CD40-stimulated B Lymphocytes Pulsed with Tumor Antigens Are Effective Antigen-presenting Cells That Can Generate Specific T Cells

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ABSTRACT

Although they are considered as antigen-presenting cells, the role of antigen-unspecific B lymphocytes in antigen presentation and T-lymphocyte stimulation remains controversial. In this paper, we tested the capacity of normal human peripheral activated B cells to stimulate T cells using melanoma antigens or melanoma cell lysates. B lymphocytes activated through CD40 ligation and then pulsed with tumor antigens efficiently processed and presented MHC class II-restricted peptides to specific CD4+ T-cell clones. This suggests that CD40-activated B cells have the functional and molecular competence to present MHC class II epitopes when pulsed with exogenous antigens, thereby making them a relevant source of antigen-presenting cells to generate T cells. To test this hypothesis, CD40-activated B cells were pulsed with a lysate prepared from melanoma cells and used to stimulate peripheral autologous T cells. Interestingly, T cells specific to melanoma antigens were generated. Additional analysis of these T-cell clones revealed that they recognized MHC class II-restricted epitopes from tyrosinase, a known melanoma tumor antigen. The efficient antigen presentation by antigen-unspecific activated B cells was correlated with a down-regulation in the expression of HLA-DO, a B cell-specific protein known to interfere with HLA-DM function. Because HLA-DM is important in MHC class II peptide loading, the observed decrease in HLA-DO may partially explain the enhanced antigen presentation after B-cell activation. Results globally suggest that when they are properly activated, antigen-unspecific B-lymphocytes can present exogenous antigens by MHC class II molecules and stimulate peripheral antigen-specific T cells. Antigen presentation by activated B cells could be exploited for immunotherapy by allowing the in vitro generation of T cells specific against antigens expressed by tumors or viruses.

INTRODUCTION

B lymphocytes have the capacity to present exogenous proteins to T cells after antigen-specific internalization by their surface immunoglobulins. This MHC class II-restricted presentation is an important step in the establishment of the humoral response (1, 2). In addition, B cells can endocytose and present antigens in an immunoglobulin-independent fashion. For example, EBV-transformed B cells have the capacity to present tumor antigens by MHC class II (3, 4). In addition, both DCs and B lymphocytes from spleen can present a peptide derived from hen egg lysozyme by MHC class II after the injection of the protein (5). However, the in vivo consequence of antigen processing by unspecific B cells remains elusive, and there is conflicting evidence whether these B cells interact with T cells in a tolerogenic or immunogenic fashion (6–10).

Previous studies have shown that in some conditions, resting B cells can be tolerogenic (7, 8, 10, 11). Activation signals may be required to condition unspecific B lymphocytes to become competent APCs for T-cell activation and antigen presentation. Stimulatory signals can be provided to B cells from different sources, such as from activated T cells expressing CD40L and secreting cytokines such as IL-4. CD40-activation of B cells increases different aspects of APC function (7, 10, 11). Specifically, activation of a B-cell line by the CD40 receptor increased MHC class II-restricted antigen presentation (12). The reason for enhanced T-cell recognition of activated B cells is not clear, but may be a function of altered expression of proteins involved in MHC class II antigen processing and presentation.

Antigen presentation by the MHC class II pathway is dependent on the involvement of several key proteins implicated in MHC class II maturation and sorting, as well as proteins favoring the loading of stable peptides. Among them, the invariant chain is known to associate with the newly synthesized MHC class II molecules in the endoplasmic reticulum to chaperone transport to the MIIC (13). In MIICs, the invariant chain is degraded leaving a small fragment called CLIP in the peptide binding site of the class II molecules (14). An accessory molecule, HLA-DM, catalyzes the removal of the CLIP fragment and peptides having low affinity to MHC molecules, which promotes the loading of peptides with higher stability (15–17). Although HLA-DM is expressed in all of the APCs, B lymphocytes also express HLA-DO, another nonclassical MHC accessory molecule (18). By interacting with HLA-DM (19) and possibly impairing its function, HLA-DO modifies the peptide repertoire associated with surface MHC class II molecules in favor of low affinity peptides (20, 21). Indeed, HLA-DO was postulated to indirectly favor presentation of antigens taken up by the surface immunoglobulins by inhibiting class II loading of nonspecific proteins in early compartments of the endocytic pathway. Because HLA-DO changes the peptide/MHC class II repertoire, its presence or absence could impact the B- and T-lymphocyte interaction.

In this study, antigen-unspecific B lymphocytes from peripheral blood lymphocytes of human donors were stimulated with CD40L and IL-4 in a primary cell culture system, and analyzed for their ability to present exogenous antigens by MHC class II. Data presented suggest that such stimulation of B cells increased the presentation of exogenous antigens by MHC class II molecules. Furthermore, B-cell activation provoked a consistent decrease in the expression of HLA-DO.

As a source of antigen, we used lysates from tumor cells, which are known to contain a number of weakly immunogenic antigens, to stringently test the immunogenicity of these B cells. Importantly, CD40-activated B cells pulsed with melanoma lysates had the capacity to generate antigen-specific CD4+ T cells from resting PBMCs. This data suggests that when properly activated, antigen-unspecific B cells can be efficient APCs.

MATERIALS AND METHODS

Cell Culture. Immortalized EBV-B cells from patient 1088, and melanoma cell lines 1088mel, 888mel, and 553mel were prepared as described previously (4). The melanoma line SK23mel and the breast cancer line MDA231 were obtained from the American Type Culture Collection (Manassas, VA). EBV-B
B lymphocytes and some of the T cells were cultured in complete medium consisting of Iscove’s Modified Dulbecco’s medium (Iscove) supplemented with 10% human AB serum (male, heat inactivated; Gemini Bio-Products, Calabasas, CA), 1 mM glutamine, 50 units/ml penicillin, 50 μg/ml streptomycin (all from Biofluids Inc., Rockville, MD), and 50 μg/ml gentamicin (Invitrogen). B lymphocytes and some of the T cells were cultured in complete medium consisting of Iscove’s Modified Dulbecco’s medium (Iscove) supplemented with 10% human AB serum (male, heat inactivated; Gemini Bio-Products, Calabasas, CA), 1 mM glutamine, 50 units/ml penicillin, 50 μg/ml streptomycin, and 50 μg/ml gentamicin (all from Invitrogen).

To generate CD40-activated B-cell cultures, purified B cells (using a negative selection system; StemCell Technologies Inc., Vancouver, British Columbia, Canada) or bulk PBMCs were cultured with either 500 ng/ml of a soluble trimeric CD40L (soluble trimeric CD40L; Immunex Corporation, Seattle, WA). Cells were cultured in complete medium complemented with 200 units/ml of recombinant human IL-4 (Peprotech, Rocky Hill, NJ). When using the StemCell enrichment system, we routinely obtained >95% B lymphocytes with no detectable T cells or monocytes. Fresh complete medium was added again on day 3 with IL-4 (and 500 ng/ml soluble trimeric CD40L if used). After the first round of proliferation (days 5–8), cells were either frozen for future use or restimulated every 2–3 days when the culture reached a density of 1.5–2 × 10^6 cells/ml. They were replated at about 3–5 × 10^5 cells/ml of medium (containing 500 ng/ml of CD40L) and 200 units/ml IL-4.

The anti-gp100 clone was prepared by limiting dilution of a bulk T-cell culture after 2 stimulations with gp100-transduced CD34-derived DCs, and were cultured and characterized as described previously (22).

**Phenotypic Analysis and MLR.** Fluorescence-activated cell sorter analyses were performed using phycoerythrin- or FITC-labeled antibodies specific to human CD3, CD14, CD19, CD20, CD21, CD40, CD80, CD86, and HLA-DR (all from Becton-Dickinson, San Jose, CA), CD83 (Immunotech, Marseille, France), or isotype-match controls (Becton Dickinson). Labeled cells were collected and analyzed using a FACScan cytometer (Becton Dickinson), and analysis was done with CellQuest software (Becton Dickinson). Propidium Iodine was used to exclude dead cells from analysis.

MLR was performed as described previously (22). Briefly, 1 × 10^5 allogeneic-enriched T cells (human T-cell immunoaffinity columns; R&D Inc., Minneapolis, MN) were cocultured with increasing numbers of irradiated (1500 rads) PBMCs or CD40 and IL-4-activated B cells prepared from the same donor in 96-well flat-bottomed tissue culture plates for 6 days. T-cell proliferation was monitored by [3H]thymidine (DuPont New England Nuclear, Boston, MA) incorporation for the last 16 h. Results were corrected for [3H]thymidine incorporation by irradiated CD40−B cells and T cells alone.

**Pulsing of CD40-activated B Cells and Recognition by CD4+ T-Cell Clones.** Cell lysates were prepared by five successive freeze/thaw cycles of 1 × 10^7 cells/ml in Iscove’s base medium containing 10% human AB serum. 10^5 autologous B cells were plated in 2 × 10^5 cells/well in 96-well plates at 100 μl or 1–2 × 10^5 cells/well in 24-well plates in 500 μl in B-cell medium without IL-4. Lysates were added for 16–20 h to reach a ratio of B cell:lysed cell of 1:1 to 1:0.5 depending on the toxicity of the lysate on B-cell viability as determined by trypan blue exclusion. For a recognition assay in 96-well plates, T cells were added for an additional 24 h. For a T-cell stimulation assay, pulsed B cells from 24-well plates were harvested and irradiated at 15 Gy.

In some experiments, chloroquine (Sigma, St. Louis, MO) was used at 100 μM for 4 h before antigen pulsing on B cells. Cells were then washed and cultured for 20 h in the presence of 5 μM of chloroquine with or without gp100. Cells were washed once and fixed with 0.5% of formaldehyde for 5 min. Cells were then washed extensively three times and cocultured with T cells in a 96-well flat-bottomed assay.

**Western Blots.** B lymphocytes were purified from PBMCs using a negative selection system (StemCell Technologies Inc.). B cells were >95% pure based on flow cytometry analysis. Cells were left untreated or stimulated with 500 ng/ml of CD40L and 200 units/ml of IL-4 in B-cell culture medium for a different period of time as indicated in “Results.” Protein extracts were prepared as described previously (23). The amount of protein in each sample was determined by the Bradford method to ensure even loading on the gels. Samples were boiled for 5 min under nonreducing conditions in SDS loading buffer and loaded on 10% SDS-PAGE. Proteins were transferred on nitrocellulose membranes (Amersham Pharmacia Biotech, Buckinghamshire, England) and saturated in tris buffer saline-tween 0.1% (TT133)-milk for 1 h as recommended by the manufacturer. Antibodies were incubated overnight at 4°C. Polyclonal antibodies were first incubated for 1 h on a blank membrane in TTBS-milk to remove background. After three washes in TTBS, a peroxi- dase-coupled antibody was added for 1 h. After three more washes, the peroxidase substrate [2,4,5-trichloro-3-(2-methoxy-4-nitro)benzene (TMB); Roche Diagnostics Corporation, Indianapolis, IN] was added, and the membranes were exposed on Fuji Medical X-ray films (Fuji Film Co., LTD, Tokyo, Japan).

Membranes were washed using the ReBlot-Western blot recycling kit (Chemicon International Inc., Temecula, CA) and reprobed using an actin-specific antibody to ascertain equal loading of proteins. The films were analyzed with Gel Pro Analyser (1993–97; Media Cybernetics, Silver Spring, MD). This software measures gray densities of protein spots found on a scanned film. Values of stimulated cells were normalized compared with those of actin. These normalized values were compared with gross values of non-stimulated cells [Normalized value = (actin value of nontreated cells/actin values of treated cells) × antigen presentation molecule values]. Ratios were then calculated with relevant combinations of molecules to observe expression differences after cell activation. Nonstimulated cell ratios were adjusted to 1, and stimulated cell ratios were calculated accordingly so they could be compared with the nonstimulated ratios.

**Antigens.** DA6.147 is a monoclonal antibody (IgG1) directed against the cytoplasmic tail of the HLA-DRα chain (24). The rabbit antisera against the cytoplasmic tails of HLA-DQα or HLA-DMβ have been described previously (25). Mouse antiavidin is a monoclonal antibody that recognizes the NH2-terminal two-thirds of the cellular actin molecules (Chemicon International, Inc.). Secondary peroxidase-coupled antibodies are directed against the Fe part of mouse immunoglobulins (used with primary monoclonal antibodies) or heavy and light chain of rabbit immunoglobulins (used with primary polyclonal antibodies; Jackson ImmunoResearch Laboratories, Inc., Mississauga, Ontario, Canada).

**Stimulation of Autologous T Lymphocytes with Pulsed CD40-activated B Cells.** In the first T-cell generation experiment, the T-cell donor was a melanoma patient from the Surgery Branch (National Cancer Institute) who previously received two i.m. injections of Vaccinia-gp100 and one injection of Fowlpox-gp100 on Institutional Review Board-approved protocols. Irradiated CD40-activated B cells (1 × 10^5) pulsed with a lysate prepared from the melanoma line 1088mel were cultured with 1 × 10^5 allogeneic PBMCs in a 96-well plate in complete Iscove’s base medium containing 10% human AB serum. Cell cultures were restimulated with 1 × 10^6 1088mel lysate-pulsed CD40-activated B cells per well on day 11. IL-2 (Chiron, Emeryville, CA) was added on day 13 and again every 3–4 days at 600 IU/ml. Individual wells were assayed by coculture with 1 × 10^5 CD40-activated B cells pulsed with lysates prepared from 1088mel or a control cell line MDA251. After 16–24 h, GM-CSF secretion was evaluated by ELISA from culture supernatants using coupled antibody pairs from (Endogen, Woburn, MA) using the protocol provided by the manufacturer. Streptavidin-peroxidase (Research Diagnostic Inc., Flanders, NJ) was added at a dilution of 1:4000 followed by peroxidase substrate (Dako, Carpinteria, CA).

In the second T-cell generation study, the T-cell donor was a melanoma patient from the Surgery Branch (National Cancer Institute), and cells were obtained before any immunizations. Irradiated washed CD40-activated B cells (2 × 10^5) pulsed with a lysate prepared from 1088mel were cocultured with purified CD4+ T cells (5 × 10^5; human T-cell immunoaffinity columns; R&D Inc.) in complete Iscove’s base medium containing 10% human AB serum in 1 well of a 24-well plate. Cultures were restimulated with 1088mel lysate-activated B cells on day 11, and IL-2 was added on day 13 and again every 3–4 days at 150 IU/ml. Cultures were restimulated again on day 17 using 1088mel lysate-pulsed B cells. Culture cells were cloned by limiting dilution on day 28 as described previously (22, 26). Clones were cultured in complete medium using AIM-V (Invitrogen) supplemented with 5% human AB serum, 1 mM glutamine, 50 units/ml penicillin, 50 μg/ml streptomycin, 50 μg/ml gentamicin, and 300 IU/ml of IL-2.

All of the recognition assays were carried out in 96-well plates (flat-bottomed) by coculture of 1 × 10^5 target cells and 1 × 10^4 responder T cells for 20–24 h in 200–250 μl. Supernatants were harvested, and IFN-γ or GM-CSF were assayed by ELISA using coupled antibody pairs from Endogen.
RESULTS

B-Cell Presentation of an Exogenous Antigen by MHC Class II.
B lymphocytes were first enriched from PBMCs prepared from normal donors. B-lymphocyte purity and phenotype were additionally confirmed by flow cytometry analyses. After the negative selection enrichment procedure, >95% of the cells expressed B-cell markers such as CD19, CD20, and CD21 (Fig. 1A), which were additionally confirmed by the absence of markers for T cells or monocytes (Fig. 1B). Before stimulation, purified B cells showed the following phenotype: CD19+/CD20+/CD21+ (B-cell markers), CD3− (T-cell markers), CD14− (monocyte/macrophage markers), CD40−, CD40L−, CD80low, CD86low, and CD83+ (APC markers). After CD40L and IL-4 stimulation, the phenotype was CD19+/CD20low/CD21low (B cell markers), CD3− (T cell markers), CD14− (monocyte/macrophage markers), HLA-DR+, CD40low, CD80low, CD86+, and CD83+ (APC markers). Importantly, CD40L/IL-4 stimulation resulted in the generation of a highly homogeneous population of CD19+/CD20+ B cells.

CD40L and IL-4-stimulated B lymphocytes were efficient in allogeneic T-cell stimulation in MLR as presented in Fig. 1D, consistent with a previous report (27).

The capacity of activated human primary B cells to present an exogenous tumor antigen by MHC class II molecules was then evaluated. B cells enriched from PBMCs were stimulated with CD40L and IL-4 individually or in combination. At a variety of time points, activated B cells were pulsed for 18–24 h with recombinant gp100, a melanoma antigen. Class II-restricted presentation of gp100 was assessed using a T-cell clone specific for an HLA-DRβ1*0701 gp100-epitope (22). Purified B cells pulsed with antigen on day 0 that were either nonstimulated (NS d0) or stimulated with IL-4 alone (IL-4 d0) were weakly recognized by the gp100-specific CD4+ T-cell clone (Fig. 2A). However, the recognition was enhanced when gp100-pulsed B cells were stimulated with CD40L alone (CD40L d0) or a combination of CD40L and IL-4 (CD40L/IL-4 d0). The recognition of B cells cultured with CD40L and IL-4 for 5 days before gp100 antigen exposure was better compared with any other condition. These observations suggest that short-term stimulation with CD40L is sufficient to increase the capacity of B cells to present exogenous antigens by MHC class II.

In a second experiment, purified B cells cultured with CD40L or a combination of CD40L and IL-4, pulsed with gp100 antigen 2 days later were efficiently recognized by gp100-specific CD4+ T cells (Fig. 2B; CD40L d2 and CD40L/IL-4 d2). The recognition was specific to gp100 since CD40L/IL-4-stimulated B cells were not recognized when pulsed with the control protein NY-ESO-1 (Fig. 2, A and B), which is a recombinant protein prepared similarly to gp100. Again, gp100-pulsed B cells that were unstimulated or cultured in IL-4 alone were not efficiently recognized. Also, no recognition was observed when HLA-DRβ1*0701 CD40-activated B cells were pulsed with gp100 (Fig. 2B), confirming the specificity of the MHC restriction element. B lymphocytes stimulated with either CD40L or IL-4 alone were not cultured for >3 days because of lack of proliferation.

Finally, to determine whether intracellular antigen processing was required for presentation by MHC class II, CD40L/IL-4-activated B cells were first treated with chloroquine, which inhibits the processing of exogenous antigen and presentation by MHC class II by neutralizing the pH of endosomes. As presented in Fig. 2C (left section), chloroquine treatment resulted in inhibition of MHC class II presentation of exogenous gp100, suggesting that intracellular processing was required for MHC class II presentation. Importantly, this inhibition was not because of chloroquine toxicity because similar treatments of tumor cell lines did not inhibit MHC class I-mediated presentation of endogenous gp100 when using a CD8+ T-cell clone specific for an HLA-A*0201 epitope.

These observations suggest that activation of human B cells with CD40L enhances the capacity to present an exogenous tumor antigen by MHC class II, and that intracellular antigen processing is required. MHC class II-restricted presentation of exogenous antigens was effective in CD40-activated B cells cultured for >16 days (data not shown). The reason for enhanced antigen presentation after B-cell activation is not clear, but may involve a differential expression of molecules involved in MHC class II antigen processing.

CD40-activated B cells pulsed with tumor lysates as APC
Differential Expression of HLA-DR, -DM, and -DO Correlates with Increased Antigen Presentation Activity. The above results show that on activation, antigen-unspecific B cells can present soluble antigens to T cells. This could be caused by multifactorial events such as increased expression of costimulatory molecules, secretion of cytokines, increase in endocytosis, or enhancement in antigen processing and presentation. Interestingly, the group of Liljedahl (28) demonstrated recently that HLA-DO expression in B cells inhibits the presentation of such soluble fluid-phase endocytosed antigens. HLA-DO is almost exclusively expressed in B lymphocytes and interacts with HLA-DM to form a heterotetrameric structure (19). As a consequence of its association with HLA-DM and the colocalization in the MHC class II peptide loading compartment (MIIC), HLA-DO modulates the MHC class II/peptide repertoire (20, 21, 29). We reasoned that CD40 activation of B cells would probably lead to a decrease in the expression of this nonclassical class II molecule in our system. To investigate this potential mechanism in the observed enhancement in antigen presentation after B-cell activation, we analyzed the expression of molecules known to be critical for MHC class II antigen processing (15–17, 30). Analysis of HLA-DR, -DM, and -DO expression was performed by Western blotting on total cell extracts from control cells and B cells activated for various periods of time. Importantly, resting B cells highly expressed HLA-DO, which probably contributes to their inability to efficiently present soluble antigens to T cells. After activation with CD40L and IL-4, a continuous decrease in the expression of HLA-DO was observed as early as 2 days after activation, with only a faint band visible 6 days after activation (Fig. 3A). Additional analysis of the autoradiograms revealed that the band intensity ratios for HLA-DR:DO and HLA-DM:DO gradually increased after B-cell activation (Fig. 3B). The ratios for nonstimulated cells were adjusted to 1 to facilitate the analysis of the results. The HLA-DR/DO band intensity ratios increased to 6 (donor A) and 4 (donor B) after 2 days of activation, and additionally increased to 50 (day 13 for donor A) and 31 (day 16 for donor B) at later time points. The HLA-DM/DO band intensity ratios also increased to 9 (day 9 for donor A) and 10 (day 6 for donor B) but at later time points compared with the HLA-DR:DO band intensity ratios.

These data suggest that the enhanced recognition of pulsed-activated B cells by the specific CD4+ T cells could be a consequence of a change in the expression of molecules involved in antigen presentation by MHC class II, such as a decrease in HLA-DO expression.

Generation of Melanoma Antigen-specific T Cells Using CD40-activated B Cells Pulsed with a Melanoma Lysate. Having demonstrated that activated B cells can present exogenous antigens by MHC class II and considering that they were efficient T-cell stimulators (Fig. 1D), B cells were next evaluated for their capacity to promote the generation of tumor-specific T cells from resting peripheral lymphocytes. As a source of antigen, we used lysates from tumor cells, which are known to express a number of weakly immunogenic self-antigens (31), thereby rigorously testing the stimulatory capacity of the activated B cells.

Fig. 2. Presentation of gp100 by MHC class II by activated B cell. Purified B cells were concomitantly pulsed with recombinant gp100 protein and activated (A) for 18–24 h or were preactivated for 2 days (C2) and pulsed with protein for an additional 18–24 h as described in "Materials and Methods." B cells preactivated with CD40L and IL-4 for 5, 7, or 9 days were subsequently pulsed with protein for 18–24 h as indicated (d5, d7, and d9, respectively). C: HLA-DR7 or -DR7. CD40L and IL-4-activated B cells (left section) or tumor cells (right section) were treated with 100 μM chloroquine (CHL) or were left untreated (NT). Cells were then washed, and B cells were pulsed with gp100 as indicated for 20 h in the presence of 5 μM of chloroquine. Cells were washed once and fixed with 0.5% formaldehyde for 5 min. Cells were then washed extensively three times. For A–C, antigen processing was assessed by coculturing the pulsed B cells with a CD4+ T-cell clone specific to an MHC-DR7-restricted epitope of gp100 (A–C, left section) or a CD8+ T-cell clone specific for an HLA-A*0201 epitope of gp100 (B, right section) for an additional 18–24 h. Culture supernatants were collected and IFN-γ was measured using an ELISA assay (pg/ml). ESO-1, NY-ESO-1 recombinant protein; gp100, gp100 recombinant protein; DR7, HLA-DRB1*0701.
CD40-activated B cells were pulsed with a lysate prepared from a melanoma cell line and were used to stimulate autologous T cells from blood. In experiment 1, PBMCs were stimulated in a 96-well plate with autologous CD40-activated B cells pulsed with a lysate prepared from the melanoma cell line 1088mel. After three stimulations with 1088mel lysate-pulsed B cells, all of the 96-wells were individually tested against CD40-activated B cells pulsed with lysates prepared from 1088mel or from the irrelevant breast cancer line MDA231. Cells from 5 of the 96 wells were more reactive against the 1088mel lysate-pulsed B cells (Fig. 4A) compared with those pulsed with the MDA231 lysate. These results suggest that T cells reactive against antigens expressed in 1088mel were generated.

In a second experiment, CD4+ T cells purified from a different donor were stimulated three times with 1088mel lysate-pulsed autologous CD40-activated B cells 10 days apart in 1 well of a 48-well plate. The T-cell culture was cloned by limiting dilution and from 80 wells, 2 were consistently reactive against CD40-stimulated B cells pulsed with 1088mel lysate but not against those pulsed with a lysate prepared from MDA231 (Fig. 4B). The recognition was specific for melanoma because both clones failed to recognize a lysate prepared from EBV-immortalized B cells generated from patient 1088, excluding the possibility of recognition of allogeneic antigens.

Both T-cell clones were expanded using B cells pulsed with 1088mel lysate as stimulators. Clone 2 specifically expanded when stimulated with autologous B cells pulsed with 1088mel lysate (Fig. 5A), but failed to proliferate when stimulated with autologous B cells pulsed with a lysate from MDA231 or allogeneic B cells pulsed with either lysates from 1088mel or MDA231. Additional characterization of these two T-cell clones revealed that the antigen recognized was shared by at least two other melanoma cell lines (SK23mel and 553mel) when lysates were pulsed on B cells (Fig. 5B). Also, the recognition of CD40-activated B cells pulsed with 1088mel lysate was inhibited using a blocking pan-MHC class II antibody or an anti-HLA-DR antibody (data not shown), suggesting that the recognition was HLA-DR restricted. HLA-restriction was additionally characterized using CD40-activated B cells from HLA-DR-matched donors pulsed with 1088mel lysates. Clone 1 recognized only pulsed CD40-activated B cells prepared from HLA-DRB1*1501 donors (Fig. 5C).

In contrast to clone 1, clone 2 recognized only pulsed CD40-activated B cells prepared from HLA-DRB1*0401 donors, demonstrating that recognition by clones 1 and 2 was, respectively, HLA-DRB1*1501 and HLA-DRB1*0401 restricted.

To determine the antigen recognized, lysates were prepared from COS cells transfected with a panel of known melanoma tumor antigens (4, 22, 32). By using these pulsed B cells as targets for clones 1 and 2, tyrosinase was identified to be the antigen recognized by both clones (Fig. 5D). This observation was confirmed using B cells pulsed with lysates prepared from COS cells infected with a fowlpox virus vector expressing tyrosinase (data not shown).

In summary, T cells reactive against melanoma were generated using CD40-activated B-lymphocytes pulsed with lysates prepared from melanoma cells. These data suggest that CD40-activated B cells are efficient APCs in the presentation of exogenous antigens by MHC class II molecules and have the capacity to generate antigen-specific T cells.

Finally, clone 2 was used to examine whether B-cell stimulation...
was required for the presentation of the HLA-DR10401-restricted tyrosinase epitope. Purified B cells were left unstimulated or were activated by CD40L, IL-4, or a combination of these. B cells were then pulsed with lysates from 1088mel and used to stimulate clone 2. Clone 2 was reactive against B cells pulsed with antigen on day 0 and stimulated with CD40L alone (Fig. 6A; CD40L d0) or a combination of CD40L and IL-4 (CD40L/IL-4 d0), but failed to recognize B cells that were either nonstimulated (NS d0) or stimulated with IL-4 alone (IL-4 d0). The recognition of B cells cultured with CD40L and IL-4 for 5 days before antigen exposure was better compared with any other condition. These observations confirmed that short-term stimulation with CD40L is sufficient to increase the capacity of B cells to present exogenous antigens by MHC class II, as presented previously in Fig. 2A.

DISCUSSION

In the present work, we studied the consequences of CD40-activation on antigen presentation and T-cell stimulatory capacity of normal human antigen-unspecific B lymphocytes. Results presented show that CD40-activation of B cells increased their ability to present exogenous antigens by MHC class II. Indeed, B lymphocytes have critical functions of antigen uptake, processing, and presentation in the context of the humoral response, where antigens are taken up by the rare B cells expressing antigen-specific surface immunoglobulins, a situation that has been extensively studied. In the present paper, we show that after CD40-activation, nontransformed normal human B lymphocytes can uptake, process, and present exogenous antigens independent of the surface immunoglobulins. Interestingly, this experimental system allows the culture of primary B cells for up to 3 weeks, and a direct correlation was found between the length of activation and their ability to stimulate T cells. These CD40-activated B cells pulsed with a lysate prepared from melanoma cells had the capacity to generate T cells specific to melanoma. Our data suggest that indeed, when properly activated, B cells can present exogenous antigens by MHC class II molecules and stimulate antigen-specific T cells.

B-cell activation modulates the activity of other processes involved in antigen presentation such as increase in fluid-phase endocytosis, increase in the expression of cosstimulatory molecules (as shown in Fig. 1C demonstrating increased CD86 expression; Refs. 9, 33), and secretion of immunomodulatory cytokines (34). It is possible that activation of B cells by CD40L results in a number of changes leading to enhanced antigen presentation and T-cell stimulatory capacity. In this study, MHC class II antigen presentation by B cells correlated with the differential expression of classical and nonclassical MHC
CD40-ACTIVATED B CELLS PULSED WITH TUMOR LYSATES AS APC

**Fig. 6.** Presentation of tyrosinase by MHC class II by activated B cell. Purified B cells were concomitantly pulsed with a lysate prepared from 1088mel and activated (d0) for 18–24 h (A) or were preactivated for 2 days (d2) and pulsed with lystate for an additional 18–24 h (B) as described in “Materials and Methods.” B cells preactivated with CD40L and IL-4 for 5 or 12 days were subsequently pulsed with lysate for 18–24 h as indicated (d5 and d12, respectively). Antigen processing was then assessed by coculturing pulsed B cells with the tyrosinase-reactive T-cell clone 2 for an additional 18–24 h. Culture supernatants were collected, and IFN-γ was measured using an ELISA assay (pg/ml).

class II molecules after activation by CD40L. The expression of HLA-DO, a molecule described previously to alter the MHC class II/peptide repertoire and to inhibit the presentation of nonspecific exogenous soluble antigens, was decreased after B-cell activation with CD40L. By using B cells stimulated with an antibody specific to IgM compared with unstimulated cells, Roucard et al. (35) have reported a 50% decrease in the expression of HLA-DO, but no functional analysis of antigen processing was performed. In contrast to our work, no decrease in HLA-DO expression was observed when stimulating B cells with IL-4 and an antibody against CD40. Perhaps the decrease in HLA-DO that we observed after activation was because of B-cell stimulation with the more physiological soluble trimeric form of CD40L compared with the use of an antibody against the receptor. An increase in the HLA-DR:DO and HLA-DM:DO ratios should allow the loading of a broader repertoire of peptides, which bind with more stability to surface HLA molecules. Additionally, antigen-unspecific B cells that become activated by T cells may also increase their capacity to load antigens in less acidic compartments because of the diminished expression of HLA-DO. The down-regulation of HLA-DO might be one of several events leading to improved APC functions in addition to other mechanisms enumerated earlier. The exact mechanisms by which DO is down-regulated and potentially influences antigen presentation are currently under investigation.

The results presented suggest that, after CD40-activation, antigen-unspecific normal human B cells can efficiently present exogenous antigens through MHC class II. CD40L is mainly expressed on activated CD4+ T cells (36). In vivo, unspecific B lymphocytes could receive CD40 stimulation from activated CD4+ T cells during an immune response. We hypothesize that antigen-unspecific B cells receive stimulatory signals from activated CD4+ T cells, and as a consequence, B cells present exogenous antigens to T cells. The involvement of antigen-unspecific B cells in T-cell activation could enhance the magnitude of the T-cell response in an immune reaction. Although we used weakly immunogenic tumor cells as a source of antigen, B cells may also play a role in presenting more immunogenic foreign antigens after viral or bacterial infections. Additional work is needed to define the relevance of T-cell activation by antigen-unspecific B cells in vivo.

CD40-activated B cells can present exogenous antigens by MHC class II, and they are effective in the generation of antigen-specific T cells. For in vitro use, B cells have a number of advantages compared with other APCs, such as DCs, including a high proliferation rate, which allows the generation of a many activated B cells, even from cryopreserved PBMCs. This would allow the generation of an effective source of APCs when a small amount of blood is available (27). Antigen-specific T cells generated from lysate-pulsed activated B cells could be used to identify new tumor antigens from common cancers for vaccine development. In addition, tumor antigen-pulsed CD40-activated B cells could be used to generate specific T cells for adoptive immunotherapy, which has been demonstrated recently to result in high response rates in melanoma patients (37).

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CD40-stimulated B Lymphocytes Pulsed with Tumor Antigens Are Effective Antigen-presenting Cells That Can Generate Specific T Cells

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