Hyperexpression of CD40 Ligand by B and T Cells in Human Lupus and Its Role in Pathogenic Autoantibody Production

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Abstract

We investigated the role of the costimulatory molecules, CD40 and its ligand CD40L, in the pathogenesis of human SLE. In comparison to normal subjects or patients in remission, PBMC from active lupus patients had a 21-fold increase in the frequency of CD40L-expressing, CD4+ T cells. However, the expression of CD40L induced in either lupus or normal T cells by mitogenic stimulation could be down-regulated equally well by CD40 molecules on autologous B cells. Active lupus patients also had a 22-fold increase in percentage of CD8+ T cells expressing CD40L, consistent with their unusual helper activity in SLE. Surprisingly, patients with active lupus had a 20.5-fold increase in B cells that spontaneously expressed high levels of CD40L, as strongly as their T cells. Although lupus patients in remission had low levels of CD40L+ cells in the range of normal subjects, mitogen-induced upregulation of CD40L expression in their T and B cells was markedly greater than normal, suggesting an intrinsic defect. A mAb to CD40L blocked significantly the ability of lymphocytes from lupus patients with active and established disease to produce the pathogenic variety of antinuclear autoantibodies in vitro, bolstering the possibility of anti-CD40L immunotherapy for lupus. Future studies on the hyperexpression of CD40L could elucidate a regulatory defect in the pathogenic T and B cells of lupus. (J. Clin. Invest. 1996. 97:2063–2073.) Key words: system lupus erythematosus • anti-DNA autoantibodies • autoimmune disease • CD40 ligand expression • immunotherapy

Introduction

Production of the pathogenic variety of antinuclear autoantibodies in SLE (1–10), is driven by certain T helper (Th) cells that are prevalent in patients with active lupus nephritis but not in normal subjects or patients in remission (11–15). We were able to clone these Th cells in vitro by selecting for their special functional property (13, 14). Among 396 “autoactive” T cell clones derived from five patients with lupus nephritis, only 59 (~15%) had the ability to selectively augment the production of pathogenic anti-DNA autoantibodies when cultured with autologous B cells (14). Therefore, the pathogenic autoantibody-inducing Th cells of lupus are a special group of autoimmune T cells that are distinct from conventional autoreactive T cells. Analogous results were obtained in lupus mice (11, 12, 16–20) and similar pathogenic autoantibody-inducing Th clones could rapidly induce lupus nephritis when transferred in vivo into preautoimmune mice (16). Although the contribution of certain double negative (CD4+/CD8+) and CD4+ Th cells are important, CD4+ Th cells are the prime movers of the pathogenic autoimmune response in SLE (13, 14, 21). The initial findings showing that the antigen binding CDR3 loops of T cell receptors expressed by these special Th cells of lupus contain a recurrent motif of charged residues led to the identification of the autoantigens that activate them: charged DNA-binding proteins in nucleosomes, namely HMG and histones, presented by MHC class II molecules (15–18). The pathogenic autoantibody-producing B cells of lupus bind DNA complexed with these charged nuclear proteins (2, 9, 15, 16, 18). Most probably these B cells then process and present the nucleosomal proteins to the pathogenic T cells which in turn drive the B cells to make pathogenic autoantibodies (14–16, 18). This cognate Th and B cell interaction is not only essential for the production of pathogenic anti-DNA autoantibodies in SLE (12–14) but other types of autoantibodies as well (22–25). On the other hand, several studies (26, 27) have indicated that an intrinsic B cell hyperactivity that is manifest in lupus from early life (28, 29) is sufficient by itself to cause disease without the help of T cells.

Nevertheless, it is also widely believed that during cognate contact-dependent help for antibody production, the CD40 ligand (CD40L or gp39) is transiently expressed on activated CD4+ Th cells, and it binds to CD40 on the antigen-specific B cells transducing a second, costimulatory signal. This signal is essential for B cell growth and differentiation and for the generation of memory B cells by preventing apoptosis of germinal center B cells that have encountered antigen (30–42). Therefore, in this study we addressed the following questions: (a) Is the interaction between CD40 and its ligand, CD40L, important for the production of pathogenic autoantibodies in human lupus? (b) Is the expression of CD40L and its regulation defective in human lupus T cells? (c) Are human lupus B cells still dependent on CD40 transduced signals or are they autonomous? Surprisingly, we found that hyperexpression of CD40L occurs not only in the T cells but also in the B cells of lupus patients.

Methods

Patients and healthy donors. Eight patients with active lupus (seven females and one male; aged 18–50 yr), seven patients in long-term remission (all females; aged 21–51 yr) and six normal (healthy) subjects
(five females and one male; aged 22–40 yr) were studied. Disease activity by Systemic Lupus Activity Measure or SLAM (43), ranged between 7 and 20 for active patients, except for one patient who mainly had severe thrombocytopenia. None of the patients in remission had detectable proteinuria or serum anti-DNA autoantibodies at the time of testing and their SLAM ranged between 0 and 4. The patients in remission had never received any cytotoxic drugs, and they were not receiving any steroids at the time their blood samples were drawn for the assays. Steroids had been discontinued for several years in the remission patients, except for two patients who had received a short course of low dose steroids (Prednisone, 10 mg/d) 2 mo before the assays.

**Antibodies.** Anti-CD3 (OKT3) and anti-CD40 (G28-5) mAb secreting hybridomas were obtained from American Type Culture Collection (Rockville, MD). The hybridomas supernatants were concentrated by 47% saturated ammonium sulfate precipitation and dialysis before use. Anti-CD28 antibody (mAb 9.3) containing ascites was kindly provided by Kathy Cabrian (Bristol-Myers Squibb, Seattle, WA). Purified mAb to human CD40L (clone 24-31), anti-human CD40L FITC, as well as isotype matched control IgG1 antibody and control IgG1 FITC were obtained from Ancell (Bayport, MN). Phycoerythrin (PE)-conjugated mAbs to human CD4, CD8, CD20, and CD16, control PE- mAb, FITC-conjugated mAb to CD3 and CD69 were purchased from Becton Dickinson & Co. (San Jose, CA). For flow cytometry, all mAbs were used at optimal saturating concentrations as recommended by the manufacturers.

**Cell preparations and short-term T cell lines.** Peripheral blood mononuclear cells (PBMC) were obtained from lupus patients and healthy donor by centrifugation of heparinized blood over Ficoll-Hypaque (Pharmacia LKB Biotechnology Inc., Piscataway, NJ) gradient (13). CD4+ cells were selected from PBMCs using anti-CD4 mAb-coupled magnetic beads (Dynal Inc., Oslo, Norway) and rested overnight to detach from beads. CD4+ short-term T cell lines were made within 10–14 d after one round of stimulation of the CD4+ T cells with plate-bound anti-CD3 and anti-CD28 mAb and rIL-2 (20 U/ml).

**Stimulation of T and B cells in PBMCs ex vivo.** PBMCs (1.5–2.0 × 10^6/ml) were cultured for 18 h at 37°C with anti-CD3 mAb (1/256 dilution of SAS cut supernatant, unless specified otherwise) coated onto culture wells to stimulate T cells in vitro. Other aliquots of the diluted of SAS cut supernatant, unless specified otherwise) coated onto culture wells to stimulate T cells in vitro. Other aliquots of the PBMCs were cultured for 18 h at 37°C with Staphylococcus aureus Cowan I strain (SAC) at 1/30,000 (Pansorbin, Calbiochem Corp., La Jolla, CA) and 20 U/ml of rIL-2 to stimulate B cells in vitro (44). Cells were harvested and stained for flow cytometry.

**Flow cytometry analysis.** Unstimulated and stimulated cells were harvested and stained as described (13). Briefly, the cells were washed in FACS® buffer containing PBS/5% FCS/0.05% sodium azide and resuspended with 10 μg of human IgG (Sigma Chemical Co., St. Louis, MO) for 30 min at 4°C to block Fc receptors. Cells were washed to remove excess IgG and were double-stained with either FITC-conjugated mAb against human CD40L or FITC-control IgG mAb and PE-conjugated mAb against CD4 or CD8 or CD20 or CD16, or PE-control IgG mAb for 30 min at 4°C. Cells were washed twice, resuspended in FACS® buffer, fixed with 1% paraformaldehyde, and analyzed by flow cytometry using Becton Dickinson FACScan. PBMCs were gated according to their forward and side scatter characteristics. Different subsets of cells (CD4+, CD8+, CD20+) were gated further based on FL2 staining. The samples were analyzed using Cellquest software (Becton Dickinson).

**Regulation of CD40L expression in CD4+ short-term T cell lines.** CD4+ T cell lines were rested for 7–10 d in medium without rIL-2. The T cells (1 × 10^6/ml) were then stimulated in complete RPMI medium (13–15) with either anti-CD3 (1/125 dilution) coated on the culture wells or with 5 ng/ml PMA (Sigma Chemical Co.) and 0.5 μg/ml ionomycin (Sigma) for 6 h at 37°C. After the 6 h stimulation, in some cases, an equal number (1 × 10^6/ml) of autologous EBV-transformed B cell line (EBV-BCL) were added to the activated CD4+ T cells and incubated for an additional 2 h at 37°C according to described protocols (45–47). The EBV-BCL were pretreated with either 10 μg/ml of anti-CD40 or control mAb for 10 min before addition to the activated T cells. The cells were harvested after incubation and double stained to detect CD40L or CD40F expression on CD4+ T cells. Separate aliquots of T cells from the lines were incubated by themselves after PMA/ionomycin stimulation without coculturing with EBV-BCL and harvested at different time points to determine the percentage of CD4+ T cells that expressed CD40L by flow cytometry.

**Helper assays for IgG autoantibody production.** PBMCs (0.3 × 10^6 cells in 200 μl/microwell) were cultured in the presence of either anti-CD40L mAb (10 μg/ml) or isotype matched control IgG1 (10 μg/ml) in complete RPMI with 10% FCS for 13 d at 37°C (13, 14). Baseline production of autoantibodies secreted by lupus B cells was measured by culturing enriched B cells alone (0.1 × 10^6 cells/microwell). These B cells were obtained after depletion of CD4+ and CD8+ T cells from PBMCs, using anti-CD4 and anti-CD8 mAb coupled magnetic beads (Dynal). To exclude autoantibodies produced by B cells that were already preactivated in vivo, 120 μl of medium from each culture was replaced with fresh medium and corresponding mAb on day 5 without disturbing the cells at the bottom of the microwells (13). At the end of 13 d, production of IgG autoantibodies to single-stranded (ss) DNA, double-stranded (ds) DNA, and nucleosomes (histone/DNA complex), as well as total polyclonal IgG was quantitated in the culture supernatants.

**ELISA.** IgG class autoantibodies to ssDNA, dsDNA, and nucleosomes (histone/DNA complexes) as well as total polyclonal IgG were measured as described (13, 18, 48). Standard curves for anti-DNA ELISA were obtained by serial dilutions of one lupus patient’s serum that had high anti-DNA and antinucleosome autoantibody titer. All sera were heat inactivated at 56°C for 30 min before use. The absorbance value (at 405 nm) of 1/100 dilution of the reference patient’s serum was considered to be equivalent to 1 U/ml of anti-ssDNA, anti-dsDNA or antinucleosome autoantibodies, and the standard curve with the same reference serum was generated with each assay. Purified human IgG (Organon Teknika-Cappel, West Chester, PA) was used to obtain standard curves for total polyclonal IgG ELISA.

**B cell purification and semi quantitative reverse transcription (RT)-PCR for CD40L expression.** B cells were positively selected and isolated from PBMC using magnetic beads coated with mAb to CD19 (Dynal) according to the manufacturer. After detachment of the beads, cell purity was analyzed by flow cytometry by staining B cells with anti-CD20 mAb conjugated to PE. 99.7% of the purified cells were B cells with <0.2% being CD3+ cells.

Total RNA was extracted from enriched B cells or from T cells from the short-term lines (described above) using TRizol (Life Technologies Inc., Gaithersburg, MD) according to the manufacturer’s instructions. Reverse transcription was performed with 0.8 μg of RNA, 2.5 μM Random Hexamers, 1 mM dNTP, 3.5 μM MgCl2, 20 U RNase inhibitor, and 50 U murine leukemia virus reverse transcriptase in 20 μl PCR buffer. The mixture was incubated at room temperature for 30 min, followed by 42°C for 30 min, 99°C for 5 min, and 5°C for 5 min. All reagents used in the cDNA synthesis and PCR were from GeneAmp RNA PCR kit (Perkin Elmer, Branchburg, NJ). The protocol for amplifying CD40L message was as described previously (15, 17, 44) with slight modifications: after reverse transcription, 2.5 U Taq DNA polymerase, 3.75 μM MgCl2, and 150 nM of each primer in 80 μl × PCR buffer were added to the cDNA mixture. After denaturation at 94°C for 5 min, 45 cycles of amplification were performed at 95°C for 1 min, 54°C for 1.5 min, and 72°C for 2.5 min per cycle. The primers for CD40L message amplification were 5’-ACATAACAACCAACTTTCTCC-3’ (sense) and 5’-AGATGTTGGTACGCTG3’ (antisense). In some cases a second step PCR was performed to further amplify 1 μl of the first PCR product with 2.5 U Taq DNA polymerase, 150 nM of each primer, and 0.2 mM dNTP in 100 μl × PCR buffer with 3.75 μM MgCl2. The first PCR product DNA was denatured at 94°C for 5 min and amplified for 30 cycles, each at 95°C for 1 min, 54°C for 1 min, and 72°C for 2 min. As an internal control for each PCR, HLA-DQ α chain message, which is constitutively expressed by human B and T cells, was also...
amplified by RT-PCR under identical conditions except the primers were as follows: (GH26) 5'-GGTGGTACGATTTGAACAATTAGTCAG-3', and (GH27) 5'-CACGGATCCGGTAGCAGCGGTAGAGTTG-3' (49).

Results

Hyperexpression of CD40L by T and B cells of human lupus. PBMCs from patients with active lupus or patients in remission or normal subjects were analyzed by two-color staining and flow cytometry for CD40L expression (Table I and Figs. 1–3). In comparison to normal subjects, PBMC from patients with active lupus had on the average a 21-fold increase in the percentage of CD40L-expressing, CD4+ T cells (P < 0.0005, two-tailed t test). On further stimulation of the PBMC with anti-CD3, the percentage of CD40L-expressing CD4+ T cells increased fivefold in normal but only 1.4-fold in active lupus. However, the frequency of CD40L+/CD4+ T cells still remained markedly higher in active lupus (P < 0.001) than in normal PBMC after stimulation (Table I, Fig. 1). The anti-CD3 antibody was titrated and used at a submaximal stimulating dose (1:625 dilution) to bring out the differences in threshold of stimulation between lupus and normal T cells. In lupus patients in remission of their disease, the percentage of CD40L+ CD4+ T cells in freshly isolated PBMC was not significantly different from normal PBMC. However, on anti-CD3 stimulation the CD40L+ CD4+ T cells increased 13-fold in the inactive lupus patients and were significantly higher than in anti-CD3 stimulated PBMC of normal subjects (P < 0.01). Indeed, in contrast to the marked difference between unstimulated PBMC from active vs. remission patients (P < 0.001), the anti-CD3-stimulated PBMC of remission patients had a frequency of CD40L+ CD4+ T cells comparable to that from active lupus (P > 0.05).

Similarly, the frequency of CD8+ T cells that expressed CD40L in freshly isolated PBMC from active lupus patients (Fig. 2) was on the average ~22-fold higher than in normal subjects (P < 0.005) and 16-fold higher than in PBMC from patients in remission (P < 0.01). On anti-CD3 stimulation, the levels of CD40L+ CD8+ T cells in PBMC of active lupus remained markedly higher than in anti-CD3 stimulated normal PBMC (P < 0.001), or remission patient’s (P < 0.01) PBMC (Table I, Fig. 2). Although the unstimulated CD8+ T cells from normal subjects and patients in remission expressed comparable levels of CD40L (P > 0.1), on anti-CD3 stimulation the remission patient’s CD8+ T cells increased CD40L expression by 11-fold as compared to a 3.6-fold increase in normal PBMC (P < 0.05).

None of the patients in remission were receiving any cytotoxic drugs or steroids at the time of these assays. The percentage of CD4+ T cells that were CD69+ in the unstimulated PBMC from the remission patients was 0.52±0.22 (mean±SEM), and on anti-CD3 stimulation the percentage increased to 27.90±5.40. The corresponding values from the remission patients for CD8+ T cells that were CD69+ in the unstimulated PBMC were 1.97±0.43 (unstimulated) and 27.50±6.35 (anti-CD3 stimulated). These values were similar to those in normal subjects (see later in Table III, reference 46, and data not shown).

Remarkably, even without any stimulation in vitro, the B cells from patients with active lupus (Table I, Fig. 3) also had a marked increase in CD40L expression: 20.5-fold over that in normal subjects (P < 0.0005) and 12.3-fold over that in patients in remission (P < 0.0005). The levels of CD40L+ B cells in unstimulated PBMC were comparable in normal subjects and patients in remission (P > 0.1). Upon stimulation with SAC and IL-2, the frequency of CD40L+ B cells was still significantly higher in active lupus PBMC as compared to normal (P < 0.01), although the increase from unstimulated levels in the case of active lupus was 2.5-fold vs. a 7.6-fold increase in normal PBMC (Table I, Fig. 3). However, on stimulation the levels of CD40L+ B cells increased 17-fold in PBMC from patients in remission bringing them to levels comparable to active lupus PBMC (P > 0.1) but markedly higher than in stimulated normal PBMC (P < 0.01). The mean fluorescence intensities of CD40L expression by the CD4+ T cells and CD8+ T cells and the B cells of active lupus patients were comparable with mean±SEM values of 64.11±5.16, 72.50±8.76, and 67.39±4.73 respectively (Figs. 1–4). Acid wash of the B cells by incubating with PBS at pH 4.1 for 3 min, as described previously (44), did not decrease CD40L staining (data not shown).

Although the lupus patients had lymphocytopenia as compared to normal subjects, they still had significant increases in the absolute numbers of CD40L+ T and B cells (Table II), similar to the increases seen when the CD40L+ cells were expressed as percent values (Table I).

To establish further that CD40L expressed by B cells is not passively acquired from activated T cells, PCR amplifications of CD40L mRNA from highly purified B cells were done. B cells from a normal subject expressed CD40L only on stimula-

Table I. Hyperexpression of CD40L by Lupus T and B Cells

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Percentage of cells expressing CD40L among:*</th>
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<tbody>
<tr>
<td></td>
<td>CD4+ T cells</td>
<td>CD8+ T cells</td>
<td>B cells (CD20+)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>unstim. anti-CD3</td>
<td>unstim. anti-CD3</td>
<td>unstim. SAC + IL-2</td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>0.83±0.14 4.22±1.32</td>
<td>0.98±0.31 3.58±2.18</td>
<td>0.56±0.08 4.26±1.04</td>
<td></td>
</tr>
<tr>
<td>Active lupus</td>
<td>17.27±2.03 23.40±2.95</td>
<td>21.17±4.01 35.35±3.32</td>
<td>11.47±0.96 28.12±5.91</td>
<td></td>
</tr>
<tr>
<td>Remission</td>
<td>1.16±0.29 14.97±2.86</td>
<td>1.32±0.53 14.81±3.62</td>
<td>0.93±0.14 15.88±2.60</td>
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</table>

*Results obtained by flow cytometry of PBMC that were stained (two color) with mAb to CD4 and CD40L, or CD8 and CD40L, or CD20 and CD40L. The PBMC were stained either as unstimulated cells ex vivo (unstim.) or after stimulation with anti-CD3 mAb (anti-CD3) or with SAC + IL-2. Background values of staining by appropriate control mAbs were subtracted.
tion by SAC and IL-2 but not by anti-CD3 stimulation, indicating that CD40L message was endogenously upregulated in B cells and was not due to any contaminating T cells (Fig. 5). The presence of CD40L mRNA could be detected in normal B cells only after SAC + IL-2 stimulation, and only then after a second PCR amplification (30 additional cycles). In contrast, CD40L message was detectable in purified B cells from patients with active lupus, without any stimulation ex vivo, and after only one round of PCR, consistent with the hyperexpression seen by flow cytometry (Figs. 3–5, Tables I and II).

Two color staining of PBMC from lupus patients for natural killer (NK) cells with anti-CD16 and anti-CD40L mAb did not show any CD40L-expressing NK cells (data not shown).

Regulation of CD40L expression in Lupus T cells. On mitogenic activation of normal human T cells, CD40L expression peaks by 6–8 h and then goes down by 40 h (45–47, 50). Even

![Figure 1](http://www.jci.org)

**Figure 1.** Two color immunofluorescence (dot plots) of “gated” CD4⁺ T cells for CD40L expression. Examples from analysis of PBMC from normal subjects and lupus patients in remission or with active disease are shown. Based on background staining (not shown), the dot-plots were divided into four quadrants. Cells staining with both mAbs are in the right upper quadrant of each dot plot with percent positive cells indicated in parenthesis. Upper panels show results with unstimulated cells, and lower panels show that with anti-CD3-stimulated cells.

![Figure 2](http://www.jci.org)

**Figure 2.** Examples of two color immunofluorescence of CD8⁺ T cells expressing CD40L. The other features of this Figure are the same as in Fig. 1 legend.
without addition of any mitogen in vitro, the levels of CD40L+
T and B cells in the PBMC mixture of active lupus patients re-
mained high even after 40 h of culture (data not shown). This
result suggested either an intrinsic defect or perpetual rounds
of stimulation due to interactions between CD40L and CD40
molecules expressed by T and B, and B and B cells in the cul-
tured PBMC of lupus. To study the T cells in isolation, we de-
rived short-term (2-wk old) CD4 T cell lines from six patients
with active lupus and three normal subjects by just one round
of anti-CD3 and anti-CD28 stimulation of their CD4-enriched
T cells. CD40 expressed by B cells is known to rapidly down-
modulate CD40L, but not CD69 expression by activated T
cells from normal subjects (45–47). Therefore, we also derived
EBV-BCL that constitutively express CD40 (45–47) from
the same subjects from which the short-term T cell lines were de-
rived. After resting the T cell lines for 7–10 d, the unstimulated
T cells from active lupus patients tended to have higher levels
of CD40L-expressing CD4 T cells than those from the normal

Figure 3. Examples of two color immuno-
fluorescence of CD20 B cells
expressing CD40L. The other features
are the same as in Fig. 1 legend, ex-
cept for the lower panels where the
cells were stimulated with SAC and
IL-2.

Figure 4. Histograms showing the hy-
perexpression of CD40L (solid black)
by gated CD4 T cells and B cells
(CD20) of patients with active lupus
(active) as compared to those from
normal subjects. Percentage CD40L+ cells are indicated in parentheses.
These examples correspond to the dot
plots shown in Figs. 1 and 3, but the
histograms show the comparable in-
tensities of CD40L hyperexpression
in the T and B cells of lupus better.
Table II. Increase in Absolute Numbers of CD40L+ T and B Cells in Lupus

<table>
<thead>
<tr>
<th>Subjects</th>
<th>CD4+ T cells</th>
<th>CD8+ T cells</th>
<th>B cells (CD20+)</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>unstim.</td>
<td>anti-CD3</td>
<td>unstim.</td>
</tr>
<tr>
<td>Normal</td>
<td>6.04±0.80</td>
<td>28.05±7.40</td>
<td>4.98±1.25</td>
</tr>
<tr>
<td>Active lupus</td>
<td>29.19±7.53</td>
<td>43.39±12.11</td>
<td>52.45±9.53</td>
</tr>
<tr>
<td>Remission</td>
<td>3.93±1.20</td>
<td>50.57±10.36</td>
<td>1.63±0.74</td>
</tr>
</tbody>
</table>

*Results show the yield of CD40L+ cells in PBMC obtained from 1 mm3 of peripheral blood, either as unstimulated PBMC ex vivo (unstim.) or after stimulation overnight with anti-CD3 mAb or with SAC + IL-2. The number of CD40L+ cells in each subject was calculated by using the total PBMC count obtained immediately after purification from blood and the percent values of cells in each subset obtained by flow cytometry of the subject's PBMC (Table I and data not shown). In unstimulated PBMC, the total number (mean±SEM per mm3 of peripheral blood) of CD4+ T cells in normal subjects was 647.30±49.41, in patients with active lupus it was 197.76±65.48, and in remission patients it was 355.12±60.78. The corresponding values for CD8+ T cells were respectively, 418.33±76.38, 210.38±12.84, and 192.16±37.22; and those for B (CD20+ cells) were respectively, 218.25±7.17, 188.68±41.49, and 355.68±132.50. The percentage of CD40L+ T and B cells can be recalculated from the data in this Table and they are concordant with, but are not identical to, the values obtained directly from flow cytometry in Table I. The minor differences are probably due to loss of cells during processing of the PBMC for flow cytometry after the initial Ficoll-Hypaque purification and cell count.

T cell lines (Table III). Upon PMA + ionomycin stimulation for 6 h, both types of lines were maximally stimulated for expression of CD40L, and parallel staining for another T-cell activation marker with anti-CD69 supported this result. The up-regulation of CD40L in the CD4+ T cells in both types of T cell lines could be markedly downmodulated by incubation with autologous EBV-BCL for 2 h (Table III, Fig. 6). Preincubation of the EBV-BCL with anti-CD40 mAb could block the downmodulatory effect indicating that CD40 molecules on the EBV-BCL were responsible (Table III, Fig. 6). Similar results were obtained when anti-CD3 antibody was used at a submaximal dose to stimulate the T cell lines. CD40 molecules on autologous EBV-BCL downmodulated CD40L expression by the anti-CD3 activated T cells in both types of lines, even below their “resting” (unstimulated) levels. Interestingly, anti-CD3 stimulation induced a much higher level of CD40L-expressing T cells in the lupus T cell lines than the normal T cell lines, but their levels of CD69 expression were comparable (Table IIII). These results are similar to those obtained with freshly isolated PBMC (Tables I and II).

The kinetics of expression of CD40L by activated CD4+ T cells from lupus and normal subjects were also similar when T cells from respective lines were cultured by themselves after PMA + ionomycin stimulation, without the addition of any EBV-BCL (Fig. 7).

Anti-CD40L inhibits the production of the pathogenic variety of autoantibodies by lupus patient’s lymphocytes. Previous work showed that the production of pathogenic autoantibodies, such as the clonally restricted and predominantly cationic IgG autoantibodies to ssDNA, dsDNA, and nucleosomes by B cells in vitro, requires T cell help (13–15, 18) and is detectable only in PBMC from patients with active lupus (13). The results from six of the patients with active lupus B and T cells were cultured together as unfractionated PBMC vs. when the B cells were cultured alone. Moreover, the mAb to CD40L, in contrast to the control mAb, could block significantly the T cell augmented production of the IgG autoantibodies by PBMC of lupus patients with active and established disease (Table IV). In the six lupus patients tested, the anti-CD40L mAb blocked (mean±SEM of percent inhibition) IgG anti-ssDNA production by 418.33±76.38, IgG anti-dsNDA by 210.38±12.84, and IgG antinucleosome autoantibodies by 192.16±37.22. All of the patients began to receive steroids and/or cytoxan therapy for their active disease soon after the blood samples were obtained for these assays. The anti-CD40L-mediated inhibition of pathogenic autoantibody production, especially autoantibodies to dsDNA and nucleosomes, could still be detected in a patient who agreed to donate blood again for a repeat assay (EB and EB-2 in Table IV).

In the case of three of the patients (EB, SC, and VC), the input of CD40L message by B cells. Lanes 1 and 2 show PCR-amplified products from anti-CD3–stimulated normal T cells from a CD4+ T cell line; lanes 3 and 4 are from an active lupus patient’s B cells that were purified but unstimulated ex vivo; lanes 5 and 6 are from purified, normal B cells that were stimulated with anti-CD3; and lanes 7 and 8 are from purified, normal B cells that were stimulated with SAC and IL-2. The amplified CD40L band is 379 bp (lanes 2, 4, 6 and 8), and the constitutively expressed HLA-DQ band is 242 bp (lanes 1, 3, 5 and 7). Lanes 1, 2, 3, 4, 5 and 7 show the results after a second PCR amplification, designated by 2x (30 additional cycles). Unstimulated T cells from the T cell lines after resting for 10 d and unstimulated B cells from the normal subject did not express CD40L message that could be detected by PCR (not shown).
Table III. Regulation of CD40L Expression in CD4+ Short-term T Cell Lines

<table>
<thead>
<tr>
<th>Culture conditions*</th>
<th>Lupus T cell line</th>
<th>Normal T cell line</th>
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<tbody>
<tr>
<td></td>
<td>CD40L</td>
<td>CD69</td>
</tr>
<tr>
<td>Unstimulated</td>
<td>9.50</td>
<td>11.84</td>
</tr>
<tr>
<td>[P+I]</td>
<td>80.80</td>
<td>96.56</td>
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<tr>
<td>[P+I]→[EBV-BCL/c-mAb]</td>
<td>11.72</td>
<td>ND</td>
</tr>
<tr>
<td>[P+I]→[EBV-BCL/α-CD40]</td>
<td>78.10</td>
<td>ND</td>
</tr>
<tr>
<td>[α-CD3]</td>
<td>25.00</td>
<td>48.69</td>
</tr>
<tr>
<td>[α-CD3]→[EBV-BCL/c-mAb]</td>
<td>0.48</td>
<td>ND</td>
</tr>
<tr>
<td>[α-CD3]→[EBV-BCL/α-CD40]</td>
<td>17.10</td>
<td>ND</td>
</tr>
</tbody>
</table>

*Short-term T cell lines derived from CD4+ T cells of lupus patients or normal subjects were rested first (unstimulated) and then used for the different culture conditions, followed by two-color staining with either anti-CD4 and anti-CD40L or anti-CD4 and anti-CD69 mAbs. Representative examples from studies on short-term lines derived from six patients with active lupus and three normal subjects are shown. [P+I] means PMA + ionomycin stimulation. As indicated, in separate cultures [P+I] stimulation was followed by (+) coculture with either EBV-BCL preincubated with anti-CD40 (α-CD40) or EBV-BCL preincubated with isotype matched control (c-mAb). Anti-CD3 (α-CD3) mAb was used at 1:125 dilution. ND means not done, as it is known that CD69 expressed by activated T cells cannot be downmodulated by CD40 on EBV-BCL (references 45–47, and data on extreme right column).

Discussion

These findings on CD40–CD40L interaction in human SLE, some of which are quite unexpected, have implications for understanding the etiologic mechanisms of this disease as well as for developing specific immunotherapy. The increase in expression of CD40L by CD4+ T cells that occurs spontaneously in the patients with active lupus, resembles our results in murine lupus (48). In contrast to normal mice, lupus mice had much higher levels of CD40L-expressing T cells even at the preautoimmune age of 1 mo (48). Moreover, as shown here, the superinduction of CD40L on mitogenic stimulation of the lupus patient’s T cells, even during remission of the disease, indicates a defect in the regulation of CD40L expression. This interpretation is consistent with abnormalities in other signal transduction pathways that have been detected in the T cells of lupus (51, 52). The hyperexpression of CD40L by the lupus T and B cells persisted during the prolonged culture of the lupus PBMC in vitro (data not shown), most probably because of continued T-B interaction and rounds of stimulation in the culture mixture. However, in short-term CD4+ T cell lines, the superinduction of CD40L by mitogenic stimulation of the lupus T cells could be downmodulated by CD40 on autologous B cell lines, indicating that this negative feedback mechanism is still intact in lupus as in normal T cells (45–47).

The augmented expression of CD40L by CD8+ T cells in active lupus is another interesting finding, because it may explain the unusual helper activity of T cells with this phenotype in lupus (13, 21). Interestingly, CD8+ T helper cells were found subsequently in patients with AIHS that produce high levels of polyclonal Ig, including autoantibodies (53).

The most unexpected finding was the hyperexpression of CD40L by lupus B cells. It is widely accepted that CD40L expressed only by activated CD4+ Th cells has any functional consequence. Indeed in normal mice, CD40L expression by T

Production of IgG autoantibodies was preferentially blocked by anti-CD40L as compared to total polyclonal IgG which is produced at much higher levels than the autoantibodies in culture (Table IV). Whereas in the other patients, the production of total IgG was also markedly inhibited along with the autoantibodies. The anti-CD40L mAb (or the control mAb) was present in the PBMC cultures throughout the 13-d culture period (see Methods).

Figure 6. CD40 on B cells down-modulates CD40L expression by lupus T cells. Representative example of a histogram showing CD40L fluorescence (solid black) of gated CD4+ T cells. CD4+ T cells from short-term T cell line from a patient with active lupus were analyzed for CD40L expression under the following conditions: (A) at rest; (B) after stimulation for 8 h with PMA and ionomycin; (C) same as in B except autologous EBV-BCL B cells (pretreated with a control mAb) were added to the T cells in the last 2 h before staining; and (D) same as in C except for pretreatment of the EBV-BCL with anti-CD40 mAb before adding to the stimulated T cells.
cells only appear to be of any functional consequence in vivo (54). Recently however, while we were making these observations in lupus, Grammer et al. reported the expression of CD40L by normal human B cells (44). Our results with normal B cells confirm those observations. In addition, herein we found important differences in lupus B cells. Normal human B cells have to be coaxed by pharmacologic manipulations, such as a combination of calcium ionophore, phorbol ester, and IL-2 to express CD40L, and even then the level of expression is markedly less than activated T cells (44). In contrast, lupus patients’ B cells spontaneously express CD40L in levels as high as activated T cells. Furthermore, mitogen-stimulated up-regulation of CD40L is markedly greater in lupus T and B cells, even during remission of the disease, suggesting that a defect in SLE has lowered the threshold for CD40L expression, in addition to autoantigen-driven activation. None of the patients in remission were receiving any cytotoxic drugs or steroids, and the levels of CD69 (activated) T cells in their peripheral blood were in the range of normal subjects. Thus the lymphocytes of remission patients were not in a state of chronic activation by this criteria, and yet CD40L was hyperinducible in these patients’ T and B cells, suggesting an intrinsic abnormality.

CD40L is endogenously produced by the B cells; it does not appear to be passively acquired from the activated Th cells. The intensities of CD40L expression were similar in the B and T cells of lupus (Figs. 1–4). Stimulation with B cell mitogen (SAC and IL-2) caused a further increase in CD40L expression by B cells from patients with active lupus and markedly more so by B cells from remission patients (Figs. 3 and 4), and the CD40L staining could not be removed by acid-wash of the B cells (data not shown, but similar to reference 44). Furthermore, NK cells in the PBMC of lupus patients did not stain positively for CD40L (data not shown), again indicating that passively acquired CD40L shed from T cells is not a significant phenomenon. When CD40-expressing EBV-BCL B cells were cocultured with activated T cells expressing CD40L, the former cells did not pick up CD40L from the T cells; all the cells that stained for CD40L in such cocultures were CD4+ (Fig. 6 and dot plots not shown). Finally, PCR for CD40L mRNA showed that highly purified B cells from normal subjects express CD40L only upon stimulation with B cell mitogens but not a T cell mitogen (Fig. 5). This result again indicates that CD40L expression is endogenously upregulated by activated B cells and is not passively acquired from T cells. Further studies will determine whether the increase in CD40L mRNA in lupus B cells (Fig. 5) is due to increased transcription or due to increased stability of CD40L mRNA (55).

In contrast to other anti–human mAb, 5c8 and 4D9-9 (44), the mAb (clone 24-31) used here can detect CD40L on human B cells ex vivo without any acid treatment, suggesting that unlike the 5c8 epitope which may be masked by soluble CD40, the 24-31 (anti-CD40L) mAb detects another epitope that remains exposed. This feature may be therapeutically important in the future.

Thus in addition to being activated and costimulated by CD40L on Th cells, the lupus B cells could amplify and sustain the pathogenic autoimmune response further by expressing CD40L and thus interacting with CD40 on other autoimmune B cells in the vicinity as well as follicular dendritic cells in the germinal center (44, 56). Moreover, a bidirectional interaction between the CD40L+ B cells and the Th cells might occur in lupus, since CD40 itself might be expressed by activated T cells (57). Nucleosomal autoantigen presentation by the autoimmune B cells to the pathogenic Th cells (15, 18) could be markedly augmented by such T–B and B–B interactions, since signaling by CD40L upregulates the B7 family of molecules on B cells which would in turn costimulate the cognate Th cells via CD28/CTLA4 (32, 58, 59). However, the hyperexpression of CD40L by lupus B cells is not sufficient by itself to mediate an increase in pathogenic autoantibody production, at least in vitro (Table IV). Probably, the focused delivery of Th cytokines at a close range to the autoantibody producing B cells during their cognate contact-dependent interaction with the lupus Th cells is still required for pathogenic autoantibody production (Table IV). Nevertheless, the hyperexpression of CD40L by lupus B cells could increase B–B cell interaction via CD40 and thus prevent apoptosis or anergy of autoimmune B cells and make them more susceptible to pathogenic T cell help. The amazingly long-term benefit of a very brief anti-CD40L therapy in a mouse model of lupus (48) is consistent with the importance of CD40L–CD40 interactions in a variety of cells in the immune system that lead to pathogenic autoantibody production in SLE. However, the increased or aberrant CD40L expression is not global in SLE, NK cells did not express CD40L in patients with active lupus (data not shown).

The long-term beneficial effect of a brief anti-CD40L therapy in lupus mice (48) is relevant to our findings in human SLE. The effect of anti-CD40L in the mice was most probably mediated by checking the expansion the autoimmune memory B cells that are committed to produce pathogenic autoantibodies, since the brief therapy did not affect the Th cells in the long term (48). In the natural history of lupus, once the autoimmune memory B cells are generated they may further am-
plify the autoimmune response by efficiently presenting nucleosomal peptides (15, 18) and subsequently by “epitope spreading” present autoantibody idiotype-derived peptides to pathogenic Th cells (60). The brief anti-CD40L therapy probably also checked this amplification process. Moreover, the anti-CD40L therapy in lupus mice (48), although brief, could render autoimmune B cells tolerogenic by diminishing their costimulatory ability and antigen-presenting cell function (32, 54, 58, 59).

These studies also suggest that the pathogenic B cells of lupus can be downregulated and that they are not totally autonomous due to some intrinsic defect. The autoimmune B cells of most murine models of lupus (except MRL-lpr), as well as human SLE, may be as susceptible to apoptosis as normal antigen-driven B cells unless they are rescued by the CD40-mediated second signal (38, 40–42, 61). However, B cells capable of producing pathogenic anti-DNA autoantibodies are not deleted in the bone marrow; they are present in the periphery of normal subjects, and they can be expanded by various means to produce nephritogenic autoantibodies (5, 7, 62–66). Thus the factors determining the peripheral drive and cognate Th cell–induced pathogenic autoantibody production are relevant to understanding the basic mechanism of SLE.

Herein, the antibody to CD40L could block in vitro the production of IgG autoantibodies by cells from lupus patients with established and active disease. The anti-CD40L mAb was present in the culture for the entire period of 13 d; thus it also inhibited total polyclonal IgG production significantly in some of the patients. Th cells specific for other exogenous antigens, in addition to the autoimmune Th cells, were probably activated in those patients at the time blood sample was obtained and again during the prolonged culture period. Nevertheless, it is encouraging that significant blocking of pathogenic anti-DNA autoantibodies occurred in all patients tested even after the autoantibody-producing B cells and their Th cells had already been triggered in vivo in these patients with active and

### Table IV. Anti-CD40L mAb Inhibits the Production of Pathogenic Autoantibodies by Cells from Active Lupus Patients

<table>
<thead>
<tr>
<th>Lupus patient</th>
<th>Culture condition</th>
<th>( \alpha )-ssDNA</th>
<th>( \alpha )-dsDNA</th>
<th>( \alpha )-Nucleosome</th>
<th>Total IgG</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( U/ml )</td>
<td>( U/ml )</td>
<td>( U/ml )</td>
<td>( \mu g/ml )</td>
<td></td>
</tr>
<tr>
<td>EB</td>
<td>B cells alone</td>
<td>0.105</td>
<td>0.038</td>
<td>0</td>
<td>0.220</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.235</td>
<td>0.099</td>
<td>0.034</td>
<td>1.148</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.185</td>
<td>0.012</td>
<td>0</td>
<td>1.094</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>21.3</td>
<td>87.9</td>
<td>100</td>
<td>4.7</td>
</tr>
<tr>
<td>SC</td>
<td>B cells alone</td>
<td>0.070</td>
<td>0</td>
<td>0</td>
<td>1.729</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.100</td>
<td>0.074</td>
<td>0.022</td>
<td>2.315</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.046</td>
<td>0.030</td>
<td>0</td>
<td>2.152</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>54</td>
<td>59.5</td>
<td>100</td>
<td>7</td>
</tr>
<tr>
<td>RC</td>
<td>B cells alone</td>
<td>0.359</td>
<td>0.046</td>
<td>0.125</td>
<td>0.361</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.700</td>
<td>0.285</td>
<td>0.883</td>
<td>2.097</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.306</td>
<td>0.024</td>
<td>0.233</td>
<td>0.043</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>56.3</td>
<td>91.6</td>
<td>73.6</td>
<td>97.9</td>
</tr>
<tr>
<td>VC</td>
<td>B cells alone</td>
<td>0.025</td>
<td>0.015</td>
<td>0</td>
<td>1.050</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.353</td>
<td>0.112</td>
<td>0.098</td>
<td>1.440</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.147</td>
<td>0.039</td>
<td>0.028</td>
<td>1.186</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>58.4</td>
<td>65.2</td>
<td>71.4</td>
<td>17.6</td>
</tr>
<tr>
<td>IH</td>
<td>B cells alone</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.101</td>
<td>0.134</td>
<td>0.094</td>
<td>10.158</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.109</td>
<td>0.035</td>
<td>0.047</td>
<td>2.439</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>0</td>
<td>73.9</td>
<td>50</td>
<td>76</td>
</tr>
<tr>
<td>BM</td>
<td>B cells alone</td>
<td>0.032</td>
<td>0.020</td>
<td>0</td>
<td>0.375</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.284</td>
<td>0.559</td>
<td>0.465</td>
<td>1.793</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.143</td>
<td>0.397</td>
<td>0.299</td>
<td>1.176</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>49.6</td>
<td>29</td>
<td>35.7</td>
<td>34.4</td>
</tr>
<tr>
<td>EB-2</td>
<td>B cells alone</td>
<td>0.029</td>
<td>0</td>
<td>0.003</td>
<td>0.311</td>
</tr>
<tr>
<td></td>
<td>PBMC + c-IgG1</td>
<td>0.144</td>
<td>0.096</td>
<td>0.102</td>
<td>2.315</td>
</tr>
<tr>
<td></td>
<td>PBMC + ( \alpha )-CD40L</td>
<td>0.135</td>
<td>0.036</td>
<td>0.060</td>
<td>2.025</td>
</tr>
<tr>
<td></td>
<td>Percent inhibition</td>
<td>6.3</td>
<td>62.5</td>
<td>41.2</td>
<td>12.5</td>
</tr>
</tbody>
</table>

\(^3 \times 10^7\) PBMC (contains both T and B cells) or 1 \times 10^5 B cells alone were cultured per microwell. c-IgG1 means control IgG1 that is isotype-matched to the anti-CD40L mAb (\( \alpha \)-CD40L). Values of percent inhibition of IgG antibody production by PBMC with \( \alpha \)-CD40L are shown. \( ^* \)IgG autoantibodies are expressed in units per milliliter based on reference serum standard. \( \alpha \)- means anti-. ND, not done. Total IgG is expressed in micrograms per milliliter as it is produced in much higher amounts than the autoantibodies in the culture.
established disease. This result suggests that the pathogenic Th and B cells of lupus undergo repeated cycles of activation and rest or different waves of activated cells participated in the production of pathogenic autoantibodies during the 13-d culture period. Interestingly, a brief anti-CD40L therapy given at early stages of lupus in mice preferentially blocks the pathogenic autoimmune response in vivo, but it does not cause generalized immunosuppression (48). This beneficial result could be due to the fact that CD40L is transiently expressed on activated T cells, and in the lupus mice, most of the T cells that are activated are the Th cells that drive B cells to make pathogenic autoantibodies. Indeed, anti-CD40L therapy was beneficial even at the time early autoimmune abnormalities were manifest in the lupus mice. This suggests that anti-CD40L therapy might be beneficial in newly diagnosed lupus patients or in patients in the early stages of disease flare-up. Interestingly, blocking B7–CTLA4/CD28 interaction by CTLA4Ig also has a beneficial effect on murine lupus (67), probably through its indirect effect on the CD40L–CD40 signaling pathway (32, 68).

These studies in human and murine SLE should stimulate the development of anti-CD40L therapy in human lupus (69). We are even more interested in understanding why CD40L expression is deregulated in the B and T cells of lupus? It appears to be an intrinsic defect and not merely or wholly due to autoimmune activation. Remarkably, the defect in the regulation to be an intrinsic defect and not merely or wholly due to autoimmune activation is deregulated in the B and T cells of lupus mice. This suggests that the pathogenic anti-DNA autoantibody-inducing T helper cell lines from patients with active lupus nephritis: Isolation of CD4+CD8+ T helper cell lines that express the 9b-T cell receptor. Proc. Natl. Acad. Sci. USA. 87:7020–7024.


CD40L Hyperexpression in Human Lupus

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