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*J Immunol* 2012; 189:1340-1348; Prepublished online 29 June 2012;
doi: 10.4049/jimmunol.1200442
http://www.jimmunol.org/content/189/3/1340

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Chronic Alcohol Consumption Impairs Distribution and Compromises Circulation of B Cells in B16BL6 Melanoma-Bearing Mice

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Accumulating research indicates that B cells are involved in anti-tumor immunity. Chronic alcohol consumption is associated with decreased survival of cancer patients. The effect of alcohol consumption on B cells in tumor-bearing hosts is unknown. Results in melanoma-bearing mice showed that chronic alcohol consumption did not alter the percentage and number of B cells in bone marrow, spleen, and lymph nodes but dramatically decreased B cells in the peripheral blood. Alcohol consumption did not alter the development of B cells in the bone marrow and did not affect follicular B cells in the spleen; however, it increased T1 B cells and decreased marginal zone B cells in the spleen. Alcohol consumption also decreased mature B cells in the blood. It did not alter the chemotactic capacity of plasma to facilitate migration of splenocytes or the chemotactic response of splenocytes to CXCL13 and CCL21. However, the response of splenocytes to sphingosine-1-phosphate was impaired in alcohol-consuming, melanoma-bearing mice. The expression of sphingosine-1-phosphate receptor-1 (S1PR1) and sphingosine-1-phosphate lyase-1 (SPL1) in splenocytes was downregulated. Taken together, these results indicate that chronic alcohol consumption decreases peripheral blood B cells by compromising B cell egress from the spleen. The downregulation of S1PR1 and SPL1 expression in alcohol-consuming, melanoma-bearing mice could be associated with compromised egress of B cells from the spleen.


B cells are a large population of lymphocytes that originate from the bone marrow (BM) and further mature in the spleen. These cells play important roles in humoral immunity. Immature B cells in the spleen are called transitional B cells and are usually divided into two subsets, T1 B cells, which phenotypically are CD19+CD93+CD23hi, and T2 B cells, which phenotypically are CD19+CD93+CD23+ (1). Mature splenic B cells can also be divided into two subsets: follicular B cells and marginal zone (MZ) B cells. Follicular B cells circulate in the body and are characterized by the following phenotype: CD19+CD21hiIgDhiIgMlo. MZ B cells are noncirculating cells. These splenic B cells are the first line of defense against blood-borne Ags. MZ B cells express activated innate immune cell features, and their signature cell surface markers are CD19+CD1dhi CD21hi (2, 3).

B cells not only play a pivotal role in humoral immunity by producing Ab but also are involved in cellular immune responses by presenting Ag to T cells. B cells are the largest population of APCs in the immune system. They play an important role in Ag presentation, especially under low protein Ag conditions (4, 5). MZ B cells express high levels of CD1d, which is the molecule that presents lipid Ags to NKT cells, and they are key players in regulating the cytokine profile and activation of NKT cells (6–8). Because B cells must capture Ag to activate themselves to produce Ab and present Ag to T cells, effective recirculation among lymphatic organs and tissues is crucial for these cells to fulfill their function. B cell circulation and redistribution in lymphoid organs are directed by the CXCL13 chemokine and its receptor, CXCR5 (9, 10). The egress of B cells from lymph nodes (LNs), spleen, and BM is controlled by sphingosine-1-phosphate (S1P) and its sphingosine-1-phosphate receptor-1 (S1PR1) (11–13).

Compared to extensive studies involving T cells in anti-tumor immunity, the knowledge of B cells in anti-tumor immune responses is limited and controversial. Some studies indicate that depletion of B cells inhibits tumor progression and enhances anti-tumor immunity, suggesting that B cells inhibit anti-tumor immune responses (14, 15). One of the reasons associated with the inhibitory effects of B cells on anti-tumor immunity is that some B cells produce high levels of IL-10, which inhibit T cell function (14). This suggests the existence of regulatory B cells. Recently, it was found that B cells exhibiting the CD5+CD1dhi phenotype produce IL-10 upon activation by LPS. These CD5+CD1dhi IL-10–producing cells are one subset of regulatory B cells called B10 cells (16, 17). Other studies indicate that B cells enhance T cell anti-tumor immunity. Depletion of B cells enhances B16 melanoma metastasis to the lung by inhibiting CD8+ T cell proliferation and Th1 cytokine production (18). Deficiency of mature B cells significantly compromises T cell-mediated anti-tumor immunity induced by the anti-glucocorticoid–induced TNFR-related protein mAb, DTA-1 (19). Adaptive transfer of CpG-activated B cells inhibits tumor progression, whereas depletion of mature B cells increases tumor burden in the lungs of CpG-treated mice (20).

Chronic alcohol consumption in the melanoma-bearing mice leads to inhibition of memory T cell and especially tumor-specific CD8+ T cell expansion. This causes T cell dysfunction in B16BL6 melanoma-bearing mice (21). The underlying mechanism remains to be elucidated. Chronic alcohol consumption decreases B cells...
in the spleen of mice not inoculated with tumor (22). The relationship between chronic alcohol consumption and B cell antitumor immunity is unknown. In this study, using a chronic alcohol-consuming mouse model, we report that alcohol consumption 1) decreases B cells in the blood but not in the spleen, LNs, or BM and 2) decreases MZ B cells and B10 cells in the spleens of mice inoculated s.c. with B16BL6 melanoma.

Materials and Methods

Mice and alcohol administration

Female C57BL/6 mice at 6–7 wk of age were purchased from National Institutes of Health/National Cancer Institute Charles River Laboratories (Wilmington, MA). After arrival, mice were housed singly in filtered-transparent polycarbonate cages in the Wegner Hall Vivarium at Washington State University. Mice were allowed ad libitum access to sterilized Milli-Q water and Purina 5001 Rodent Laboratory Chow. After 1 wk of acclimation, mice were randomly divided into two groups. One group was continuously given Milli-Q water and laboratory chow. The other group was given filter-sterilized 20% (v/v) alcohol (Everclear, St. Louis, MO) and laboratory chow. The animal protocol used in this research was approved by the Institutional Animal Care and Use Committee at Washington State University.

Measurement of daily alcohol intake and blood alcohol concentration

The amount of daily alcohol intake was determined by measuring the starting weight and ending weight of alcohol in the feeding tube every other day. These values were then converted to milliliters by dividing the weight of alcohol consumed by the density of alcohol. The blood alcohol concentration was determined by using the NAD-ADH Reagent Multiple Test Vial kit (Sigma-Aldrich, St. Louis, MO) following the manufacturer’s instructions.

Tumor cell line and tumor cell inoculation

B16BL6 melanoma cells were cultured in DMEM supplemented with 10% FBS, 1% penicillin and streptomycin, and incubated at 37°C with 5% CO2. Cells were harvested when they reached 50–70% confluence and suspended in PBS at 1 × 10^7 cells/ml. After 3–6 mo of alcohol consumption, mice were inoculated s.c. with 2 × 10^6 melanoma cells in the right hip area.

Abs and reagents

The following PE-, FITC-, and PerCP-conjugated anti-mouse Abs were purchased from BD Biosciences (San Diego, CA) and BioLegend (San Diego, CA): anti-CD1d (1D1, B1), anti-CD3 (145-2C11), anti-CD4 (RM4-5), anti-CD5 (5-7.3), anti-CD8 (56-6.7), anti-CD19 (6D6), anti-CD21 (7E9), anti-CD23 (B3B4), anti-CD93 (AA.4.1, NK1.1, PK136), anti-CXCR5 (2G8), anti-CCR7 (4B12), anti-IgD (11-25.2a), and anti-IgM (RMM-1). Recombinant mouse CCL21 and CXCL13 were purchased from PeproTech (Rocky Hill, NJ). S1P was purchased from Avanti Polar Lipids (Alabaster, AL).

Leukocyte isolation

Leukocytes from BM, spleen, LNs, and peripheral blood were isolated following previous procedures (23). The leukocyte number was determined with the aid of the Vi-CELL cell viability analyzer (Beckman Coulter, Brea, CA). Cells were collected by centrifugation and resuspended in an appropriate amount of PBS plus BSA.

Phenotype analysis

The phenotype of leukocytes from different organs was determined by flow cytometry following previously published methods (22). Briefly, 10^6 cells were preincubated with anti-CD16/CD32 mAb (2.4G2) at 4°C for 5 min to block nonspecific Fc-receptor binding. Cells were stained with the appropriate mAb at 4°C for 20 min. After washing with FACS buffer (PBS plus 0.1% BSA plus 0.1% NaN3), cells were analyzed using a Becton Dickinson FACScan Flow Cytometer. Data were collected and analyzed with CellQuest software.

Transwell migration assay

The Transwell migration assay was conducted with the Costar 24-well plate Transwell system using an insert with a diameter of 6.5 mm and pore size of 5 μm. Splenocytes were suspended in serum-free RPMI 1640 medium, prewarmed in an incubator at 37°C for 30 min before conducting the Transwell migration assay, and 10^6 cells were added to the upper chamber of the insert. The lower chamber contained one of four different chemotactic attractants: 125 nM S1P, 500 ng/ml CCL21, 1 μg/ml CXCL13, or 300 μl plasma in a total volume of 600 μl serum-free RPMI 1640 medium. A blank well containing 600 μl serum-free RPMI 1640 medium without added chemotactic agent was used as a control. The plates were incubated at 37°C in a CO2 incubator for 4–5 h and then placed on ice for 10 min before cells in the lower chamber were collected and counted using a Vi-CELL viability analyzer (Beckman Coulter). The migration rate was calculated using the following formula: Migration rate = (number of migrated cells – number of migrated cells in the control) × 100/10^6.

RNA extraction and RT-PCR

Total RNA was extracted from the spleen using TRIzol reagent (Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions. First-strand cDNA was prepared using M-MLV reverse transcriptase (Promega, Madison, WI) and adhering to the manufacturer’s protocol. The following primers were used to amplify S1PR1 and sphingosine-1-phosphate lyase-1 (SPL1) cDNA fragment, respectively. S1PR1 forward primer, 5'-GAGCATTGGTGTCGACCGCTGCT-3'; S1PR1 reverse primer, 5'-ACAGCAGCTCGCTCAAGCC-3'. SPL1 forward primer, 5'-AGCCCGGAGCTCGATCTGCG-3'; SPL1 reverse primer, 5'-GCAGCAGTGACACTCGCG-3'. GAPDH was used as an internal control of gene expression. Two primers, GAPDH1 5'-ACCACGCTCTACCACAC-3' and GAPDH2 5'-TACAGACAGGGTGTTGGA-3', were used to amplify GAPDH. PCR was conducted with 1 cycle of predenaturation at 94°C for 5 min, followed by 30 cycles of denaturation at 94°C for 30 s, annealing at 58°C for 30 s and elongation at 72°C for 40 s, and 1 cycle of final elongation at 72°C for 7 min. PCR products were separated by 2% agarose gel electrophoresis with Tris-Acetate EDTA buffer. The absorbance of the PCR products was analyzed by using the UVP BioImaging Systems and Labworks gel analysis software.

Statistical analysis

The data were analyzed by Microsoft Excel software or GraphPad Prism 5 and expressed as mean ± SD. The two-tailed Student t test was used to analyze differences between the two groups, and differences were considered significant at p < 0.05. Except where specifically indicated, each group contained 7–10 mice.

Results

Daily alcohol intake and blood alcohol concentration

The daily intake of 20% w/v alcohol was 5.6 ± 1.0 ml/day in the nonmelanoma-injected mice and 6.1 ± 1.2 ml/day in the tumor-bearing mice. There was no significant difference in alcohol consumption between nonmelanoma and melanoma-injected mice. The blood alcohol concentration determined 2 wk after melanoma injection was 0.028 ± 0.004%. This is consistent with previous determinations of alcohol blood concentration in mice not injected with melanoma (24, 25).

Chronic alcohol consumption induces lymphopenia in melanoma-bearing mice but not in nontumor-inoculated mice

We previously found that chronic alcohol consumption decreases cellularity in the spleen of nontumor-inoculated mice (22). In this study, we examined if chronic alcohol consumption affected leukocyte numbers in the spleen and blood of melanoma-bearing mice. Results indicated that although the cellularity is still lower in the spleen of melanoma-bearing, alcohol-consuming mice compared with respective water-drinking mice, the difference is not significantly different (Fig. 1A). However, the numbers of PBLs are significantly lower in alcohol-consuming, melanoma-bearing mice than in their water-drinking counterparts (Fig. 1B). We then examined if alcohol consumption decreased PBLs in nonmelanoma-inoculated mice. The results indicated no effect of chronic alcohol consumption on total PBL number (Fig. 1C); however, alcohol consumption decreased the numbers of CD8^+ T cells and NK (CD3^+ NK1.1^+) cells in the blood of these nonmelanoma-inoculated mice (Fig. 1D). The percentage of CD19^+ B cells increased and the percentage of CD8^+ T cells decreased in...
the PBLs (Fig. 1E). These results indicated that chronic alcohol consumption induces lymphopenia in the melanoma-bearing mice but not in mice that are not inoculated with melanoma.

Chronic alcohol consumption selectively decreases B cells and T cells in blood but not in BM, LNs, and spleen

We further determined if the decrease of PBLs in melanoma-bearing, alcohol-consuming mice was associated with a general decline in all types of lymphocytes or due to selective reduction in specific cell types. CD4+ T (CD4+NK1.1−) cells, CD8+ T cells, B (CD19+) cells, NKT (CD3+NK1.1+) cells, and NK (CD3−NK1.1+) cells were examined. The results indicated that chronic alcohol consumption significantly decreased the percentage of B cells (Fig. 2C), increased the percentage of NKT cells (Fig. 2D), and did not significantly alter the percentage of CD4+ T cells (Fig. 2A). The presence of melanoma tumors had varying effects on CD8+ T cells (Fig. 2) and NK cells (Fig. 2E). On day 14, the percentages of CD8+ T cells were significantly lower and NK cells were significantly higher in the alcohol-consuming mice than in water-drinking mice, but not different on days 11 and 17.

Overall, the major effects associated with lymphopenia in melanoma-inoculated mice were observed in B and T (CD4+ and CD8+) cells, where the numbers of these cells were decreased 2- to 4-fold (Fig. 2F–H). The greatest degree of decrease was in...
B cells (Fig. 2H) followed by CD8+ T cells (Fig. 2G) and CD4+ T cells (Fig. 2F).

As a peripheral reservoir of T cells and B cells, the spleen plays an important role in adjusting these lymphocytes in the blood. Alcohol consumption decreases T and B cells in the spleen of nontumor-inoculated mice (22). Therefore, we next examined if these cells were further altered in melanoma-inoculated mice. The results indicated that alcohol consumption had little effect on the percentage of T, B, NK and NKT cells in the spleen (Fig. 3A–E). CD4+ T cells (Fig. 3A) were increased whereas CD8+ T cells (Fig. 3B) were decreased in alcohol-consuming mice, but only on day 11 after melanoma inoculation.

The numbers of CD4+ T cells (Fig. 3F) and NKT cells (Fig. 3I) were not modulated by alcohol consumption and melanoma inoculation. B cell numbers did not significantly change during the early time points after melanoma inoculation; however, they were significantly lower in alcohol-consuming mice on day 17 (Fig. 3H).

Chronic alcohol consumption decreases the numbers of CD4+ T cells and B cells in the spleen, LNs, and BM are largely not affected by alcohol consumption and melanoma inoculation. However, alcohol consumption and melanoma inoculation decrease the numbers of CD8+ T cells in the spleen and LNs.

**FIGURE 3.** Effects of chronic alcohol consumption on different types of lymphocytes in the spleen of B16BL6 melanoma-bearing mice. (A–E) Percentage of the indicated lymphocytes in the splenocytes of melanoma-bearing mice at the indicated days after s.c. tumor inoculation. (F–J) Number of the indicated lymphocytes in the splenocytes of the water-drinking and alcohol-consuming mice at the indicated days after s.c. tumor inoculation. Each group contained 7–10 mice. Experiments were repeated once with similar results. *p < 0.05 (water-drinking group different from alcohol-consuming group on the indicated day). ETOH. Alcohol-consuming, melanoma-bearing mice; Water, water-drinking, melanoma-bearing mice.
CD1d+CD5+ regulatory B cells were significantly lower in the alcohol-consuming, melanoma-bearing mice (Fig. 6E). Collectively, these results indicate that the decrease of T and B cells in the blood is not caused by apoptosis. Chronic alcohol consumption does not compromise the chemotactic ability of plasma from melanoma-bearing mice to facilitate splenocyte migration.

Because the cell number and differentiation of B cells in the BM and spleen were normal in the alcohol-consuming, melanoma-bearing mice, the decrease of B cells in the blood is not caused by apoptosis. We also found that the decrease in peripheral blood B cells is mainly found in the mature B cell portion. Therefore, the decrease in the PBLs is likely due to an inability of splenic T and B cells to migrate. One possibility is that the blood is deficient in chemotactic factors to attract T cells and B cells from spleen. To examine this possibility, we used a Transwell cell migration system to test the chemotactic ability of plasma from alcohol-consuming, melanoma-bearing mice and water-drinking, melanoma-bearing mice to facilitate splenocyte migration in vitro. The results indicated that there was no significant difference in migration of splenocytes indicating that neither alcohol nor melanoma affects the chemotactic ability of plasma to attract splenocytes (Fig. 8).

Chronic alcohol consumption does not impair the response of splenocytes from melanoma-bearing mice to CCL21 and CXCL13.

Given that the chemotactic ability of plasma is intact in alcohol-consuming, melanoma-bearing mice, it is possible that the inability of splenic T cells and B cells to enter the blood could result from a defect in the cells themselves to respond to chemotactic factors. It is well established that the most important chemokines and receptors that control T cell migration and homing are CCL21, CXCL13, and their receptor, CCR7 (26, 27). The chemokine and its receptor that are involved in B cell migration and redistribution in the follicle of the spleen is CXCL13 and CXCR5 (9). Therefore, we examined if alcohol consumption impaired the expression of the chemokine receptors CCR7 and CXCR5 on splenic T cells and B cells, respectively. We found that chronic alcohol consumption did not alter the expression of these receptors on splenic T and B cells from melanoma-bearing mice (Fig. 9A–D). Consistent with chemokine receptor expression, the ability of splenocytes to respond to CCL21 and CXCL13 was also not altered in the alcohol-consuming, melanoma-bearing mice (Fig. 9E).

Chronic alcohol consumption does not induce T cell and B cell apoptosis in the blood.

One factor that could explain the decreased number of T and B cells in the blood is loss of these cells via apoptosis facilitated by melanoma and alcohol consumption. To test this hypothesis, we determined the percentage of annexin V+ T and B cells in the blood. The results indicated that the percentage of annexin V+ cells in the CD8+ T cells and CD19+ B cells was not significantly different between the alcohol-consuming, melanoma-bearing mice and water-drinking, melanoma-bearing mice (Fig. 7A, 7B). These results indicate that the decrease of T and B cells in the blood of alcohol-consuming, melanoma-bearing mice is not caused by apoptosis.

Chronic alcohol consumption decreases MZ B cells and CD1d+CD5+ B cells and downregulates CD1d expression on MZ B cells in melanoma-bearing mice.

Alcohol consumption did not affect the numbers of follicular B cells in the spleen of melanoma-bearing mice. We next examined if MZ B cells and regulatory B cells were affected. We found that the percentages and numbers of MZ B cells (CD21hiCD1dhi) and CD1d+CD5+ regulatory B cells were significantly lower in the alcohol-consuming, melanoma-bearing mice than in the respective water-drinking counterparts (Fig. 6A–D). Although the overall numbers of MZ B cells were not significantly different between the alcohol-consuming and water-drinking groups, the expression of CD1d on these cells was significantly downregulated in the alcohol-consuming, melanoma-bearing mice (Fig. 6E).
mice (data not shown). S1P levels in the blood of mice given the alcohol-containing Leiber–DeCarli liquid diet are also not altered by chronic alcohol consumption (29). Therefore, the ability of splenocytes to respond to S1P chemotaxis could be compromised by alcohol consumption in melanoma-bearing mice. To examine this possibility, we determined migration of splenocytes from water-drinking and alcohol-consuming, melanoma-bearing mice in response to S1P. We found that chronic alcohol consumption significantly decreased S1P-induced splenocyte migration (Fig. 10A). S1PR1 controls the response of T and B cells to S1P. Cell surface S1PR1 is controlled by S1PR1 gene expression and also by S1P concentration in the microenvironment. High concentrations of S1P will induce S1PR1 internalization (30). We used RT-PCR to examine the gene expression of S1PR1 in splenocytes from water-drinking and alcohol-consuming, melanoma-bearing mice. Expression of S1PR1 was significantly downregulated in splenocytes from alcohol-consuming, melanoma-bearing mice compared with their water-drinking counterparts (Fig. 10B). Gene expression of SPL1, an enzyme catalyzing the degradation of S1P to hexadecanaldehyde and phosphoethanolamine, was also downregulated in the alcohol-consuming, melanoma-bearing mice (Fig. 10C). These data indicate that alcohol consumption compromises the splenocyte response to S1P via decreasing S1PR1 expression on splenocytes.

Discussion
The data demonstrate that alcohol consumption differentially affects lymphocyte distribution in the spleen and blood of normal and melanoma-bearing mice. Alcohol consumption in the nontumor-inoculated mice results in a general decrease in lymphocytes in the spleen (22) but does not alter PBLs except for CD8+ T cells, which are decreased in both spleen and blood (Fig. 1). In the melanoma-bearing mice, however, alcohol consumption dramatically decreases PBLs, but does not significantly decrease lymphocytes in the spleen except for CD8+ T cells (Figs. 2, 3). The effects of alcohol consumption on each type of lymphocyte in the blood of melanoma-bearing mice were different. Alcohol does not significantly affect NK cells, increases NKT cells, only moderately decreases CD4+NK1.1+ T cells, and dramatically decreases B cells and CD8+ T cells. Compared to B cells, CD8+ T cells are a small percentage of the total PBL population. B cells compose ∼45% of the PBLs; thus, the decrease in B cells contributes more profoundly to the lymphopenia in the alcohol-consuming, melanoma-bearing mice.

B cells circulate in the blood, and a decrease could result from the defective production and/or release of these cells from the BM or from impaired egress of these cells from LNs and spleen. Because alcohol consumption neither alters the total numbers of B cells in the BM (Fig. 4B) nor significantly affects the development of B cells in the BM (Fig. 5B), the decreases in circulating B cells in the blood are not due to compromised B cell development, BM production, or the release from BM. Thus, the decreases of B cells in the blood highly likely result from compromised circulation of these cells among the secondary lymphoid organs such as LNs and spleen.

The moderate increase in CD93+CD23− cells suggests that alcohol consumption might impair B cell maturation from the transitional stage into mature follicular B cells or MZ B cells. However, the increase in the percentage of CD23+ B cells (Fig. 5G) suggests that follicular B cells accumulate abnormally in the spleen. If the follicular B cells fail to circulate in the blood and accumulate in the spleen, then the percentage of CD23+ follicular

FIGURE 5. Effects of chronic alcohol consumption on B cell development and distribution of B cell subsets of in BM, blood, and spleen of melanoma-bearing mice at 2 wk after s.c. tumor inoculation. (A) Dot plot of IgMlo IgDlo B cells (R4), IgMhi IgDlo B cells (R5), and IgMhi IgDhi B cells (R3) in gated BM CD19+ B cells. (B) Percentage of indicated B cell subsets in the BM CD19+ B cells. (C) Dot plot indicating IgMlo IgDlo immature B cells in the gated CD19+ peripheral blood B cells. (D) Percentage of IgMlo IgDlo immature B cells in CD19+ peripheral blood B cells. (E) Dot plot indicating CD23 and CD93 defined subsets of B cells in gated splenic CD19+ B cells. (F) Percentage of indicated subsets of B cells in splenic CD19+ B cells. (G) Percentage of CD23+CD19+ B cells in the indicated organs. (H) Number of CD23+CD19+ B cells in the indicated organs. Each group contained 7–10 mice. Experiments were repeated once with similar results. *p < 0.05, **p < 0.01, ***p < 0.001 (ETOH group different from water group). ETOH, Alcohol-consuming, melanoma-bearing mice; Water, water-drinking, melanoma-bearing mice.
B cells will increase and the percentage of MZ B cells, which are noncirculating cells, will decrease in the splenic B cell population as we observed (Fig. 6). Because the BM exports immature B cells into the blood, the majority of mature B cells in the blood come from secondary lymphoid organs. If the production and release of B cells from the BM is normal and the egress of B cells from spleen is compromised, then the portion of immature B cells in blood would be expected to increase, which is also what we showed (Fig. 5D, 5G, 5H). Therefore, these data indicate an alcohol–melanoma interaction that influences the number and maturity of B cells in the blood.

If the alcohol–melanoma interaction impairs B cell egress from the spleen, then the number of B cells should be elevated in the spleen; however, we did not observe this (Fig. 3H). Alcohol consumption significantly decreases B cells in the spleen of nontumor-injected mice (22); however, the effect is diminished in melanoma-bearing mice where no significant decreases are observed until day 17 after tumor inoculation. Thus, it is likely that the impaired egress of B cells is obscured by the more profound effect of alcohol consumption on B cell numbers before melanoma inoculation. It is worthy to note that alcohol consumption downregulates the expression of CD1d on MZ B cells. CD1d is a nonclassical MHC class I-like molecule that presents lipid Ag to NKT cells and plays an important role in NKT cell activation. MZ B cells are involved in NKT cell activation and the regulation of cytokine production (6, 8). Alcohol consumption increases NKT cells in the blood of melanoma-bearing mice (Fig. 2D, 2I). Whether the decrease in MZ B cells and the downregulation of these cells affects NKT cell function in the alcohol-consuming, melanoma-bearing mice is an interesting question that remains to be elucidated.

The current study indicates that the interaction between alcohol and melanoma does not affect the chemotactic capacity of plasma to induce migration of T and B cells nor does the interaction alter the expression of CXCR5 and CCR7 on these cells. Alcohol consumption downregulates the expression of S1PR1 and SPL1 in splenocytes from melanoma-bearing mice. Therefore, the impaired egress of T cells and B cells from the spleen is associated with a compromised S1PR1 signaling pathway. This is evidenced not only by the decreased response of splenocytes from alcohol-consuming, melanoma-bearing mice to S1P-induced migration.

**FIGURE 6.** Effects of chronic alcohol consumption on MZ B cells and CD5+CD1dhi regulatory B cells in the spleen of melanoma-bearing mice at 2 wk after s.c. melanoma inoculation. (A) Dot plot indicating CD21hi CD1dhi MZ B cells (R1) in gated splenic CD19hi B cells. (B) Contour plot indicating CD5+CD1dhi regulatory B cells in gated splenic CD19hi B cells. (C) Percentage of MZ B cells (CD21hiCD1dhi) and regulatory B cells (CD5+CD1dhi) in splenic CD19hi B cells. (D) Number of MZ B cells (CD21hiCD1dhi) and regulatory B cells (CD5+CD1dhi) in the spleen. (E) Mean fluorescence intensity of CD1d in CD21hiCD1dhi MZ B cells. Each group contained 7–10 mice. Experiments were repeated once with similar results. *p < 0.05, **p < 0.01 (ETOH group different from water group). ETOH, Alcohol-consuming, melanoma-bearing mice; Water, water-drinking, melanoma-bearing mice.

**FIGURE 7.** Effects of chronic alcohol consumption on the apoptosis of CD8hi T cells and CD19hi B cells in the blood of tumor-bearing mice. Mice were inoculated with B16BL6 melanoma s.c. and euthanized 2 wk after tumor inoculation. PBLs were stained with anti-mouse CD19–PerCP, anti-mouse CD8–PE, and annexin V–FITC. (A) Dot plot indicating annexin V+ cells in CD19hi B cells. (B) Percentage of annexin V+ cells in CD8hi T cells and CD19hi B cells. Each group contained 7–10 mice. ETOH, Alcohol-consuming mice; Water, water-drinking mice.

**FIGURE 8.** Effects of chronic alcohol consumption on chemotactic capacity of plasma in melanoma-bearing mice. Plasma was collected 2 wk after s.c. melanoma inoculation. The chemotactic capacity of plasma was examined in a Transwell migration experiment as detailed in Materials and Methods. The figure shows the percentage of migrated cells compared with the total number of seeded cells. The bars of the graph represent the mean ± SD from six individual migration experiments conducted for six mice from each group. ETOH, Alcohol-consuming, melanoma-bearing mice; Water, water-drinking, melanoma-bearing mice.

B cells from the BM is normal and the egress of B cells from spleen is compromised, then the portion of immature B cells in blood would be expected to increase, which is also what we showed (Fig. 5D, 5G, 5H). Therefore, these data indicate an alcohol–melanoma interaction that influences the number and maturity of B cells in the blood.

If the alcohol–melanoma interaction impairs B cell egress from the spleen, then the number of B cells should be elevated in the spleen; however, we did not observe this (Fig. 3H). Alcohol consumption significantly decreases B cells in the spleen of nontumor-injected mice (22); however, the effect is diminished in melanoma-bearing mice where no significant decreases are observed until day 17 after tumor inoculation. Thus, it is likely that the impaired egress of B cells is obscured by the more profound effect of alcohol consumption on B cell numbers before melanoma inoculation. It is worthy to note that alcohol consumption downregulates the expression of CD1d on MZ B cells. CD1d is a nonclassical MHC class I-like molecule that presents lipid Ag to NKT cells and plays an important role in NKT cell activation. MZ B cells are involved in NKT cell activation and the regulation of cytokine production (6, 8). Alcohol consumption increases NKT cells in the blood of melanoma-bearing mice (Fig. 2D, 2I). Whether the decrease in MZ B cells and the downregulation of CD1d on these cells affects NKT cell function in the alcohol-consuming, melanoma-bearing mice is an interesting question that remains to be elucidated.

The current study indicates that the interaction between alcohol and melanoma does not affect the chemotactic capacity of plasma to induce migration of T and B cells nor does the interaction alter the expression of CXCR5 and CCR7 on these cells. Alcohol consumption downregulates the expression of S1PR1 and SPL1 in splenocytes from melanoma-bearing mice. Therefore, the impaired egress of T cells and B cells from the spleen is associated with a compromised S1PR1 signaling pathway. This is evidenced not only by the decreased response of splenocytes from alcohol-consuming, melanoma-bearing mice to S1P-induced migration.
but also by the fact that only B and T cells and not NKT and NK cells decrease in the blood. It is known that activation of the S1P–S1PR1 signaling pathway affects the cytokine production of NKT cells but does not affect the egress and migration of these cells (31). NK cell egress from BM and LNs is regulated by the S1P–S1PR5 pathway and not the S1P–S1PR1 pathway (32, 33).

The egress of B cells from the BM is also controlled by the S1P–S1PR1 pathway (28). We found that alcohol consumption did not impair B cell egress from BM of melanoma-bearing mice. This suggests that the expression of S1PR1 in BM-derived B cells is not affected, which is consistent with our observations that alcohol consumption leads to downregulation of S1PR1 expression in spleen but not in BM (data not shown). This further indicates an organ-specific effect associated with alcohol consumption in the melanoma-bearing mice. The cell surface expression of S1PR1 is regulated by the s1pr1 gene and also regulated by the concentration of S1P in the microenvironment (34). We found that this gene is downregulated in the spleen of alcohol-consuming, melanoma-bearing mice. We also found that the expression of SPL1 is downregulated in the spleen. SPL1 catalyzes the degradation of S1P. The decrease of SPL1 could result in accumulation of S1P. The elevation of S1P will induce the internalization of S1PR1, which in turn could compromise the response to S1P-induced chemotaxis. One of the products of S1P degradation catalyzed by SPL1 is phosphoethanolamine. Chronic alcohol consumption alters the metabolism of lipids, including phosphoethanolamine (35, 36). It is likely that alcohol consumption interacting with tumor cells changes the metabolism of lipids, which lead to organ specific alternations in S1P and S1PR1. We are investigating this hypothesis and the mechanism of how alcohol consumption interacts with melanoma to regulate S1PR1 expression.

The effects of B cells in tumor immunity have attracted more attention in recent years. Some reports indicate that depletion of B cells enhances anti-tumor immunity, implicating B cells in the suppression of anti-tumor immune responses (14, 15). However, in B16 melanoma, the lack of B cells, especially mature B cells, compromises T cell function and enhances tumor progression (18). Also in B16 melanoma, depletion of CD20+ mature B cells increases lung tumor growth and burden in CpG-treated mice (20). It is also reported that the high density of infiltrated B cells in melanoma correlates with a significant survival advantage and positive prognostic outcome in melanoma patients (37). Understanding the relationship among impaired B cell circulation, dysfunction of CD8+ T cells, and decreased survival in alcohol-consuming melanoma-bearing mice will not only advance our knowledge on the effects of B cells in anti-tumor immunity but also provide new insights into how to recover immune function in alcoholics with melanoma and possibly other cancers. Research in this area is ongoing.

In summary, we conclude that chronic alcohol consumption compromises the egress of B and T cells from the spleen via impairment of the S1P–S1PR1 pathway, which in turn induces severe lymphopenia in B16BL6 melanoma-bearing mice. Alcohol consumption neither compromises the development nor affects the egress of B cells from BM. The compromised egress of B cells from the spleen results in an immature B cell-dominant circulating population in the blood of melanoma-bearing mice.
DiLillo, D. J., K. Yanaba, and T. F. Tedder. 2010. B cells are required for optimal


Zhang, H., and G. G. Meadows. 2005. Chronic alcohol consumption in mice increases the proportion of peripheral memory T cells by homeostatic prolifer-


Gunn, M. D., S. Kyuwa, C. Tam, K. Kakiuchi, A. Matsuzawa, L. T. Williams, and H. Nakano. 1999. Mice lacking expression of secondary lymphoid organ che-


