Conditional Vascular Cell Adhesion Molecule 1 Deletion in Mice: Impaired Lymphocyte Migration to Bone Marrow

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Abstract
We generated vascular cell adhesion molecule (VCAM)-1 “knock-in” mice and Cre recombinase transgenic mice to delete the VCAM-1 gene (vcam-1) in whole mice, thereby overcoming the embryonic lethality seen with conventional vcam-1–deficient mice. vcam-1 knock-in mice expressed normal levels of VCAM-1 but showed loss of VCAM-1 on endothelial and hematopoietic cells when interbred with a “TIE2Cre” transgene. Analysis of peripheral blood from conditional vcam-1–deficient mice revealed mild leukocytosis, including elevated immature B cell numbers. Conversely, the bone marrow (BM) had reduced immature B cell numbers, but normal numbers of pro-B cells. vcam-1–deficient mice also had reduced mature IgD1 B and T cells in BM and a greatly reduced capacity to support short-term migration of transferred B cells, CD4+ T cells, CD8+ T cells, and preactivated CD4+ T cells to the BM. Thus, we report an until now unappreciated dominant role for VCAM-1 in lymphocyte homing to BM.

Key words: VCAM-1 • Cre recombinase • knockout mice • bone marrow • lymphocyte migration

Introduction
Leukocyte trafficking is a dynamic orchestration of molecular cues. Numerous molecules participate in one or more steps in an integrated multistep process of leukocyte rolling on the blood vessel wall, followed by “activation,” firm adhesion, and transmigration into tissue (1–3). One of the players in this scenario is vascular cell adhesion molecule (VCAM)-1, which mediates both rolling on endothelial cells (ECs) and firm adhesion (4–7). VCAM-1 was originally identified on the surface of activated human umbilical vein ECs as a mediator of adhesion with melanoma cells and lymphocytes (8–10). VCAM-1 expression can be induced by numerous factors, including TNF-α, IL-1, IL-4, IL-13, and intercellular adhesion molecule (ICAM)-1 cross-linking (8–14). The ability of VCAM-1 to facilitate leukocyte adherence to ECs is thought to be a critical factor in the initiation and/or perpetuation of inflammation and autoimmunity. Upregulated VCAM-1 expression on ECs is associated with several diseases including multiple sclerosis, allograft rejection, atherogenesis, rheumatoid arthritis, appendicitis, dermatitis, and inflammatory bowel disease (15–23).

Besides inducible expression on ECs, VCAM-1 is expressed on peripheral LN (PLN) and mesenteric LN (MLN) high endothelial venules (HEVs; reference 24), bone marrow (BM) stromal cells (25), BM microvasculature (7, 26, 27), thymic epithelial cells (28), spleen stromal cells (29, 30), and spleen red pulp macrophages and dendritic cells (DCs) (31, 32). Although not normally expressed by T cells, VCAM-1 is reportedly expressed by thymocytes and T cells undergoing apoptosis (33). VCAM-1 on HEVs has an apparently redundant role in lymphocyte migration to LNs (24), whereas VCAM-1 on BM microvessels has an overlapping role with selectins in hematopoietic progenitor cell recruitment into the BM compartment (7, 34). VCAM-1 on BM stromal cells might have a role in B cell development. Anti–VCAM-1 antibody greatly reduced B lymphocyte formation in long-term BM
cultures (25). This is seemingly in contrast to studies with BM stromal cell clones from VCAM-1 null mice, where long-term maintenance and proliferation of clonal pre-B cells, cobblestone formation, and differentiation to IgM-secreting mature B cells were equally possible on VCAM-1+ and VCAM-1− BM stromal cells (35). Studies of α4 integrin–deficient hematopoietic progenitors in chimeric mice revealed a defect in α4 integrin–deficient B cell development at the pre-B cell stage (36), reinforcing the principle of a role in B cell development for VCAM-1 or some other α4 integrin receptor. That is, the heterodimeric integrins αβ1 (very late antigen [VLA]-4) and αβ2 bind VCAM-1, but VLA-4 (the principal VCAM-1 ligand) also binds fibronectin and itself via homotypic aggregation (37–44).

Recent studies also suggest that VCAM-1 might be more promiscuous than at first appreciated, in that αβ2 (expressed by a variety of leukocytes) and αβ1 (expressed by neutrophils) also bind to VCAM-1 (45–47). Thus, VCAM-1 may have roles besides the potential roles revealed by α4 integrin–deficient hematopoietic progenitors. Attempts by others to further investigate the roles of VCAM-1 using genetically deficient mice met with limited success because vcam-1 deficiency caused embryonic lethality in two independent studies (31, 48), although a few VCAM-1 null mice did survive embryonic lethality (31, 35). This hurdle has been overcome here by the generation of vcam-1 “knock-in” mice and “TIE2Cre” Cre recombinase transgenic mice. vcam-1 knock-in mice express normal levels of VCAM-1 but allow deletion of the vcam-1 gene promoter and first exon, using the Cre recombinase/loxP system. When intercrossed with TIE2Cre transgenes, vcam-1 knock-in mice showed virtually complete loss of VCAM-1 on ECs and hematopoietic cells. These semiregulated vcam-1−deficient mice had reduced immature B cells and mature lymphocytes in BM, and a greatly reduced capacity to recruit B cells and T cells to BM in short-term migration assays.

Materials and Methods

Generation of VCAM-1 Knock-in Mice. The VCAM-1 gene (vcam-1) was isolated from a λ Fix II 129/Ola library. The targeted region spans a 7-kb region between an upstream BamHI site and the 3′ end of the pGEM4 clone (see Fig. 1). This 3′ end does not include the BamHI site, which is expected to be slightly further downstream in vcam-1 (49–51). The 7-kb targeted region was subcloned into pBluescript II (Stratagene) as two separate halves: an upstream BamHI-EcoRI region and a downstream HindIII site and the 3′ end of the upstream EcoRI site (see Fig. 1). This modified upstream vcam-1 fragment after Klenow end-filling. A loxp site oligonucleotide duplex was designed to target the loxP site inserted into the desired HindIII site in intron 1 without loss of the small HindIII intron 1 fragment (see Fig. 1) were identified by digestion with BamHI, HindIII, XhoI, and combinations thereof.

A loxp-flanked neomycin resistance cassette was then inserted into the newly introduced XhoI site of the modified upstream vcam-1 clone as an XhoI-Sall fragment from pLox2neoA (see below). The desired orientation of the insert (see Fig. 1) was identified by digestion with BamHI, EcoRI, XhoI, and combinations thereof. This modified downstream vcam-1 clone constituted the right arm of the targeting construct. The construct pLox2neoA (Koni, P., and R. Flavell, unpublished results) contains a neomycin resistance cassette from pMC1neoA (Stratagene) flanked by loxp sites in pBluescript II (Stratagene), pLox2neoA was created with Sall and XhoI sites at the 5′ and 3′ ends (relative to the neomycin resistance cassette), respectively (as well as several other sites).

The left and right arms of the targeting construct were then joined by first excising the left arm, by partial digestion with EcoRI and then complete digestion at the pBluescript II polylinker NotI site. The full-length 4.5-kb left arm was then inserted into the right arm construct between the upstream polylinker NotI and EcoRI sites. Both the left and right arms of the targeting construct were therefore ~2.7 kb in size (excluding the 1.6-kb loxp-flanked vcam-1 promoter/exon 1 region). The targeting vector was linearized at the 3′ Sall site and 25 μg was used to electroporate 107 W9.5 embryonic stem (ES) cells. ES cells were then plated onto mitomycin C–treated primary embryonic fibroblasts. Double drug selection for homologous recombinants was begun 24 h later with 2 μM gancyclovir (Syntex) and 0.3 mg/ml G418 (GIBCO BRL).

ES cell colonies and subsequent mice were screened by BamHI digest Southern blot analysis using probes A and B (see Fig. 1). Probe A was a 1.0-kb EcoRI-EcoRV fragment. Probe B was a 1.0-kb SphI-Sall fragment at the 3′ end of the genomic clone. All probes were products of 32P incorporation by random priming using [32P]dCTP (Amersham Pharmacia Biotech) and a Prime-II kit (Stratagene). Homologous recombinant ES cells were injected into C57BL/6 blastocysts and chimeric males were bred to C57BL/6 females. Heterozygous targeted mice still bearing the neomycin resistance cassette in vcam-1 intron 1 (vcam-1+/neo) were then interbred to obtain vcam-1−/−neo mice and wild-type littersmates. All mice were housed in specific pathogen-free conditions in accordance with institutional animal care and use guidelines.

To avoid any possible interference with VCAM-1 expression,
the neomycin resistance cassette was then deleted to create the vcam-1lox allele (see Fig. 1 B for definitions). This “partial” deletion was achieved by use of the splicer mouse (see below).

**Generation of Cre Recombinase Transgenic Mice.** Splicer mice were generated with a transgene from pTet-Cre, which contains the Cre recombinase coding sequence from pBS185 (GIBCO BRL) cloned into the EcoRV site of pTet-Splice (GIBCO BRL) as a Klenow-blunted MluI-XhoI fragment. TIE2Cre transgenes were generated with a TIE2 kinase promoter/enhancer cassette described previously (53). The construct pStP.tT2PXK (pg54) (a gift from Thomas Sato, Beth Israel Hospital, Boston, MA) contained the TIE2 kinase promoter and enhancer with HindIII and NotI sites between the two. These sites allowed us to directionally clone a HindIII–NotI fragment from pTet-Cre, thus introducing the Cre recombinase coding sequence, intronic donor/acceptor sequences, and polyadenylation signal sequences into pg54. This TIE2Cre transgene was then excised from the vector backbone using SalI.

All transgenic mice were generated on a (C3H × C57BL/6)F₂ background. Screening of tail DNA for Cre recombinase transgene presence was by PCR with the following primers: forward, 5'-CGATGCAACGAGTGATGAGG-3' and reverse, 5'–CGCATACACGTGAAACGCG-3'. Positive founder mouse lines were then crossed with C57BL/6 mice for two generations before interbreeding with VCAM-1 knock-in mice.

**Lung Challenge and Spleen Immunohistochemistry.** Mice were given an intraperitoneal injection of alun-precipitated antigen consisting of 20 μg OVA (grade V; Sigma-Aldrich) adsorbed onto 2 mg aluminum hydroxide in 0.1 ml PBS. A booster injection was given 5 d later. Control animals received precipitated alun in PBS only. 12 d after the first sensitization, mice were challenged by exposure to an aerosol of 0.5% OVA in PBS, twice for 1 h each with a 4-h interval. This was done by placing the mice in a Plexiglass chamber attached to an ultrasonic nebulizer (1–5 particles/μm) and then dialyzed against PBS. Mice 6–8 wk of age were then challenged intraperitoneally with 50 μg of NP₆C₆G adsorted to alun in 0.1 ml PBS. Spleens were harvested at day 10 after the challenge, frozen in Tissue-Tek OCT compound using a dry ice/methylbutane bath, and stored at −70°C until cutting. Sections of 7-μm thickness were cut onto silanized glass slides, fixed in cold acetone for 10 min, and then stored at −70°C until use. Slides were then stained as above after thawing and rehydration in PBS.

**Humoral Challenge and Spleen Immunohistochemistry.** Chicken γ-globulin (CG; Sigma-Aldrich) was conjugated with (4-hydroxy-3-nitrophenyl) NP acetyl succinimide ester (Calbiochem) in 0.1 M sodium borate, pH 9.2, to an NP/C₆G molar ratio of 13:1 (NP₆C₆G), and then dialyzed against PBS. Mice 6–8 wk of age were then challenged intraperitoneally with 50 μg of NP₆C₆C₆G adsorbed to alun in 0.1 ml PBS. Spleens were harvested at day 10 after the challenge, frozen in Tissue-Tek OCT compound using a dry ice/methylbutane bath, and stored at −70°C until cutting. Sections of 7-μm thickness were cut onto silanized glass slides, fixed in cold acetone for 10 min, air dried, and then stored at −70°C until use. For staining, sections were thawed for 30 min and then rehydrated in PBS for 20 min. Endogenous peroxidase was quenched with 0.3% hydrogen peroxide for 5 min. Sections were washed in PBS for 10 min and then preblocked with PBS/3% BSA/0.1% Tween 20 for 30 min in a humidified chamber. Staining for IgG was with rat anti-IgG (11–26) (Southern Biotechnology Associates, Inc.) and then horseradish peroxidase–conjugated polyclonal goat anti-rat IgG (Southern Biotechnology Associates, Inc.). Germinal centers were stained with biotin–conjugated peanut agglutinin (PNA) (EY Laboratories) and then rehydrated in PBS. Mice 6–8 wk of age were then challenged intraperitoneally with 50 μg of NP₆C₆G adsorbed to alun in 0.1 ml PBS. Spleens were harvested at day 10 after the challenge, frozen in Tissue-Tek OCT compound using a dry ice/methylbutane bath, and stored at −70°C until cutting. Sections of 7-μm thickness were cut onto silanized glass slides, fixed in cold acetone for 10 min, air dried, and then stored at −70°C until use. For staining, sections were thawed for 30 min and then rehydrated in PBS for 20 min. Endogenous peroxidase was quenched with 0.3% hydrogen peroxide for 5 min. Sections were washed in PBS for 10 min and then preblocked with PBS/3% BSA/0.1% Tween 20 for 30 min in a humidified chamber. Staining for IgD was with rat anti-IgD (11–26) (Southern Biotechnology Associates, Inc.) and then horseradish peroxidase–conjugated polyclonal goat anti-rat IgG (Southern Biotechnology Associates, Inc.). Germinal centers were stained with biotin–conjugated peanut agglutinin (PNA) (EY Labs; reference 55). The presence of VCAM-1 was assessed with anti–VCAM-1 (M/K-2, rat IgG1, κ–biotin (Southern Biotechnology Associates, Inc.). Anti-CD88.2 (53–5.8, rat IgG1, κ–biotin was used as a negative control for VCAM-1 staining. All biotin conjugates employed a secondary step of alkaline phosphatase–conjugated streptavidin (Zymed Laboratories). Incubations were done overnight at 37°C for 1 h. Following washing steps, RNAs were hybridized with biotinylated DNA probes specific for VCAM-1 (Zymed Laboratories) and then rehydrated in PBS for 20 min. Substrates for horseradish peroxidase and alkaline phosphatase were diaminobenzidine (DAB; Zymed Laboratories) and 5-bromo-4-chloro-3-indolyl phosphate (BCIP) (Zymed Laboratories). Mounting was with Clearmount (Zymed Laboratories).

**Spleen Collagenase Digestion.** Spleens were harvested into 2 ml digest buffer in a 35-mm dish on ice. Digest buffer was calcium-free HBSS with 1% FCS, 2 mm l-glutamine, 0.5 mg/ml collagenase type IV (Sigma-Aldrich), and 0.1 mg/ml deoxyribonuclease type I (Sigma-Aldrich). Spleens were washed with 23-gauge needles before incubation at 37°C for 30 min. The cell suspension was further disrupted by pipetting before being made up to 10 ml with PBS/5 mM EDTA. Splenocytes were then centrifuged at 1,000 rpm for 5 min and resuspended into 2 ml ACK erythrocyte lysis buffer (BioWhittaker). 1 min later, the suspension was again made up to 10 ml with PBS/5 mM EDTA and re-centrifuged. Finally, cells were resuspended in PBS/1% FCS and filtered through 0.1-mm nylon mesh (Millipore).

**Fluorocytometry.** For fluorocytometry of DCs, cells were pre-
pared from spleen by collagenase digestion as described above. Otherwise, cells were recovered into 5 ml PBS/1% FCS from thymus, Peyer’s patches, spleen, and/or LNs by using the plunger of a syringe to tease the tissue between two pieces of 0.1-mm nylon mesh (Millipore). Total BM was collected from femurs and tibias. Splenocytes were further treated by centrifugation at 1,000 rpm for 5 min, resuspension into 2 ml ACK erythroid cell lysis buffer (Bio-Whittaker), and then addition of 10 ml PBS 1 min later. All cell suspensions were then centrifuged at 1,000 rpm for 5 min. Finally, cells were resuspended into PBS/1% FCS and filtered through 0.1-mm nylon mesh. BM was not subjected to erythroid cell lysis, but total nucleated cell numbers in both BM and EDTA-treated blood were determined by counting in Turk’s solution (0.01% wt/vol gentian violet in 3% vol/vol glacial acetic acid).

Aliquots of 10⁶ nucleated cells were made into 0.2 ml PBS/1% FCS supplemented with 5 𝜇g/ml FcBlock (BD Pharmingen). Samples were left on ice for 30 min before primary antibodies were added and left on ice in the dark for another 1 h. Samples were washed by the addition of 1 ml PBS/1% FCS and centrifugation at 1,000 rpm, 4°C for 5 min. Secondary antibody incubation and washing were as above. Four-color fluorocytometry employed a FACSCalibur™ with argon and red diode lasers (Becton Dickinson). Fluorocytometry of spleen DCs, BM cells, and 5-chloromethylfluorescein diacetate (CMFDA)-labeled cells (see below) was done by counting 250,000 events. Otherwise, 50,000 events were collected.

Anti–mouse IgM (donkey polyclonal)-Cy5 and antidigoxin (mouse IgG polyclonal)-Cy5 were from Jackson ImmunoResearch Laboratories. Anti-IgD (11-26c.2a)-digoxigenin was made (mouse IgG polyclonal)-Cy5 were from Jackson ImmunoResearch Laboratories. Anti–mouse IgM (donkey polyclonal)-Cy5 and antidigoxin (mouse IgG polyclonal)-Cy5 were from Jackson ImmunoResearch Laboratories. Anti-IgD (11-26c.2a)-digoxigenin was made and employed by us as described previously (56). All other reagents were from BD Pharmingen.

**In Vivo Lymphocyte Homing Assay.** C57BL/6J and tcr⁻⁻⁻⁻ mouse were purchased from The Jackson Laboratory and breeding colonies were maintained in our animal facility according to institutional animal care and use guidelines. Cells to be labeled were either LN (axillary, brachial, inguinal, and mesenteric) cells from C57BL/6J mice, splenocytes from tcr⁻⁻⁻⁻ mice (after erythroid cell lysis and washing as described earlier), or preactivated T cells. The latter were prepared by culturing LN and spleen cells from C57BL/6J mice (after erythroid cell lysis and washing as described above) for 3 d at 5 × 10⁶ cells/ml in DMEM supplemented with 2 mM glutamine, 50 𝜇M 2-mercaptoethanol, 100 IU/ml penicillin, 0.1 mg/ml streptomycin, 5% FCS, 1 𝜇g/ml anti-CD3 antibody (clone 2C11; Southern Biotechnology Associates, Inc.), and 50 U/ml IL-2. Cells were then washed and rested for a further 3 d in the same medium but without anti-CD3 antibody. Finally, dead cells were removed by centrifugation through lymphocyte separation medium (BioWhittaker).

Cells were labeled with 1 𝜇M CMFDA succinimidyl ester (Molecular Probes) in PBS/0.1% BSA for 5 min at 37°C, and then washed three times with PBS by centrifugation at 1,000 rpm for 5–10 min. Finally, labeled cells were made to 2 × 10⁶/ml in PBS and each 8–10-wk-old recipient was given 0.1 ml intravenously. 2 h later, peripheral blood, spleen, PLNs (axillary, brachial, and inguinal LNs), MLNs, and BM (femurs and tibias) were harvested for fluorocytometry as described above, with 250,000 events being counted per sample.

**Figure 1.** Targeting strategy and conditional deletion of the *vcam-1* allele. (A) VCAM-1 knock-in mice (bottom locus map) contain Cre recombine sites of recombination (loxP sites; black arrowheads). The *vcam-1* coding sequence exons are depicted as striped bars, with the 5’ untranslated region as a white bar. Probes A and B are shown above the top map line as black bars. Endonuclease sites shown are BamHI (B), EcoRI (R), HindIII (H), SphI (S), and XhoI (X). (B) The *vcam-1* allele (neo) is shown along with two of the possible outcomes of Cre recombinase-mediated deletion, the *vcam-1* allele (floxed) and the *vcam-1* allele (Δ). (C and D) BamHI Southern blot analysis of tail DNA using probes A and B, respectively. Deletion of the *vcam-1* allele to generate the *vcam-1* allele (floxed) and the *vcam-1* allele (Δ) in whole mice was achieved by interbreeding with the *TIE2Cre* transgene. (E) Southern blot analysis of mouse tail DNA as in D. A *vcam-1* allele (track 1) is shown alongside two *TIE2Cre* mice born of *vcam-1* allele (floxed) parents, but with the *TIE2Cre* parent being the father (track 2) or mother (tracks 1 and 3). (F and G) Southern blot analysis as in D of genomic DNA from a *vcam-1* allele (floxed)/*TIE2Cre* mouse and a *vcam-1* allele (floxed)/*TIE2Cre* mouse, respectively. Tissues analyzed were: BM; Br, brain (cortex); H, heart; Ki, kidney; Li, liver; Lu, lung; LNs; M, muscle (thigh); Pa, pancreas; and Sp, spleen.
Results

Generation of Conditional vcam-1−deficient Mice. Mice bearing Cre recombinase loxP sites of recombination were generated by homologous recombination in 129/Ola ES cells using conventional techniques (Fig. 1). The vcam-1 region flanked by loxP sites includes the defined cytokine-responsive promoter region (49) and exon 1, with the loxP-flanked neomycin resistance cassette in intron 1. Exon 1 contains the signal peptide sequence and is critical to all the alternatively spliced forms of vcam-1 (49–51). This region of vcam-1 was previously deleted in mice using conventional techniques and resulted in embryonic lethality, as described by others (31, 48). Thus, the targeting strategy employed here was believed to be appropriate for achieving vcam-1 inactivation, once the loxP-flanked region is deleted. The neomycin resistance cassette was removed by use of the splicer mouse (see Materials and Methods) to avoid any possible interference of VCAM-1 expression. Indeed, vcam-1fl/fl mice had four- to fivefold lower VCAM-1 on spleen DCs compared with vcam-1flox/+ mice and C57BL/6j mice (data not shown).

Conditional deletion of the vcam-1 flox allele (see Fig. 1 B for definitions) was achieved with a TIE2Cre Cre recombinase transgene. The latter was generated with the TIE2 kinase promoter/enhancer expression cassette, which gives uniform expression in all ECs during both embryogenesis and adulthood (53). When both parents had a vcam-1 flox/flox genotype and the TIE2Cre + parent was a female, all of the progeny had a vcam-1 flox/Δ genotype regardless of whether or not they themselves were TIE2Cre + (Fig. 1 E). Thus, the TIE2Cre transgene appeared to delete the vcam-1 allele in the germline of the mother.

Both vcam-1 flox/flox/TIE2Cre + mice (born with the TIE2Cre + parent being the father) and vcam-1 flox/Δ/TIE2Cre + mice had extensive deletion of the vcam-1 flox allele in several tissues, but especially in tissues rich in hematopoietic cells (Fig. 1, F and G). This was anticipated because de novo TIE2 kinase is expressed by hematopoietic progenitors and ECs (57, 58). Virtually 100% deletion was seen in the BM of vcam-1 flox/Δ/TIE2Cre + mice (Fig. 1 G).

The high degree of deletion in lung tissue of vcam-1 flox/flox/TIE2Cre + mice was reflected by a complete lack of VCAM-1 on lung ECs of vcam-1 flox/flox/TIE2Cre + mice after OVA sensitization and aerosol challenge, which upregulates VCAM-1 expression in the lung (Fig. 2 A). VCAM-1 expression was also examined on BM myeloid lineage cells, spleen myeloid DCs, and spleen lymphoid DCs by fluorocytometry. This revealed relatively normal levels of VCAM-1 in vcam-1 flox/flox mice (Fig. 2 B), whereas vcam-1 flox/Δ mice had four- to fivefold lower VCAM-1 (data not shown). VCAM-1 levels were greatly reduced in vcam-1 flox/Δ/TIE2Cre + mice, but complete deletion was achieved only in vcam-1 flox/Δ/TIE2Cre + mice (Fig. 2 B). As illustrated below (see Fig. 6), VCAM-1 was also deleted from spleen red pulp macrophages.

Finally, ROSA26R mice (54) were employed to further visualize deletion by the TIE2Cre transgene. ROSA26R mice express β-galactosidase only after Cre recombinase–mediated deletion of a “STOP” signal, and thus allow a histological evaluation of sites of Cre recombinase activity by staining for β-galactosidase activity (54). Thus, male TIE2Cre + mice were crossed with ROSA26R + mice (54) to obtain double-positive and ROSA26R +/TIE2Cre − progeny. Double-positive mice clearly revealed Cre recombinase activity as blue staining in the ECs of brain, kid-
ney, LNs, spleen, and thymus (Fig. 3), while no staining was seen in ROSA26R+/TIE2Cre− mice (data not shown). The blue staining of capillaries seen in the brain and thymus sections was typical. The intense blue staining of HEVs in the axillary LNs shown was typical of HEVs, and was also representative of thymic LNs and MLNs (data not shown). The relatively high staining intensity seen with HEV cells is presumably a reflection of the degree of β-galactosidase expression driven by the ROSA26R reporter promoter. It is not a reflection of more or less Cre recombinase–mediated deletion of the ROSA26R reporter allele because there is only one ROSA26R allele per cell. The completeness of deletion among HEV cells is witnessed at least by the blue coloration of the entire circumference of HEVs in every example seen. There was also consistent staining of arterioles and larger vessels in spleen sections and uniform staining of B cell follicles, in agreement with the earlier observation of complete deletion of the vcam−/flox allele in hematopoietic cells. The apparent lack of staining in T cell areas of LNs and spleen as well as the thymus medulla may be due to the fact that T cells have a β-galactosidase inhibitory activity, as noted by others (59).

Altered Lymphocyte Populations in Peripheral Blood and the BM Compartment. Fluorocytometry of cells from thymus, spleen, MLNs, PLNs (axillary, brachial, and inguinal), and Peyer’s patches of vcam−/flox and vcam−/flox/TIE2Cre+ mice did not reveal any significant differences in leukocyte types or absolute numbers (data not shown). However, analysis of peripheral blood revealed mild leukocytosis in vcam−/flox/TIE2Cre+ mice, including elevated levels of B220+IgDloIgMhi immature B cells (Fig. 4). Peripheral blood granuloid cells were also elevated in vcam−/flox/TIE2Cre+ mice compared with their vcam−/flox/TIE2Cre− littermates, although this was not statistically significant compared with the age- and sex-matched vcam−/flox mice.

Examination of BM from vcam−/flox/TIE2Cre+ mice revealed a significant reduction in IgDloIgMhi immature B cells (Fig. 5). However, there was no significant difference in numbers of B220+CD4+CD8− pro-B cells (60), which represented ~5% of total nucleated BM cells in both vcam−/flox mice and vcam−/flox/TIE2Cre+ mice (Fig. 5). IgD+ B cells and CD8+ T cells were also reduced compared with vcam−/flox mice, with a small but statistically insignificant reduction in total B220+ B cells and CD4+ T cells (Fig. 5). Total nucleated BM cell numbers were very similar in the

Figure 3. The TIE2Cre transgene activates the ROSA26R reporter allele and results in β-galactosidase activity in brain (Br), kidney (Ki), liver (Li), axillary LNs (LN), spleen (Sp), and thymus (Th), revealed by blue X-Gal staining (original magnification: X100). Brain sections were 30 µm thick while all others were 10 µm. By way of example, thick and thin arrows in the spleen section indicate some larger vessels and arterioles, respectively.

Figure 4. Peripheral blood analysis reveals leukocytosis in vcam−/flox/TIE2Cre+ mice. A representative experiment (n = 5 per group) with age- and sex-matched vcam−/flox mice (black bars), vcam−/flox/TIE2Cre+ mice (striped bars), and their vcam−/flox/TIE2Cre+ littermates (white bars). Total leukocyte numbers were determined from EDTA-treated blood in Turk’s solution and are shown in millions of cells per milliliter of blood. Differential counts were established by fluorocytometry. PMNs were defined on the basis of high side-scatter, with the other cells defined as mononuclear (mono). The numbers of CD4+ T cells, IgD+IgMlo immature B cells, and IgD+ B cells shown on the right are not in addition to the mononuclear cell counts but are part of the latter. Significant differences between vcam−/flox mice and vcam−/flox/TIE2Cre+ mice are indicated (Student’s t test, *P < 0.05; **P < 0.01).
numbers in peripheral blood, *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ mice showed relatively normal spleen B cell follicles with robust germinal center (GC) formation upon intraperitoneal challenge with NP\(_{10}\)CG adsorbed to alum (Fig. 6). There was also no significant difference in anti-NP IgG1 levels in the serum of *vcam-1*\[^{-}\]/\[^{\Delta}\]/ mice (\(n = 4\)) versus *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ mice (\(n = 8\)) at days 10 and 35 after challenge (data not shown). Finally, the latter mice were given a booster injection of 0.2 mg soluble NP\(_{10}\)CG in PBS at day 35 after the challenge and serum anti-NP IgG1 was determined 6 d later. A significant difference was not seen between the two groups of mice (data not shown).

Of note is the fact that *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ mice still had VCAM-1 on follicular dendritic cells (FDCs) (Fig. 6). Also, VCAM-1 was not completely absent from the spleen red pulp compared with the negative staining by the anti-CD8b control (Fig. 6), presumably a result of VCAM-1 expression by spleen stromal cells (29, 30).

**Impaired Lymphocyte Migration to BM.** Having observed reduced lymphocyte numbers in BM, short-term migration assays were performed with CMFDA-labeled lymphocytes from LNs of C57BL/6J donor mice. The absolute numbers of CMFDA-labeled cells recovered from the spleens of *vcam-1*\[^{-}\]/\[^{\Delta}\]/ recipients versus *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ recipients were not significantly different (data not shown). The fractions of CMFDA-labeled cells recovered in spleen, MLNs, and PLNs were also similar in the two groups of mice, but *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ mice clearly had greatly reduced CMFDA-labeled cells in BM (Fig. 7). Short-term migration of IgD^+^, CD4^+^, and CD8^+^ cells to the BM compartment of *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ recipients was reduced on average by ~93, 74, and 77% compared with the *vcam-1*\[^{-}\]/\[^{\Delta}\]/ recipients, respectively. The actual frequencies of CMFDA-labeled IgD^+^ cells among total nucleated BM cells in *vcam-1*\[^{-}\]/\[^{\Delta}\]/ recipients and *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ recipients were 0.118 ± 0.022 and 0.008 ± 0.002%, respectively.

Short-term migration assays were also performed with CMFDA-labeled splenocytes from *tcra^\(-/-\)* donors, revealing that the reduced short-term migration to BM by IgD^+^ B cells was also true of other B cell subsets (Fig. 8). This reduced short-term migration to BM was reflected in elevated cell numbers in peripheral blood (Fig. 8). The exact identity of the cells among the IgD^+^IgM^hi^ subset that had either homed to BM or remained in peripheral blood was not determined, but these cells might have included marginal zone memory B cells and/or immature “transitional” B cells.

To further dissect the requirement for VCAM-1 in CD4^+^ T cell migration, short-term migration assays were then performed with preactivated/experienced T cells (Fig. 9). This revealed greatly impaired short-term migration to BM of *vcam-1*\[^{-}\]/\[^{\Delta}\]/TIE2Cre^\+ mice by both CD4^+^CD62L\[^{lo}\] T cells and CD4^+^CD62L\[^{hi}\] T cells (75 and 70% reduction, respectively). The absolute numbers of CMFDA-labeled cells in spleen and PLNs mirrored the earlier results, with no apparent difference (Fig. 9). Also, as might be expected, CD4^+^CD62L\[^{lo}\] cells preferentially homed to PLNs compared with CD4^+^CD62L\[^{hi}\] cells (Fig. 9).
Discussion

We have employed the Cre recombinase/loxP system to generate \textit{vcam-1} knock-in (\textit{vcam-1}^{flx/flx}) mice. We also generated a Cre recombinase transgene (TIE2Cre) to delete this conditional \textit{vcam-1} allele in whole mice. Analysis of sites of TIE2Cre activity with the ROSA26R reporter strain (54) showed the TIE2Cre transgene to be capable of mediating deletion in ECs of at least brain, kidney, LNs, spleen, and thymus. The β-galactosidase staining of HEV in LNs, for example, revealed TIE2Cre-mediated deletion in every single HEV cell (Fig. 3), suggesting 100% penetration by the transgene. Furthermore, virtually complete loss of VCAM-1 was seen on lung ECs, BM myeloid cells, spleen myeloid DCs, and spleen lymphoid DCs of \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} mice. The apparent absence of VCAM-1 cannot be explained as being a result of loss of only the anti–VCAM-1 antibody epitope (rather than VCAM-1 per se), because the deletion strategy employed does not remove any of the exons encoding the mature VCAM-1 protein. These data were then represented as the percentage of homed cells, where 100% is the total number of recovered CMFDA-labeled cells in each individual mouse (\textit{n} = 3 per group). *Significantly different from controls (Student’s \textit{t} test, \textit{P}<0.02). The slightly elevated migration to PLNs of \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} recipients was not statistically significant.

Figure 6. Relatively normal B cell responses in \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} mice. Serial spleen sections from a \textit{vcam-1}^{flx/flx} mouse (WT) and a \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} mouse (KO) 10 d after intraperitoneal challenge with NP\textsubscript{3}CG adsorbed to alum (original magnifications: ×100). All sections were stained with anti- IgD antibody (seen in brown) and either PNA, anti-CD8b, or anti–VCAM-1. PNA, CD8b, and VCAM-1 staining are in purple. Arrows in the KO section indicate typical VCAM-1 staining on FDCs within GCs.

Figure 7. Reduced short-term lymphocyte migration to BM in \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} mice. A representative experiment with age- and sex-matched \textit{vcam-1}^{flx/flx} recipients (black bars) and \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} recipients (white bars). The frequency of CMFDA-labeled cells (IgD\textsuperscript{+}, CD4\textsuperscript{+}, and CD8\textsuperscript{a+}) was used to determine the absolute numbers of each type of CMFDA-labeled cell in each lymphoid organ. These data were then represented as the percentage of homed cells, where 100% is the total number of recovered CMFDA-labeled cells in each individual mouse (\textit{n} = 3 per group). *Significantly different from controls (Student’s \textit{t} test, \textit{P}<0.02). The slightly elevated migration to PLNs of \textit{vcam-1}^{flx/flx}/TIE2Cre \textsuperscript{+/} recipients was not statistically significant.
polypeptide. Rather, we have deleted the VCAM-1 promoter region and exon 1. The latter contains the signal peptide sequence and is critical to all of the alternatively spliced forms of vam-1 (49–51).

Our studies reveal a dominant role for VCAM-1 in lymphocyte migration to BM. Having said this, we have not yet definitively demonstrated that VCAM-1 is actually absent from BM ECs in vam-1<sup>flox/Δ</sup>/TIE2Cre<sup>+</sup> mice. Although we observed virtually 100% deletion of the vam-1<sup>flox</sup> allele in BM (Fig. 1 G), the vast majority of the BM genomic content is undoubtedly derived from hematopoietic cells. This issue will be addressed by the generation of BM stromal cell lines. Nonetheless, short-term migration of B cells, CD4<sup>+</sup> T cells, and CD8<sup>+</sup> T cells to the BM of vam-1<sup>flox/Δ</sup>/TIE2Cre<sup>+</sup> mice was reduced on average by ∼80, 74, and 77%, respectively. Also, short-term migration by CD4<sup>+</sup>CD62L<sup>hi</sup> and CD4<sup>+</sup>CD62L<sup>lo</sup> preactivated/experienced T cells was reduced on average by ∼75 and 70%, respectively.

It is conceivable that the low levels of lymphocytes still found in the BM of vam-1<sup>flox/Δ</sup>/TIE2Cre<sup>+</sup> mice are a result of alternate or complimentary mechanisms of migration. Certainly, this would not be unprecedented. For example, ICAM-1–deficient mice show reduced but substantial contact hypersensitivity and neutrophil migration (61, 62). Likewise, LFA-1–deficient lymphocytes show reduced but not absent lymphocyte migration to PLNs, MLNs, and Peyer’s patches (24, 63). Also, P/E-selectin double deficiency reduces hematopoietic progenitor cell rolling at BM microvessels to 30–40% of control levels (7). This selectin-independent rolling is further reduced ∼70% by anti–VCAM-1 antibody, while anti–VCAM-1 antibody treatment of wild-type BM microvessels reduces hematopoietic progenitor cell rolling by only ∼30% (7). Finally, anti–VCAM-1 antibody greatly reduces the migration of LFA-1–deficient lymphocytes to BM (24). We have shown that greatly reduced migration can also be achieved with VCAM-1 deficiency alone, and that this applies to all three major lymphocyte subsets. Taken together, these observations suggest that both LFA-1 receptors and VCAM-1 are involved in lymphocyte migration to BM, but that VCAM-1 plays a dominant role.

We have not yet definitively determined whether or not VCAM-1 is deleted from BM stromal cells of vam-1<sup>flox/Δ</sup>/TIE2Cre<sup>+</sup> mice. This is currently the subject of further investigation by the generation of BM stromal cell lines. This is of potential interest because VCAM-1 on BM stromal cells might prove to have a role in B cell development and/or humoral response beyond those that have been revealed by our vam-1<sup>flox/Δ</sup>/TIE2Cre<sup>+</sup> mice. For example, α<sub>i</sub> integrin–deficient hematopoietic progenitor cell chimeric mice had greatly reduced levels of α<sub>j</sub> integrin–deficient pre-B cells (36), and it has not yet been determined whether the α<sub>i</sub> integrin receptor involved in pre-B cell development is VCAM-1, fibronectin, or both. Also, anti–VCAM-1 antibody greatly reduced B lymphocyte formation in long-term bone marrow cultures (25). However, this is in contrast to studies with BM stromal cell clones from VCAM-1
null mice where long-term maintenance and proliferation of clonal pre-B cells, cobblestone formation, and differentiation to IgM-secreting mature B cells were equally possible on VCAM-1+ and VCAM-1− stromal cells (35).

In this issue, Leuker et al. (64) have also generated conditional vcam-1−deficient mice using an IFN-inducible Cre recombinase system. These authors show that their vcam-1−deficient mice have a phenotype similar to our mice. In addition, they show relatively low antigen-specific serum antibody several weeks after challenge, after secondary challenge (64). We did not find reduced humoral responses in our mice (data not shown), perhaps due to differences in humoral challenge. Having said this, we anticipate that this difference in humoral response will be determined to be because of the nature of our respective conditional VCAM-1 deletion strategies. That is, the IFN-induced vcam-1−deficient mice of Leuker et al. might lack VCAM-1 on cells that our mice do not (e.g., FDCs, spleen stromal cells, and/or BM stromal cells). First, in vitro studies suggest that VCAM-1 on FDCs participates in adherence with, and prevents apoptosis of, GC B cells (65). Second, human spleen-derived stromal cells promote B cell blast differentiation and survival resulting in enhanced antibody secretion, in a manner that can be partly blocked by anti–VCAM-1 antibody (66). Third, BM fibroblasts can rescue cells from apoptosis (67), conceivably in a VCAM-1–dependent manner. As mentioned below, plasma cells are known to express the VCAM-1 ligand, VLA-4. Thus, antibody-secreting cell differentiation and/or longevity might be perturbed in mice lacking VCAM-1 on spleen and/or BM stromal cells.

Humoral responses in our mice are being further investigated, but clearly any difference between our mice and those of Leuker et al. (64) is not because of a lack of deletion of VCAM-1 on conventional DCs in our mice (Fig. 2 B).

In conclusion, we have generated conditional vcam-1−deficient mice and found them to have a reduced capacity to recruit lymphocytes to BM. Our short-term lymphocyte homing studies showed that migration to wild-type mouse BM (femur and tibia) represents ~3–5% of the total homed cells among the organs examined. Given that this represents only BM from femur and tibia, the total migration to BM probably represents a substantial fraction comparable in size to that seen in MLNs and PLNs. As suggested by others (24), perhaps BM should be considered as a major part of the lymphocyte recirculation network. Besides being a home for antibody secreting cells, the physiological relevance of lymphocyte recirculation to BM is not fully understood. The BM compartment is capable of functioning as a site of primary immune function under conditions of disrupted lymphocyte trafficking to spleen and LNs (68). It remains to be seen whether or not the BM compartment can act as a priming site in other physiological conditions. The pathophysiology of lymphocytes in BM, however, is perhaps better appreciated. BM is a major site of involvement in B and T cell malignancy (69), and metastasis to BM by lymphomas is often a very poor prognostic indicator. Both normal human plasma cells and myelomas express VLA-4 (70), and myelomas home to the BM where they induce massive osteoclastic bone destruction (71, 72). Anti–VCAM-1 antibody also greatly suppresses myeloma cell adhesion to BM ECs in vitro (73). Furthermore, direct cell–cell contact between myeloma cells and BM stromal cells via VCAM-1/VLA-4 is critical in some models to the production of destructive bone resorption (74). Finally, the most common symptoms in childhood acute leukemia, for example, are fever, infection, and bleeding resulting from neutropenia, anemia, and thrombocytopenia, which are secondary consequences of BM infiltration by leukemic cells (69). Thus, it might be beneficial to be able to prevent such cells from homing to and/or flourishing in the BM compartment.

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