission, 410–470 nm). For publication, images were scaled to 8-bit RGB identically, with a linear LUT and exported in the TIFF format. Figures were made from TIFF images in Adobe Illustrator without any change in resolution. Scale bars are provided for each image.

**Quantification of GC-resident fCD8 and Tfh cells in fixed LNs**

Ki67 staining defined GCs, as both light and dark zones have a high cell proliferation rate (35). The total area of individual GCs was determined by analyzing the area of Ki67+ and quantification of fCD8 and CD4+ Tfh cells in GCs was performed as described (36). Cells expressing CD4 or CD8 were extracted from the entire confocal images by first using the interactive segmentation software ilastik (v1.1.5) (37) to separate cells from the background based on features defined by color/intensity, edges, and texture in all image channels. Although the ilastik segmentation efficiently delineated cells from the image background, groups of cells were not adequately separated from each other. Therefore, the segmentation mask, along with the raw confocal images, was imported into custom software written in MATLAB (R2015a; Mathworks, Natick, MA), with use of the image processing toolbox. The custom software performed morphological erosion on the segmentation mask to isolate individual cells, which were then each given a unique identification number, and then returned to their original size by use of morphological dilation. The boundary of the GC was then outlined manually based on the Ki67 and PD-1 expression, and the number of cells in each GC was counted. To aid comparisons between different conditions, the normalized cell count in each image was obtained by dividing the number of cells in the GC by the area (in square millimeters) of the GC.

**Statistical analysis**

The Mann–Whitney U test was used for comparisons between different groups of animals, and the Wilcoxon signed rank test was used for paired differences within the same group of animals. Correlations were assessed using the Spearman rank correlation test. Exact permutation p values were calculated. When small numbers of observed values yielded inadequate power for nonparametric tests, outcomes combined across groups were modeled using ANOVA after appropriate transformations to meet distributional assumptions. Numbers of cells in multiple GCs per animal were analyzed by repeated measures ANOVA, and p values for the multiple pairwise comparisons of the groups were corrected using the Tukey method. In all other figures, p values reported are not corrected for multiple comparisons. Analyses were performed using GraphPad Prism (GraphPad Software) and SAS/STAT software version 12.1 (SAS Institute, Cary, NC).

**Results**

CD8+ and CD4+ cells accumulate in LN follicular regions during the course of SIV infection

The number of CD8+ and CD4+ cells in LN GCs was determined immunohistochemically. Representative staining of LNs from acutely and chronically infected (LVL and HVL, viral loads shown in Fig. 1) and naive macaques is shown (Fig. 2). Eighteen GCs (7, 2, and 9, respectively) of three naive, 19 GCs (7, 8, 2, 1, and 1, respectively) of five acute, 19 GCs of four LVL (5, 6, 6, and 2, respectively) and 28 GCs (4, 5, 6, 6, and 7, respectively) of five HVL animals were analyzed. Some of the animals had less defined GCs in the tissue sections examined. We chose more defined areas for analysis, which resulted in lower GC counts in 1 of the naive and 4 of the 14 infected animals. A higher number of CD8+ cells accumulated in GCs of the acutely infected compared with naive animals, with even higher accumulation in the chronically infected animals. However, no difference in the number of CD8+ cells between LVL and HVL animals was observed (Fig. 3A). Compared to LN of naive macaques, more CD4+ cells were seen in acutely and chronically infected animals (Fig. 3B). A positive correlation was observed between CD8+ and CD4+ cells/mm² of tissue in the acute and LVL animals (Fig. 3C, 3D), suggesting that the interaction between CD8+ and CD4+ starts early after exposure and continues during chronic infection in LVL but not HVL animals. Thus, CD4+ cells might contribute to the proliferation of CD8+ cells in the acute and LVL animals.
SIV-specific fCD8 cells accumulate in B cell follicles of SIV-infected LNs

LNs from all macaque groups (see Materials and Methods) were further analyzed by flow cytometry to detect SIV-specific fCD8 cells during the course of infection. The relative expression of CCR7 and CXCR5 on CD8+ T cells determines their localization within LNs (38, 39). CD8+ T cells expressing high CXCR5 and low CCR7, defined as fCD8 cells, are localized in B cell follicles and were gated as shown (Fig. 4A). Increased frequencies of fCD8 cells were observed in LNs of acutely and chronically infected macaques, but no difference in the level of fCD8 cells was seen between LVL and HVL groups (Fig. 4B). As CD3 cells were not stained in the immunohistochemical analysis, we examined flow cytometry data from the same LNs to determine the fraction of cells that were CD3+ as well as both CD4+ and CD8+. The percentage of CD3+ T cells in the follicular CD4+ population ranged from 78.8 to 97.5%; the percentage in the follicular CD8+ population ranged from 70.9 to 98.9%. The percentage of CD4+CD8+ T cells among the total follicular CD4+ and CD8+ cells ranged from 0.3 to 4.5%. Overall, the percentage of fCD8 cells was highest in the chronic phase of infection regardless of viral load as also observed for CD8+ cells in GCs immunohistochemically (Fig. 3A).

To further investigate the expansion of fCD8 cells during the course of SIV infection, the frequency of SIV Env- and Gag-specific fCD8 cells was determined by ELISPOT assay in acutely and chronically infected animals. Cells of both specificities were observed in infected animals, with greater levels seen in chronically infected animals. However, no significant difference was obtained between LVL and HVL groups (Fig. 4C, 4D). To assess the potential impact of greater Gag-specific granzyme B fCD8 cell responses in LVL macaques on viremia control, correlation analyses were performed between fCD8 cells and median plasma viremia during chronic infection (week 12 to time of necropsy of HVL and LVL macaques and viremia at the time of necropsy of acutely infected macaques. A significant negative correlation was observed between fCD8 cells in B cell follicles and median viral load in LVL animals (Fig. 4J), but not in the other groups of animals (data not shown), suggesting that fCD8 cells might contribute to viremia control in LVL animals.

Killing capacity was next investigated as a mechanism by which fCD8 cells might contribute to viremia control. CD8+CXCR5+ and autologous CD4+ cells were sorted from LNs of acute, LVL, and HVL animals. CD4+ cells pulsed with SIVmac239 Gag peptides were used as targets and CD8+CXCR5+ fCD8 cells were used as effectors. fCD8 cells from LVL animals killed target cells more efficiently compared with those from HVL and acutely infected

FIGURE 1. Virological and immunological status of the study animals. (A and B) Viral load of HVL and LVL animals, respectively, during the course of infection. (C) Median viral load of the HVL and LVL groups after infection. (D and E) CD4 counts of HVL and LVL animals, respectively, after infection. (F) Median CD4 counts of the HVL and LVL groups after infection. (G) LN viral loads of the macaques. LN cells from 8 LVL and 11 HVL macaques were available for analysis. (H) Correlation between LN viral loads and median plasma viral load. (I) Correlation between LN viral loads and plasma viral loads at the time of LN collection. Viral loads at time of LN collection for two HVL macaques were not determined. Data of (G) were analyzed by the Mann-Whitney U test. Data of (H) and (I) were analyzed by the Spearman correlation test. ***p < 0.001.
CD8+ T cells control virus replication with the help of CD4+ T cells, which provide IL-21, essential for maintaining CD8+ T cell functionality (40, 41). The control might be compromised in the absence of CD4+ T cell help (42). An increase in Tfh cells in LNs occurs following both HIV and SIV infection (11, 43); however, IL-21 production by the cells has been reported to be significantly reduced (44). Thus, we investigated LN Tfh cell–fCD8 cell interactions. Tfh cells were gated as shown (Fig. 5A) and analyzed in the four macaque groups. Overall Tfh cell

animals (Fig. 4K), further supporting the notion that fCD8 cells contributed to control of viral load in the LVL animals.

**Tfh cells are associated with fCD8 activity in SIV-infected animals**

CD8+ T cells control virus replication with the help of CD4+ T cells, which provide IL-21, essential for maintaining CD8+ T cell functionality (40, 41). The control might be compromised in the absence of CD4+ T cell help (42). An increase in Tfh cells in LNs occurs following both HIV and SIV infection (11, 43); however, IL-21 production by the cells has been reported to be significantly reduced (44). Thus, we investigated LN Tfh cell–fCD8 cell interactions. Tfh cells were gated as shown (Fig. 5A) and analyzed in the four macaque groups. Overall Tfh cell
percentages were elevated in acutely and chronically SIV-infected animals compared with naive macaques; highest frequencies were observed in HVL animals (Fig. 5B). Comparable data were obtained on GC CD4+ cells immunohistochemically (Fig. 3B). Percentages of Env- (Fig. 5C) and Gag-specific IL-21+ Tfh cells (Fig. 5D) were highest in chronically infected macaques with no difference between HVL and LVL groups. IL-21 expression levels in response to both Env and Gag peptide stimulation, reported as MFI, although not significant tended to be higher in the LVL macaques compared with the HVL macaques (Fig. 5E, 5F). Notably in LVL macaques, but not in the other infected animals, percentages of Tfh and fCD8 cells were significantly correlated (Fig. 5G). Percentages of both Env- and Gag-specific IL-21+ Tfh cells in the LVL group, but not in the other groups, also correlated inversely with viral load (Fig. 5H, 5I). These correlations were lost in the chronic phase of infection, suggesting that Tfh play a positive role on Tfh cells in the early phase of infection.

We also investigated relationships between Tfh and viremia levels. A strong negative correlation between Tfh and chronic median viral load was observed in LVL animals (Fig. 6H). In view of the positive correlation between fCD8 cells and Tfh in LVL macaques (Fig. 6C), this result is consistent with Tfh contributing to viremia suppression. However, the percentage of Tfh in HVL macaques showed a positive trend with the median viral load (Fig. 6I), and the number of Tfh was also positively correlated with viremia (Fig. 6J). The trend of negative correlation between fCD8 and Tfh frequencies (Fig. 6D), together with the trend of positive correlation of Tfh frequency with viral load and the significant association of Tfh number with viral load in HVL macaques (Fig. 6F, 6J), suggests that Tfh contribute to the high viremia in these animals.

**Higher frequency and function of SIV-specific fCD8 cells compared with non-fCD8 cells**

As antiviral CD8+ T cells are important in the control of HIV/SIV replication and Tfh in B cell follicles are key sites of productive HIV/SIV infection (8–12, 52), we compared fCD8 cells within the B cell follicle with non-fCD8 (CD8+CXCR5+) T cells with regard to their impact on viral replication in vivo and viral load control. Higher frequencies of SIV Gag- and Env-specific fCD8 cells compared with non-fCD8 T cells were observed in SIV-infected macaques by ELISPOT assay (Fig. 7A, 7B). To investigate functionality, we examined frequencies of granzyme B+ and perforin+ cells in each compartment by flow cytometry. A previous study reported equivocal results in chronically SIV-infected macaques. Higher frequencies of granzyme B+ and perforin+ cells were found flow cytometrically in CXCR5+ CTL, including CCR7+ cells, indicating the fCD8 population, whereas perforin+ cells detected by in situ tetramer staining were more prevalent in the non-fCD8 cell population (14). In the present study, flow cytometric analysis revealed higher percentages of perforin+ SIV Gag- and Env-specific fCD8 cells compared with non-fCD8 cells in LNs of all SIV-infected macaques (Fig. 7C, 7D) together with
greater frequencies of granzyme B+ SIV Gag- and Env-specific fCD8 cells in LVL and HVL animals (Fig. 7E, 7F). The data suggest that SIV-specific CD8+ T cells accumulate during the course of infection in B cell follicles where they are available to combat virus-infected cells.

A caveat in this regard involves the PD-1/PD-L1 pathway, which contributes to the exhaustion of HIV-specific CD8+ T cells. In both HIV- (53) and SIV-specific CD8+ T cells (54), PD-1 is upregulated and positively correlated with high plasma viremia and negatively correlated with CD4+ T cell counts. Blocking of the PD-1/PD-L1 interaction in the SIV-macaque model has reversed exhaustion of Ag-specific CD8+ T cells (54). Moreover, inhibitory PD-1 has been associated with lower functionality of CD8+ T cells (55) and with decreased HIV-specific CD8+ T cell proliferation (56). Whereas PD-1+ T cells have been associated with poor functionality, these “exhausted” CD8+ T cells have also been shown to play a vital role in controlling viral replication (57). In this study, we assessed PD-1 expression on fCD8 cells at different stages of SIV infection. Elevated frequencies of PD-1+ fCD8 cells were seen in chronically infected animals, being highest in HVL animals (Fig. 8A). This result suggested that fCD8 cells of HVL macaques might have lower functionality compared with fCD8 cells of the other macaque groups. In fact, fCD8 cells of LVL macaques exhibited greater SIV-specific killing compared with both acutely infected and HVL macaques (Fig. 4K).

We further compared frequencies of PD-1+ fCD8 cells and non-fCD8 cells. In all macaque groups, greater percentages of PD-1+ cells were observed in the fCD8 populations (Fig. 8B). Killing assays performed using fCD8 or non-fCD8 cells as effectors with Gag peptide–pulsed autologous CD4+ T targets revealed a higher percentage of killing by the fCD8 cells compared with the non-fCD8 cells, regardless of the viral load or duration of infection (Fig. 8C), and despite displaying high PD-1 frequencies, as PD-1 expression has been reported to be unreliable as a marker for immune exhaustion in chronic SIV infection (58). Higher frequencies of SIV Env- and Gag-specific non-fCD8 cells were
observed in chronically infected animals compared with acutely infected ones (Fig. 8D, 8E), with no difference between LVL and HVL macaques and no difference in their killing capacity (Fig. 8F). Unlike fCD8 cells of LVL macaques, the frequency of non-fCD8 cells in all the LVL animals did not correlate negatively with viral load (data not shown). Therefore, killing attributed to non-fCD8 cells might not be a major contributing factor in viremia control of the LVL animals studied.

Recent data following in vivo CD8+ T cell depletion of SIV-infected macaques showed that whereas SIV+ cells increased in both follicular and extrafollicular areas, the increase was ~2-fold greater in the extrafollicular region (59). The authors concluded that both follicular and extrafollicular CD8+ T cells can suppress viral replication in vivo. Whether this difference from our findings results from the fact that our animals were vaccinated whereas the CD8-depleted macaques were not, or because the latter macaques had only been infected for 59 d whereas our macaques were studied after 40–50 wk of infection will require further investigation.

Discussion

Our investigation of three LN follicular T cell populations of thésus macaques revealed correlative interactions and associations with maintenance of low viremia during SIV chronic infection. As recently reported for HIV-infected patients (31) and SIV-infected macaques (30), fCD8 cells were elevated in this study in chronically infected macaques. Although similar fCD8 cell frequencies between LVL and HVL macaques and similar frequencies of SIV Gag- and Env-specific cells and granzyme B+ and perforin+ cells were seen, granzyme B expression tended to be greater in fCD8 cells of LVL macaques. These cells also exhibited significantly greater killing compared with HVL macaques (Fig. 4K). This greater functionality suggests a mechanism by which LVL macaques maintain viremia control, supported by the negative correlation of their fCD8 cell frequency and viral load (Fig. 4J). Connick et al. (14) in an earlier study reported using a mixed model analysis that SIV-specific CTL in both follicular and extrafollicular compartments of combined LN and spleen were

FIGURE 5. Association of Tfh and fCD8 cells in B cell follicles. (A) Representative gating of Tfh cells. (B) Frequency of Tfh cells among CD4+ cells in LN of the four macaque groups. (C and D) Percentage of IL-21+ Tfh cells in response to Gag (C) and Env (D) peptide stimulation. (E and F) IL-21 expression on Env- (E) and Gag-specific (F) Tfh cells determined by MFI in LVL and HVL animals. Frequencies and MFI were calculated as the peptide-stimulated response minus the unstimulated response. An MFI calculated as 0 was assigned a value 0.1 for plotting. (G) Correlation between Tfh and fCD8 cell frequencies among T cells (CD3+ CXCR5+) in B cell follicles of LVL macaques. (H and I) Correlation between Env- (H) and Gag-specific (I) IL-21+ Tfh cells and median chronic viral load of LVL animals. The data of (B)–(F) were analyzed by a Mann–Whitney U test and (G)–(I) were analyzed by a Spearman correlation test. Horizontal and vertical bars denote mean and SD. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.
inversely correlated with chronic tissue viral load of all chronically SIV-infected macaques. That study is not strictly comparable, however, to the present study. The animal populations were somewhat different, as the earlier study used macaques infected a median of 22 wk whereas our macaque samples were obtained 40–50 wk postinfection. More important differences include the fact that we correlated total fCD8 cells with plasma viral loads, whereas Connick et al. correlated dominant SIV-specific CTL in follicular and extrafollicular compartments with SIV RNA-positive cells, perhaps contributing to the different results.

Tfh cells in macaque LNs exhibited elevated frequencies associated with increasing viremia, as reported earlier (10, 12). Notably, however, Env-specific IL-21⁺ Tfh cells were significantly higher in LVL macaques and tended to be higher in HVL animals (Fig. 5C) whereas Gag-specific IL-21⁺ Tfh cells tended to be higher in both LVL and HVL animals compared with the acute group. Tfh and fCD8 cell frequencies were directly correlated in LVL (Fig. 5G) but not HVL macaques. Importantly, Env- and Gag-specific IL-21⁺ Tfh cells of LVL animals were significantly negatively correlated with viral loads (Fig. 5H, 5I). These results in LVL but not HVL animals.

We confirmed an increased abundance of Tfreg during the course of infection (50), although no difference was observed between LVL and HVL animals. Tfreg have shown negative effects on Tfh cells in HIV and SIV infection, including inhibition of proliferation, ICOS expression, and IL-4 and IL-21 production (60). Although a positive correlation between Tfreg and Tfh cells in acutely infected animals was seen (Fig. 6E), it was lost in chronic infection. A significant positive correlation between Tfreg and fCD8 cells in the LVL animals (Fig. 6C), as well as a negative correlation between Tfreg and viral load (Fig. 6H), suggested that Tfreg positively impact fCD8 cells, which in turn control viremia. Importantly, in chronic viral infections, Treg have been shown to suppress CD8⁺ T cell activity in a contact-dependent manner (61). Moreover, in situ staining of LN tissue sections for SIV-specific CD8⁺ T cells and Foxp3⁺ Tregs revealed significant numbers of these cells in direct contact (59), further suggesting cell interactions. Strikingly, however, a significant negative correlation was seen between Tfreg and ifCD8 cell frequencies in HVL macaques (Fig. 6D). Additionally, a strong positive trend was seen between Tfreg frequency in HVL animals and their viral loads (Fig. 6I), along with a significant positive correlation between Tfreg number and viral load (Fig. 6J). These data suggest suppression of ifCD8

**FIGURE 6.** Association of Tfreg with ifCD8 and Tfh cells in B cell follicles. (A) Representative gating of Tfreg. (B) Frequency of Tfreg among CD4⁺ cells in LN of four animal groups. (C and D) Correlation between Tfreg and ifCD8 cell frequencies of LVL (C) or HVL (D) animals. (E) Correlation between Tfreg and Tfh cell frequencies in acutely infected macaques. (F and G) Correlation between Tfreg number and Tfh cell number (F) or IL-21⁺ Tfh cell number (G) in acutely infected macaques. (H and I) Correlation between the frequencies of Tfreg in LVL (H) or HVL (I) macaques and chronic viral loads. (J) Correlation between Tfreg number and chronic viral load in HVL macaques. The percentages in (C)–(E), (H), and (I) were taken among T cells (CD3⁺CXCR5⁺) in B cell follicles of the macaques. Numbers of Tfreg, Tfh cells, and IL-21⁺ Tfh cells in (F), (G), and (J) were normalized to 1 million CD3⁺ cells. The data of (B) were analyzed by a Mann–Whitney U test; (C)–(J) were analyzed by a Spearman correlation test. Horizontal and vertical bars denote mean and SD. *p < 0.05, ***p < 0.001.
cells by Tfreg might contribute to the loss of viremia control in the HVL animals. Although we focused on CD4+ Tfreg in the present study, CD8+ Tfreg have been shown to make up most of the CD8 T cells within follicles of both humans and macaques (62) and to impair both Tfh and GC B cell responses. As they have been characterized as having limited cytotoxic potential, they could be critical in maintaining persistent HIV and SIV infection of lymphoid follicles.

Overall, these results illustrate the complexity of cellular activities within LN follicles, and they indicate that assessment of a single T cell population and associated molecular mechanisms will be insufficient to fully understand pathogen control. Detection of iCD8 cells in B cell follicles of secondary lymphoid organs in humans, as well as in murine and macaque models (14, 16, 17, 27–31), suggested that iCD8 cells enter B cell follicles and contribute to eradication of infected cells. Although HIV-specific killing in vitro was reported in the presence of bispecific Ab (31), to our knowledge, this is the first report showing SIV peptide–specific killing of target cells by iCD8 cells isolated from LN. Although iCD8 cells of LVL animals had higher killing capacity and were negatively correlated with viremia, iCD8 cells were also detected in HVL animals. Their higher expression of PD-1 is not necessarily firm evidence of immune exhaustion (58), and other mechanisms may contribute to their impaired viremia control. Although Tfh cell frequency was highest in HVL animals, these cells tended to have lower frequencies and/or expression of IL-21, suggesting dysfunction and diminished support of iCD8 cells. In fact, unlike LVL animals, the percentage of IL-21+ Tfh cells in HVL animals did not correlate with reduced viral load or iCD8 cell frequency. Additionally, a negative impact of Tfreg on iCD8 cells, suggested by the negative correlation in HVL macaques (Fig. 6D), is consistent with lower viral control by iCD8 cells in these animals.

The frequencies of SIV Env- and Gag-specific iCD8 cells compared with non-iCD8 cells were also detected in HVL animals. Their higher expression of PD-1 is not necessarily firm evidence of immune exhaustion (58), and other mechanisms may contribute to their impaired viremia control. Although Th cell frequency was highest in HVL animals, these cells tended to have lower frequencies and/or expression of IL-21, suggesting dysfunction and diminished support of iCD8 cells. In fact, unlike LVL animals, the percentage of IL-21+ Th cells in HVL animals did not correlate with reduced viral load or iCD8 cell frequency. Additionally, a negative impact of Tfreg on iCD8 cells, suggested by the negative correlation in HVL macaques (Fig. 6D), is consistent with lower viral control by iCD8 cells in these animals.

The frequencies of SIV Env- and Gag-specific iCD8 cells compared with non-iCD8 cells were also detected in LNs of infected animals (Fig. 7A, 7B). Furthermore, the frequencies of SIV-specific granzyme B+ and perforin+ cells were higher in iCD8 cells compared with non-iCD8 populations (Fig. 7C, 7D). These data explain the higher SIV-specific killing by iCD8 cells compared with non-iCD8 cells, and they are consistent with results obtained in the LCMV mouse model, where iCD8
cells exhibited more potent cytotoxicity than does the CXCR5− subset (29). SIV-specific killing in non-fCD8 cell populations was higher in the chronic phase compared with the acute phase, but no difference in killing was observed between LVL and HVL animals. Moreover, the frequency of SIV-specific non-fCD8 cells did not correlate with viral loads, suggesting that the activity of fCD8 cells might be more important for viremia control.

Overall, our results suggest that elimination of virus-infected cells in B cell follicles does not only depend on the cytotoxicity of fCD8 cells but also interaction of fCD8 cells with other follicular cells, including Tfh cells and Tfreg. Note that all but one in each group of the chronically infected HVL and LVL macaques in our study had been previously vaccinated, so our results cannot be generalized to what might occur in natural infection. Whether the vaccination regimen had long-term effects on follicular T cell populations in the macaques 40–50 wk postinfection remains to be determined. A recent abstract reported that a DNA/MVA SHIV vaccine regimen elicited SHIV-specific CXCR5+CD8+ T cells in the blood of rhesus macaques and were enhanced when the regimen was adjuvanted with CD40L (63). The frequency of these cells was inversely correlated with peak viral load following SHIV infection. Future studies will be critical in determining whether immunization can induce potent fCD8 cells leading to protection or enhanced control of viremia. In addition to prophylactic vaccines, further understanding of how to direct the differentiation and function of fCD8 and other follicular T cell populations might lead to the design of new strategies for eliminating HIV/SIV reservoirs.

Acknowledgments
We thank the staffs at the National Cancer Institute animal facility and Advanced BioScience Laboratories, Inc., and Tanya Hoang and Sophia Brown (Vaccine Branch, National Cancer Institute) for laboratory and flow cytometry support. PE anti-CXCR5 was obtained through the National Institutes of Health Nonhuman Primate Reagent Resource. The following reagents were obtained through the AIDS Research and Reference Reagent Program (Division of AIDS, National Institute of Allergy and Infectious Diseases, National Institutes of Health): SIVmac239 Env and SIVmac239 Gag peptides (complete sets).

Disclosures
The authors have no financial conflicts of interest.
Recognized by the monoclonal antibody T1A-1 are present in CD8+ lymphocytes in lymph nodes of human immunodeficiency virus-1-infected patients. Am. J. Pathol. 142: 1750–1758.


