In Vivo–activated CD4 T Cells Upregulate CXC Chemokine Receptor 5 and Reprogram Their Response to Lymphoid Chemokines

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Summary

Migration of antigen-activated CD4 T cells to B cell areas of lymphoid tissues is important for mounting T cell–dependent antibody responses. Here we show that CXC chemokine receptor (CXCR)5, the receptor for B lymphocyte chemoattractant (BLC), is upregulated on antigen-specific CD4 T cells in vivo when animals are immunized under conditions that promote T cell migration to follicles. In situ hybridization of secondary follicles for BLC showed high expression in mantle zones and low expression in germinal centers. When tested directly ex vivo, CXCR5 hi T cells exhibited a vigorous chemotactic response to BLC. At the same time, the CXCR5 hi cells showed reduced responsiveness to the T zone chemokines, Epstein-Barr virus-induced molecule 1 (EBI-1) ligand chemokine (ELC) and secondary lymphoid tissue chemokine (SLC). After adoptive transfer, CXCR5 hi CD4 T cells did not migrate to follicles, indicating that additional changes may occur after immunization that help direct T cells to follicles. To further explore whether T cells could acquire an intrinsic ability to migrate to follicles, CD42CD82 double negative (DN) T cells from MRL-lpr mice were studied. These T cells normally accumulate within follicles of MRL-lpr mice. Upon transfer to wild-type recipients, DN T cells migrated to follicle proximal regions in all secondary lymphoid tissues. Taken together, our findings indicate that reprogramming of responsiveness to constitutively expressed lymphoid tissue chemokines plays an important role in T cell migration to the B cell compartment of lymphoid tissues.

Key words: chemokine • CXCR 5 • ELC • follicle • T lymphocyte

Cell–dependent antibody responses require cognate interaction between antigen-presenting B cells and antigen-specific T cells (1, 2). Antigen-mediated cross-linking of the B cell receptor promotes B cell localization at the boundary between T zone and follicles (for a review, see reference 3), the location of the earliest encounters between antigen-bearing B cells and antigen-specific T cells (4–6). After receiving signals from activated CD4 T cells, some B cells undergo differentiation into plasmablasts, which migrate to the red pulp of spleen or medullary cords of LNs and become antibody-secreting plasma cells. Other activated B cells move into the follicle and form a germinal center (GC) (1). The GC is strongly dependent on help from antigen-specific T cells that must migrate into the follicle to support this response (7–12). Movement of antigen-specific T cells has been studied within lymphoid tissues using an adoptive T cell transfer model (13). When recipients of OVA-specific TCR-transgenic CD41 T cells are immunized with OVA in adjuvant, there is a short period of transgenic T cell proliferation in the T zone, and then large numbers of these cells are induced to migrate into follicles (13). T cell migration to follicles has also been studied in nontransgenic mice during the response to pigeon cytochrome c (PCC) (7, 9, 14). The predominant responding T cell population expresses TCR containing Vα11 and Vβ3 (15, 16). When mice are immunized with PCC in adjuvant, a protocol that induces a strong GC response, Vα11Vβ3-expressing T cells are found to begin localizing in follicles by day 5 and to peak in number at day 10, with many of the cells at this time being resident in GCs (7, 9, 10, 14, 17). In contrast to these findings, when mice are given antigen in saline intravenously, T cells become activated but fail to migrate into follicles. Many of the cells are quickly eliminated, and those...
that remain respond poorly to subsequent antigen exposure (13). Therefore, CD4 T cell migration into follicles has been suggested to be important for supporting not only B cell responses, but also a fully developed T cell response and induction of T cell memory.

The understanding of how antigen-specific T cells migrate to follicles is poorly developed. Recently, several chemokines have been characterized that are constitutively expressed in lymphoid tissues and help direct the movements of resting and activated lymphocytes. B lymphocyte chemo-attractant (BLC) (18), also called B cell-attracting chemokine 1 (BCA-1; 19)) is a CXC chemokine made by stromal cells in lymphoid follicles that functions as a ligand for CXCR5 chemokine receptor 5 (CXCR5, previously called Burkitt's lymphoma receptor [BLR1]) (20). Gene knockout studies have established that CXCR5 is required for B cell homing to follicles in spleen and Peyer's patches (21). In vitro, BLC was shown to be an efficacious attractant of B cells while attracting few T cells and no myeloid cells (18, 19). Two CC chemokines, secondary lymphoid tissue chemokine (SLC, also called 6-Ckine) and EBV-induced molecule 1 (EBI-1) ligand chemokine (ELC, also called macrophage inflammatory protein [MIP]-3β), are constitutively expressed by cells in the T zone (22, 23). SLC and ELC are ligands for CC chemokine receptor 7 and both strongly attract resting and in vitro-activated T cells (22-30). Mice that lack SLC expression (and have reduced ELC expression) have defective in vitro-activated T cells (22–30). Mice that lack SLC expression (and have reduced ELC expression) have defective in vitro-activated T cells (22–30). Mice that lack SLC expression (and have reduced ELC expression) have defective in vitro-activated T cells (22–30).

Mice. Six 10-wk-old BAb/cAnN mice were obtained from Charles River Laboratories. C57BL/6 (B6), MRL/MpJ (MRL), and MRL/Mp-lpr/lpr (MRL-lpr) mice were from The Jackson Laboratory. DO11.10 TCR-transgenic mice (32) on the BAb/c background, and aged and control B6 mice for memory T cell analysis were maintained in the University of California at San Francisco animal care facility. Eight 10-wk-old, specific pathogen-free, male B10.BR mice (The Jackson Laboratory) were housed under barrier conditions at the Duke University Vivarium.

Chemokines. H1S2-tagged murine BLC was prepared by PCR-based insertion of six histidine codons preceding the BLC stop codon. The BLC-his6 construct was inserted into the CMV-based mammalian expression vector pRK5 (33) and stably transfected into HEK-293 cells using the Lipotax mammalian transfection kit (Stratagene) according to the manufacturer's instructions. H1S2-BLC was purified from tissue culture supernatants using an NTA column (Qiagen). The protein was eluted in 100 μM imidazole (Fisher Scientific Co.). SDS-PAGE separation of the eluate revealed a band representing >90% of total protein corresponding to the recombinant H1S2-BLC. H1S2-ELC was produced in bacteria and purified as described (23). A similarly constructed vector for bacterial H1S2-SLC production was a gift from M. Gunn (Duke University). Stromal cell-derived factor (SDF)α (N33A) produced by chemical ligation (Gryphon Sciences), H1S2-BLC, -ELC, and -SLC were used in all chemokinesis assays, except for MRL-lpr chemokinesis assays, where, due to availability at the time of the experiments, non-H1S2-tagged murine SLC (gift of M. Gunn) and human ELC (R&D Systems) were used. We have not observed significant differences in the response of mouse cells to mouse or human ELC.

Materials and Methods

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was suspended in 2 μg/ml propidium iodide (for dead cell exclusion) in PBS with 5% FCS. The cells were analyzed using a dual laser modified FACStarPLUS® (Becton Dickinson Immunocytometry Systems; an argon laser as the primary, a tunable dye laser as the secondary) capable of seven-parameter collection. Files were acquired using CELLQuest™ software (Becton Dickinson) and analyzed using FlowJo software (Tree Star, Inc.).

Chemotaxis Assays. Chemotaxis assays were performed as described (23) using 10⁶ total cells per 5-μm transwell (Corning Costar Corp.). To identify migrating populations, a fraction of transmigrated cells was stained and analyzed by flow cytometry. Transmigrated LN cells from Balb/c recipients of OVA-specific T cells were stained with KJ1-26–FITC and anti-CD4–TRICOLOR (Caltag), or with anti-CD4–PE (Caltag) and KJ1-26–biotin followed by streptavidin-Cychrome (PharMingen). Because BLC causes reversible internalization of CXCR5 (36), transmigrated splenocytes from aged B6 mice were washed twice and incubated in RPMI plus 0.5% BSA for 1 h at 37°C, 5% CO₂ to allow CXCR5 reexpression before staining with anti-CXCR5 rabbit antiserum/goat anti-rabbit IgG/biotin/streptavidin–CyChrome–anti-CXCR5–FITC (Caltag), and anti-CD62L–PE (PharMingen). LN suspensions from 5–8-mo-old MRL-lpr mice were stained with anti-B220–PE, anti-Thy1.2–biotin (Caltag), and anti-CXCR5 rabbit antiserum followed by goat anti-rabbit IgG/FITC (Caltag) and streptavidin–CyChrome. To provide an internal control, MRL-lpr splenocytes (70% Thy1.2⁺, 9% Thy1.2⁻) were mixed 3:1 with B6 splenocytes (<1% Thy1.2⁺, 27% Thy1.2⁻) for chemotaxis assays. Transmigrated cells were stained with anti-Thy1.2–FITC (Caltag), anti-B220–PE, and anti-CD21–biotin/streptavidin–CyChrome–anti-CXCR5–FITC.

MRL-lpr DN T cell and aged B6 CXCR5hi T cell adoptive transfers. DN T cells were purified from LN s of 5–8-mo-old MRL-lpr mice. Total LN cells were incubated with biotinylated mAbs against CD22 (PharMingen), CD4, and CD8 (Caltag) followed by streptavidin–Cychrome–anti-CD22 (PharMingen). Memory phenotype splenocytes from 14-mo-old mice were enriched by MACS⁺ column (Miltenyi Biotec). Memory phenotype splenocytes from 14-mo-old mice were enriched by MACS⁺ depletion with biotinylated mAbs against CD8, B220, and CD11b (Caltag). For all transfers, 2 × 10⁶ cells were labeled with 5- and 6-carboxyfluorescein succinimidyl ester (CFSE; Molecular Probes) as described (35) and transferred by intravenous injection into appropriate syngeneic (MRL or B6) recipients. After 10 days, recipients were killed, and spleen, LNs, and Peyer’s patches were fixed and stained as described previously (35) with the following mAbs against CD22 (PharMingen), CD4, and CD8 (Caltag) followed by streptavidin–Cychrome. To provide an internal control, MRL-lpr splenocytes (70% Thy1.2⁺, 9% Thy1.2⁻) were mixed 3:1 with B6 splenocytes ( Thy1.2⁻, 27% Thy1.2⁻) for chemotaxis assays. Transmigrated cells were stained with anti-Thy1.2–FITC (Caltag), anti-B220–PE, and anti-CD21–biotin/streptavidin–CyChrome–anti-CXCR5–FITC.

Results

Immunization Conditions that Promote Antigen-specific CD4 T Cell Homing to Follicles Cause Rapid Upregulation of CXCR5 Expression. Mice that had received an inoculum of OVA-specific TCR-transgenic (DO11.10) T cells were injected with OVA323–399 peptide either subcutaneously in CFA to promote T cell trafficking to follicles, or intravenously in PBS to promote transient T cell activation without migration to follicles (13). At days 2, 3, 5, and 10 after immunization, LN s were isolated and analyzed by flow cytometry with a clonotypic antibody, KJ1-26, that recognizes the transferred OVA-specific T cells and with an antiserum specific for CXCR5 (35, 36). Before immunization, transferred OVA-specific T cells were uniformly CXCR5⁻ (Fig. 1 A). However, within 2 d of immunization with peptide in CFA, when OVA-specific T cell numbers in the draining LN s start to increase (13), a subpopulation of CXCR5⁺ cells could be identified (Fig. 1 B). By day 3, when the OVA-specific T cell frequency had increased ~30-fold, as in previous studies (13), most of the antigen-specific cells expressed high levels of CXCR5 (fluorescence intensity at least 10-fold greater than the staining control), and after 5 d the cells were uniformly CXCR5⁺ (Fig. 1 C, 1 C and D). Appearance of T cells in follicles followed similar kinetics to the CXCR5 upregulation (Fig. 2). Consistent with previous reports (13, 34), KJ1-26⁺ OVA-specific T cells began appearing in follicles by day 3 after immunization (Fig. 2 A) and reached maximal numbers by day 5 (Fig. 2 B). An enlargement of draining LN B cell areas occurred over this time period, and by day 5 many of the follicles contained nascent GCs (Fig. 2 B). Immunization of mice with OVA protein in LPS, a protocol that has been shown to promote T cell trafficking to follicles (34), also led to increased expression of CXCR5 on CD4 T cells (Fig. 1 F). In contrast to these effects, intravenous injection of OVA peptide in saline led to only weak induction of CXCR5 on OVA-specific T cells in LN s (Fig. 1 A–E) and spleen (data not shown), and did not promote KJ1-26⁺ T cell migration into follicles (Fig. 2 C). When mice immunized with OVA peptide in CFA were followed for longer times, a decline in CXCR5 expression was found to occur, although a significant proportion of KJ1-26⁺ cells remained CXCR5⁺ at day 10 after immunization (Fig. 1 E), and CXCR5⁺ cells could still be detected at day 25 (data not shown). By day 10, many B cell areas had become secondary follicles, comprising a well-developed GC and a sur-
CXCR5 upregulation by in vivo–activated OVA-specific T cells (A–E) Histograms depict CXCR5 expression on non-OVA-specific CD4 T cells (KJ1-26+) and on gated KJ1-26+B220+CD8+ cells from mice that were treated as follows (A) unimmunized; (B–E) immunized with OVA peptide subcutaneously in CFA (sc. KJ1-26+) or intravenously in saline (i.v. KJ1-26+); (F) immunized with OVA protein in LPS. In A, the level of CXCR5 expression on B cells is shown for comparison (dashed line). In B–F, the number of days elapsed between immunization and analysis is displayed in each panel. No Ag, no antigen injected. The experiment shown in F was performed at a different time from A–E, and has a different level of background staining. Results are representative of at least three independent experiments at each time point except panel F, which is representative of two experiments.

Figure 1. CXCR5 upregulation by in vivo–activated OVA-specific T cells (A–E) Histograms depict CXCR5 expression on non-OVA-specific CD4 T cells (KJ1-26+) and on gated KJ1-26+B220+CD8+ cells from mice that were treated as follows (A) unimmunized; (B–E) immunized with OVA peptide subcutaneously in CFA (sc. KJ1-26+) or intravenously in saline (i.v. KJ1-26+); (F) immunized with OVA protein in LPS. In A, the level of CXCR5 expression on B cells is shown for comparison (dashed line). In B–F, the number of days elapsed between immunization and analysis is displayed in each panel. No Ag, no antigen injected. The experiment shown in F was performed at a different time from A–E, and has a different level of background staining. Results are representative of at least three independent experiments at each time point except panel F, which is representative of two experiments.

CXCR5 upregulation on Vα11Vβ3-expressing cells during the PCC response. To test whether CXCR5 was upregulated on nontransgenic T cells under conditions not involving T cell transfer, the phenotype of PCC-specific Vα11Vβ3-expressing CD4 T cells was followed in mice immunized subcutaneously with PCC protein in adjuvant. Vα11Vβ3-expressing T cells responding to PCC were detected in immunized, but not unimmunized, animals by upregulation of CXCR5 on T cells after injection of antigen in adjuvant (Fig. 1 C). These kinetics of CXCR5 expression are in close accord with the kinetics of Vα11Vβ3-expressing T cell accumulation in follicular mantle zones and GCs during the response to PCC (9, 17). A subpopulation of the responding CD4+Vα11Vβ3-expressing T cells did not upregulate CXCR5 (Fig. 3). Such bimodality was not observed in the response of the monoclonal D011.10 T cells to OVA peptide (Fig. 1 D) and may indicate that T cells with differing affinity for peptide/MHC differ in their propensity to upregulate CXCR5.

Increased responsiveness of CXCR5-expressing T cells to the Follicular Chemokine BLC. The rapid upregulation of CXCR5 on T cells after injection of antigen in adjuvant and the migration of a fraction of the cells into follicles suggested that these cells had acquired responsiveness to BLC. This was tested directly ex vivo in transwell migration assays. OVA-specific T cells from draining LN s of adoptive transfer recipients immunized 7 d earlier with OVA peptide subcutaneously in CFA showed that a strong dose-dependent response to BLC (Fig. 4 A). By contrast, OVA-specific cells from recipients given the antigen intravenously in the absence of adjuvant did not respond to BLC (Fig. 4 A). Kinetic analysis using cells from mice immunized with OVA peptide in CFA showed the BLC response was detectable by day 2 and was maximal between days 5 and 10 (Fig. 4 B). In addition, flow cytometric analysis of transmigrated OVA-specific cells on day 3 revealed that while most input cells expressed CXCR5 (Fig. 1 C), there was an enrichment for cells expressing the highest levels of CXCR5 in the responding population (Fig. 4 C). These findings provide evidence for a direct relationship between CXCR5 expression on T cells and acquisition of BLC responsiveness, although they do not establish whether the activation state of the cells also affects their ability to respond. To investigate the BLC responsiveness of CXCR5-expressing T cells with a resting phenotype, we took advantage of previous observations that a subset of memory CD4+ T cells expresses CXCR5 (20, 21). CD4 T cells from aged (>1 yr) mice were characterized as a source of memory phenotype T cells, and a great majority of the L-selectinlo CD4+ T cells from these animals were found to express CXCR5 (Fig. 4 D). In addition to low L-selectin expression, the majority of CXCR5hi T cells in young and old mice expressed high levels of CD44 and reduced amounts of CD45R B (data not shown). Most of the CXCR5hi cells were also negative for the activation markers CD69 and CD25, further supporting their designation as memory cells. Interestingly, though a significant proportion of memory phenotype T cells in young mice expresses CXCR5, this proportion was consistently increased in aged mice (Fig. 4 D). In in vitro chemotaxis assays, the CXCR5hi memory T cells showed a very similar dose-sensitive BLC response to the activated OVA-specific T cells characterized above (Fig. 4 E). These results establish that both resting and activated CXCR5hi T cells respond to BLC. Interestingly, although the magnitude of the T cell response to BLC was lower than that observed for B cells, the T cells responded maximally to lower concentrations of BLC than did B cells (Fig. 4 E). This finding is similar to that made
previously with transfected Jurkat T cells (18), suggesting that T cells are intrinsically more sensitive than recirculating B cells to CXCR5 signaling. CD4 T cells express severalfold less surface CXCR5 than B cells (Fig. 1), demonstrating that higher surface chemokine receptor expression does not equate to higher chemokine sensitivity.

CXCR5-expressing T cells have reduced responsiveness to the T zone chemokines ELC and SLC. T cells have been shown to respond strongly to ELC and SLC in in vitro chemotaxis assays (22–24, 27–30). Since these chemokines are expressed in the T zone and might be able to counteract the ability of a cell to respond to a chemokine made in follicles, we tested whether CXCR5-expressing T cells were altered in their responsiveness to ELC and SLC. In striking contrast to the elevated ELC and SLC responsiveness of in vitro-activated T cells (28–30), OVA-specific T cells activated in vivo by subcutaneous peptide/CFA injection showed a significant downregulation in responsiveness to these T zone chemokines (Fig. 5 A). The time course of this decreased responsiveness to ELC (Fig. 5 B) was similar to the time course over which the cells became responsive to BLC (Fig. 4 B). In contrast, OVA-specific T cells from mice immunized intravenously with peptide in saline maintained their ability to respond to ELC throughout the 10-d time course (Fig. 5 B). Responsiveness to SDF1 was also decreased in the in vivo-activated T cells (Fig. 5 A), consistent with the recent in vitro finding that anti-CD3 stimulation reduces responsiveness to SDF1 (38). CXCR5-expressing T cells have reduced responsiveness to the T zone chemokines ELC and SLC.

Figure 2. Follicular homing of in vivo-activated OVA-specific T cells and expression pattern of BLC in secondary follicles. (A–C) Staining of LN sections to detect KJ1-26+ OVA-specific T cells in red and CD4 and CD8 in brown. (D) Staining with peanut agglutinin (PNA) to detect GCs in red, and with KJ1-26 to detect OVA-specific T cells in brown. Sections are from draining LNs of mice that received OVA-specific CD4 T cells and were immunized with OVA peptide subcutaneously in CFA 3 (A), 5 (B), or 10 (D) d before isolation, or intravenously in saline (C) 3 d before isolation. (E and F) In situ hybridization analysis of BLC expression pattern in LN sections containing secondary follicles. BLC hybridization is seen as dark blue staining, and sections are costained in brown for B220 (E) or PNA (F). The capsular staining in E was also seen in controls and is nonspecific. FO, follicle; T, T cell zone; FM, follicular mantle region. Original magnifications: A, C, E, and F, ×10; B and D, ×5.
Chemokine Responsiveness of Follicle-homing T Cells

Figure 3. Antigen-specific CD4 T cell expression of CXCR5 during a primary immune response. (A) Representative probability contours for CD44 and CXCR5 expression on Vα11Vβ3-expressing T cells over the course of a primary response. All profiles are presented as 5% probability contours with outliers, and are propidium iodide^2CD8^2B220^2CD11b^2Vα11^1Vβ3^1. The day after antigen administration is displayed above each panel. The quadrants are defined by the horizontal dotted lines, and indicate the limits of CD44 and CXCR5 regulation that were used to calculate the frequencies of cellular subsets. (B) Frequencies of Vα11Vβ3-expressing T cells that have upregulated CD44 and are either CXCR5^2 (left panel) or CXCR5^1 (right panel). (C) The total number of antigen-activated Vα11Vβ3-expressing T cells in the draining LNs as calculated using the frequencies obtained by flow cytometry and total cell counts estimated when organs were harvested; the left panel represents the CXCR5^2 subset, the right panel represents the CXCR5^1 subset. Each estimation (B and C) is presented as the mean from at least three separate animals ± SEM.

memory phenotype cells from aged mice showed similarly low responsiveness to ELC and SLC (Fig. 5 C). Interestingly, in these cells the responsiveness to SDF1 was elevated (Fig. 5 C).

T Cells Can Acquire the Intrinsic Ability to Migrate to Regions of Lymphoid Tissues Proximal to Follicles. The studies above demonstrate a tight relationship between CD4 T cell upregulation of CXCR5 expression, acquisition of BLC responsiveness, and migration into lymphoid follicles during an immune response. However, they do not establish whether intrinsic changes in the T cell are sufficient to direct these cells to follicles or whether additional (extrinsic) changes that accompany the adjuvant-induced immune response are also needed. Although many of the CD4 T cells in aged mice express CXCR5 (see Fig. 4 C), immuno-histochemical analysis did not reveal a significantly greater number of T cells in follicles in aged mice compared with young mice (data not shown). When CXCR5^hi CD4 cells were transferred from aged to young mice, they were found to localize within the T zone, with only occasional cells migrating into follicles (Fig. 6 A). These findings indicate that expression of CXCR5 is not sufficient to direct all types of T cells into B cell follicles. However, in contrast to aged normal mice, aged MRL-lpr mice contain very large numbers of T cells in a follicular distribution (39, 40). These CD3^+ T cells are unusual in lacking CD4 and CD8 and in expressing B220 (39, 40). Flow cytometric analysis showed that they also express high surface CXCR5 (Fig. 7 A), and in in vitro chemotaxis assays they demonstrated a robust response to BLC (Fig. 7 B) and a reduced response to ELC and SLC (Fig. 7 C). Importantly, the BLC dose–response curve of the DN T cells was typical of CXCR5^1 CD4 T cells (compare Fig. 7 B and Fig. 4, A and D) and not of B cells (see Fig. 4 D). Although the follicular location of the DN T cells in MRL-lpr mice suggests these cells have acquired the intrinsic ability to migrate to follicles, MRL-lpr mice have multiple immunological abnormalities, and it was possible that homing of DN T cells to follicles was dependent on extrinsic changes in the lymphoid tissues. To test this directly, DN T cells were purified from LN s of MRL-lpr mice, labeled with CFSE, and transferred to normal syngeneic MRL mice. Strikingly, the transferred T cells migrated to regions proximal to B cell areas in all secondary lymphoid tissues of the recipient mice (Fig. 6). Differences in the distribution of the cells were noted in the different tissues. In the spleen, the T cells homed to the outer rim of the follicles, especially near the marginal zone bridging channels, and often the cells appeared in contact with marginal metallophilic macrophages (Fig. 6 B). In LN s the T cells homed to perifollicular and interfollicular locations (Fig. 6 C), and in Peyer's patches the cells were seen to circle the whole follicular area (Fig. 6 D). These observations demonstrate that T cells can acquire the intrinsic ability to migrate to the boundaries of lymphoid follicles.
Discussion

The findings above establish that immunization with antigen in adjuvant causes antigen-specific T cells to upregulate CXCR5 expression and acquire responsiveness to the follicular chemokine, BLC, while simultaneously becoming less responsive to the T zone chemokines, ELC and SLC. We propose that this reprogramming of responsiveness to B and T zone chemokines is part of the mechanism by which antigen-activated T cells home to follicles to help initiate T-dependent antibody responses.

In the adoptive transfer studies of Jenkins and co-workers, it was observed that antigen needed to be injected in adjuvant for activated T cells to migrate to follicles (13, 34). When antigen was injected in the absence of adjuvant, T cell activation was transient and the activated cells failed to home to follicles. Our results provide a basis for understanding the different trafficking patterns of the activated cells as they show that CXCR5 upregulation and acquisition of BLC responsiveness only occurs after injection of antigen in adjuvant.

Similar results, with progressive accumulation of L-selectinlo and CXCR5hi T cells, were obtained for more than 10 animals <3 or >12 mo of age. (E) Chemotactic response of CXCR5lo/2 (white bars) and CXCR5hi (black bars) CD41 T cells, and of B cells (black bars) from the spleen of a 21-mo-old mouse. The y-axis on the left refers to T cells, and on the right to B cells. Results are representative of eight independent experiments.

Figure 4. BLC chemotactic response of in vivo-activated OVA-specific T cells and memory phenotype CXCR5-expressing T cells. Results are expressed as percentage of transmigrated input cells. Bars represent means of duplicate transwells. (A) Chemotactic response of KJ1-26+ CD41 cells from draining LNs of Balb/c recipients of OVA-specific CD41 T cells immunized 7 d before analysis with OVA peptide subcutaneously in CFA (●) or intravenously in saline (▲). R esults were similar at day 5 after immunization and are representative of five independent experiments. (B) Kinetic analysis of OVA-specific T cell acquisition of BLC responsiveness. The day after immunization with OVA peptide subcutaneously in CFA is indicated in each panel. Differences in basal migration levels were not reproducible and reflect assay to assay variability. Results are representative of at least two independent experiments at each time point. (C) CXCR5 expression on KJ1-26+ CD41 cells from day 3 draining LNs showing the total input population (Input) and the cells that migrated to BLC (Migrated). (D) CXCR5 and L-selectin (CD62L) expression on CD41 splenocytes from young (left panel) and 21-mo-old (right panel) B6 mice. Numbers represent the percentage of CD41+ and lymphocyte size-gated cells in each quadrant.

Figure 5. Chemotactic response profiles of in vivo-activated OVA-specific T cells and memory phenotype CXCR5-expressing T cells. Results are expressed as percentage of transmigrated input cells for each chemokine. (A) Chemotaxis of KJ1-26+ OVA-specific (black bars) and nonspecific CD41 (white bars) cells to 2.0 μg/ml LEC. Cells are from draining LNs of transfer recipients immunized with OVA peptide subcutaneously in CFA (solid bars) or intravenously in saline (hatched bars). The day after immunization is indicated on the x-axis. Data at day 5 are representative of five experiments for subcutaneous immunization in saline and of six experiments for intravenous immunization. (B) Response of KJ1-26+ OVA-specific (black bars) and nonspecific CD41 (white bars) cells to 0.2 μg/ml LEC. Cells are from draining LNs of transfer recipients immunized 7 d previously with OVA peptide subcutaneously in CFA. Chemokine concentrations were: BLC, 2 μg/ml; ELC, 0.2 μg/ml; SLC, 0.2 μg/ml; and SDF1, 0.3 μg/ml. Results were similar at day 5 after immunization and are representative of at least three independent experiments for each chemokine. (C) CXCR5 expression on KJ1-26+ cells from day 3 draining LNs showing the total input population (Input) and the cells that migrated to BLC (Migrated). Results are representative of five independent experiments. (D) Chemotaxis of CXCR5lo/2 (white bars) and CXCR5hi (black bars) CD41+ T cells from the spleen of a 21-mo-old mouse. Chemokine concentrations were: BLC, 2 μg/ml; ELC, 0.5 μg/ml; SLC, 0.8 μg/ml; and SDF1, 0.3 μg/ml. Results are representative of at least four independent experiments for each chemokine. CXCR5hi T cells exhibited reduced responsiveness to ELC (0.02–1.5 μg/ml) and SLC (0.08–1.2 μg/ml) at all chemokine concentrations tested.
antigen in adjuvant. Many studies have indicated that the effectiveness of adjuvants is through their potent activation of dendritic cells (DCs [41]), and it is therefore reasonable to suggest that effective induction of CXCR5 expression on T cells requires interaction with appropriately activated antigen-presenting DCs within the lymphoid tissue. OX40L is expressed by a subset of activated DCs (42), and recent studies by Lane and co-workers provide evidence that stimulation of T cells through OX40 can promote upregulation of CXCR5 (43) and homing of T cells to follicles (44). Further studies are needed to define whether additional costimulatory molecules can regulate CXCR5 expression on T cells.

In vivo activation by antigen in adjuvant decreases T cell responsiveness to ELC and SLC at the same time as increasing responsiveness to BLC. This contrasts with findings in vitro, where PHA- and IL-2-activated T cells responded more strongly than unactivated cells to SLC and ELC (28–30) and again indicates that the mode of T cell activation can strongly influence chemokine responsiveness. Recently, it has been shown that plt/plt mice, which exhibit defective homing of T cells to splenic T zones and LN s (45), have a compound defect that causes a deficiency in SLC expression and markedly reduced ELC expression (31). This finding provided strong evidence that SLC and ELC are needed for T cell homing to lymphoid T cell areas. Therefore, reduced responsiveness of CXCR5hi T cells to ELC and SLC may allow the cells to more readily leave the T zone and enter follicles. Reciprocally, the failure of T cells activated after intravenous injection of antigen to downregulate their SLC and ELC response might contribute to their inability to migrate to follicles. Since SLC appears important for cells to enter LN s via high endothelial venules or lymphatics (22, 31, 46), decreased responsiveness to this chemokine is also likely to influence the recirculation pattern of the cells.

**Figure 6.** Transferred MRL-lpr DN T cells, but not memory phenotype CXCR5hi T cells, home towards B cell follicles in secondary lymphoid organs of uninmunized nonautoimmune mice. Spleen (A and B), LN (C), and Peyer’s patch (PP; D) sections from syngeneic recipients of CFSE-labeled (A) CD4+CXCR5hi enriched splenocytes from 12–15-mo-old mice or (B–D) purified DN T cells from 8-mo-old MRL-lpr mice. Recipient tissues were isolated 1 d after cell transfer. CFSE-labeled cells are green. T cells are stained in red using mAbs to CD3e and Thy1.2 (A and B), CD8 (C), or CD4 (D). MOMA-1+ marginal zone metallophilic macrophages are stained blue. A is representative of five and B–D of three mice. F, follicle; T, T cell zone; MZ, marginal zone. Original magnification: ×10.
Our studies provide strong evidence that altered responsiveness to constitutively expressed chemokines is part of the mechanism by which antigen-activated CD4 T cells migrate towards and into B cell follicles. This conclusion is also supported by the transfer experiments showing that DN T cells from MRL-lpr mice are intrinsically capable of migrating to areas proximal to follicles (Fig. 6). However, the failure of CXCR5-high CD4 T cells to migrate to follicles after adoptive transfer suggests that additional factors might normally help guide antigen-activated CD4 T cells. The migration of only a subset of OVA-activated T cells to follicles also suggests that CXCR5-high cells may be heterogeneous in their responsiveness to these factors. Several studies have shown that B cell receptor-stimulated B cells upregulate expression of chemokines, including MIP-1α and MIP-1β (47, 48), and macrophage-derived chemokine (MDC) (49), that can attract subsets of activated T cells (50-53). Since antigen-activated B cells migrate to the boundary of B and T zones (3), it is likely that chemokines produced by activated B cells work together with constitutively expressed chemokines to bring antigen-activated T cells to the outer T zone. Whether further cues are needed to direct cells from the outer T zone into follicles remains unclear, although the failure of MRL-lpr DN T cells to migrate to the inner regions of follicles suggests this is the case. Possibilities include changes in the responsiveness of the T cells to unknown chemokines, and changes in the relative adhesiveness of the cells for features of the B or T cell compartments. Several examples of cell sorting occurring as a result of differential adhesiveness of cells have been reported (54-56).

A major role of T cells inside follicles is to support the GC response (1, 2). 9 d after immunization with PCC in adjuvant, a large majority of the PCC-responsive V<sub>α</sub>11Vβ3-expressing T cells express CXCR5 (Fig. 3), and at this time point, ~75% of the cells are localized within GCs (17). Therefore, at least a subset of PCC-responsive CXCR5-high T cells acquires the ability to enter GCs. Similarly, in the DO11.10 adoptive transfer system, 10 d after immunization with OVA peptide in CFA, the majority of KJ1-26<sup>+</sup> T cells were CXCR5-high, and many cells were found in follicular mantle zones and GCs (2 Fig. 2 D). The strong expression of BLC in primary follicles (18) and in follicular mantle zones of secondary follicles (Fig. 2, E, and F) is consistent with BLC playing a role in attracting T cells to these sites. The presence of only small numbers of BLC-expressing cells within GCs, predominantly in the region distal to the T zone (most likely corresponding to the GC light zone [1]) suggests that while BLC/CXCR5 might have a role in helping position cells within GCs, additional cues are likely to be needed. These findings also establish that there is substantial heterogeneity among follicular stromal cells in terms of BLC expression levels, with GC follicular DCs expressing relatively little of this chemokine. Perhaps by being concentrated predominantly outside the T zone-distal pole of the GC, BLC helps polarize the GC light and dark zone compartments. The notion that cues other than BLC play important roles in GC organization is supported by the finding that GCs are able to form in CXCR5-deficient mice (21). Furthermore, although spleens of CXCR5-deficient mice lack polarized follicles and contain aberrantly located GCs, the follicular disruption in LN s appeared to be minimal (21). Although this suggests that the role of CXCR5/BLC in B and T cell homing to LN s is redundant to other chemokine/receptor systems, studies in mice lacking lymphotoxin or TNF have shown that effects on follicular organization in LN s are more difficult to detect than in spleen or Peyer’s patches (57, 58), yet these effects can still be substantial (59, 60). As we have shown here, BLC is expressed in LN follicles, and CXCR5 is strongly upregulated on activated T cells in LN s. Future studies of BLC-deficient mice should help further dissect the contribution of BLC and CXCR5 to follicular organization and GC formation in LN s.

T cell homing to follicles may be important not only for providing help to B cells, but also for providing activated T cells with growth and survival signals. This possibility is suggested by the finding that antigen injected in the absence of adjuvant fails to promote T cell migration to follicles, and also fails to promote survival of activated or memory T cells (13, 34). The selective accumulation of the CXCR5-expressing subset of CD4 T cells in aged mice (Fig. 4 C) and in HIV-infected humans during disease progression (61) is consistent with the notion that trafficking through B cell areas plays a role in long-term survival of memory T cells. This possibility is also supported by the finding of a defect in CD4 T cell memory in B cell-deficient mice (62). Although we did not observe trafficking of CXCR5-high memory T cells to follicles in short-term transfer experiments,
studies in rats have suggested that memory T cells migrate through follicles at greater frequency than naive T cells (63). In summary, our findings suggest a model for how helper T cells home to follicles. After engagement of peptide/MHC complexes on appropriately activated T zone DCs, CD4 T cells upregulate CXCR5, acquire responsiveness to the follicular chemokine BLC, and simultaneously downregulate responsiveness to the T zone chemokines ELC and SLC. Together with additional presently undefined changes, this reprogrammed chemokine responsiveness helps propel T cells toward B cell areas. Further cues, such as those emanating from activated B cells and from GC cells, can then act upon these T cells to more precisely control their positioning and facilitate their ability to act as B cell helpers.

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