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Abatacept Limits Breach of Self-Tolerance in a Murine Model of Arthritis via Effects on the Generation of T Follicular Helper Cells

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Abatacept modulates CD28-mediated T cell costimulation and is efficacious in the treatment of rheumatoid arthritis (RA). Its mechanism of action has not been fully elucidated but will likely reveal critical pathologic pathways in RA. We show that abatacept substantially modulated Ag-specific T and B cell responses in vivo. Ag-specific T cell proliferation was reduced, and the acquisition of an activated phenotype, characterized by upregulation of CD69, OX40, ICOS, and programmed death-1 and downregulation of CD62L, was suppressed. Furthermore, abatacept suppressed the production of inflammatory cytokines, such as IFN-γ and IL-17. These effects were associated with a failure of Ag-specific T cells to acquire the CXCR5+ICOS+ T follicular helper cell phenotype. This, in turn, led to a failure of these cells to enter B cell follicles, resulting in reduced specific Ab responses, despite normal B cell clonal expansion. To test the pathologic significance of this, we used a novel model of RA associated with breach of self-tolerance to self-Ag and demonstrated that abatacept prevented the emergence of self-reactivity. Thus, CD28-dependent signaling is required for optimal T follicular helper cell maturation and expansion, and its inhibition prevents loss of self-tolerance in a model of articular pathology. Thus, we provide a novel mode of action for abatacept with profound implications for its potential usefulness in early inflammatory arthropathies associated with autoantibody expression. The Journal of Immunology, 2010, 185: 000–000.
Ag-specific lymphocyte behavior in vivo. We used transgenic (Tg) T cell and B cell co-transfer models to dissect the effects of abatacept on specific T cell responses and T-dependent B cell responses using trackable lymphocyte populations. Abatacept profoundly suppressed Ag-specific T cell proliferation and phenotypic activation. Furthermore, the production of inflammatory mediators, such as IFN-γ, IL-17, and MIP-1α, was reduced. Crucially, these effects were associated with a failure of lymph node (LN) Ag-specific T cells to acquire a CXCR5+ICOS+ T follicular helper (TFH) cell phenotype. This led to a failure of TFH cell migration to B cell follicles, resulting in reduced specific Ab responses, despite normal B cell clonal expansion. The clinical relevance of these data was clearly demonstrated in a novel model of RA in which abatacept prevented breach of self-tolerance, reducing anti-self Ab responses.

Materials and Methods

Mice

IgHb BALB/c (H-2d, IgM+) mice were bred in-house (Central Research Facilities, University of Glasgow). Mice homozygous for the chicken OVA peptide (323–339)-TcR-specific DO11.10 TCR transgenes (detected using the clonotypic mAb KJ1.26) on the BALB/c background were used as donors, as were mice heterozygous for the anti- hen egg lysozyme (HEL) IgM+ and IgD+ transgenes on the BALB/c background (M4). BALB/c or IgHb mice 6–12 wk old were used as recipients. All animals were specified pathogen-free and were maintained under standard animal house conditions in accordance with local and home office regulations.

Reagents

Abatacept was provided by Bristol-Myers Squibb. Chi L6 is a chimeric fusion protein consisting of the V region of murine L6 Ag, combined with a human IgG1 C region; it was used as a control fusion protein (control Ig) in these studies. Abatacept and control Ig were injected i.p. at a dose of 10 mg/kg.

Preparation of cell suspensions for adoptive transfer

Peripheral LNs (pLNs) and spleens from MD4 BALB/c and DO11.10 BALB/c mice were pooled and prepared as described previously (20). Cells were rendered with 8.5 μM CFSE in HBSS for 10 min at 37°C, as described previously (20), before cell suspensions containing 1 X 10^6 CFSE-labeled Tg T cells and 1 x 10^6 unlabeled Tg T or B cells were injected i.v. into nonirradiated, age-matched IgHb BALB/c recipients, as previously described (20). Alternately, 5 x 10^5 cells were labeled with 5 μM Cell Tracker Red (CMTPX) in CO2-independent media (Invitrogen, Paisley, Scotland, U.K.) for 40 min at 37°C for multiphoton imaging.

Adoptive transfer of OVA-specific Th1 cells and arthritis model

Arthritis was induced in BALB/c recipients by i.v. transfer of 2 x 10^5 in vitro, Th1-polarized, KJ1.26+ T cells, followed 1 d later by restimulation in vivo with 100 μg OVA in CFA in the scruff of the neck; 10 d later, 100 μg heat-aggregated OVA (HAO) alone was given by footprint injection. Mice were monitored for development of arthritis, as described previously (21).

Immunofluorescence analysis of tissue sections and confocal microscopy

Ag-specific T cell localization in the LN was determined by immunofluorescence as described previously (20), using B220-PE to stain B cell areas, anti-CD4 Alexa Fluor 450 for T cell areas, and biotinylated-KJ1.26 to detect OVA-specific cells and was visualized using streptavidin-Alexa Fluor 647 (Molecular Probes, Eugene, OR). All photographs were taken at x20 or x40 magnification.

Flow cytometry

Aliquots containing 1 x 10^5 cells were incubated in 200 μl FACS buffer (PBS/2% FCS) for 15 min at 4°C with purified anti-mouse CD16/CD32 (BD Biosciences, San Jose, CA), washed, and incubated with the appropriate fluorochrome-conjugated or biotinylated primary Abs for 30 min at 4°C. The cells were washed, resuspended in FACS buffer, and, where appropriate, biotinylated Abs were detected by fluorochrome-conjugated streptavidin for an additional 15 min. Cells were then analyzed on a FACS-Canto Flow Cytometer (BD Biosciences) using FlowJo software.

In vitro Ag-specific cytokine production

Cytokine levels were determined in cultures of total draining LN cells stimulated with 5 μM OVA (Sigma-Aldrich, St. Louis, MO) for 48 h. Cells were cultured at 2.5 X 10^6 cells/200 μl in round-bottom 96-well plates. Supernatants were assayed for levels of fibroblast growth factor; GM-CSF; IFN-γ; IL-1α/β; 2, 4, -4, 5, 6, -10, -12 (p40/g70), -13, and -17; CXCL10; CXCL1; MCP-1; CXCL9; MIP-1; TNF; and vascular endothelial growth factor by Luminex (Invitrogen) and analyzed using a Luminex XMAP system, according to the manufacturer’s instructions.

In vitro restimulation assay

PLN cells were cultured with medium or 1 mg/ml OVA. Proliferation was analyzed at 96 h by flow-cytometric staining for EdU (5-ethyl-2-deoxyuridine) incorporation (Invitrogen).

In vivo challenge

One day after adoptive transfer of naïve T cells, recipient BALB/c mice were challenged in the footpad with 100 μg OVA in CFA or CFA only as an adjuvant control (or PBS, because no difference in KJ1.26 expansion between PBS and CFA challenge was seen). Chicken OVA (fraction V) was obtained from Sigma-Aldrich, and HEL was obtained from Biozyme (Gwent, U.K.). Conjugated OVA-HEL was prepared as previously described (20). Animals were injected s.c. with 150 μg chicken OVA-HEL or 100 μg OVA, both in CFA (Sigma-Aldrich).

Quantification of follicular migration of Ag-specific T cells

The proportion of OVA-specific (KJ1.26+) cells was quantified using Volocity software. The localization of Tg T cells and B cell follicles was plotted. Using these tissue maps, the number of Tg T cells in defined gates was calculated for three gates for the total section and three B cell follicle gates per section. Data are plotted as the mean proportion of transgenic T cells (KJ1.26+) within the follicle (B220+), calculated by dividing the number of Tg T cells in the B220+ follicle by the total number of KJ1.26+ cells within the total field of view. Data shown are the mean of triplicate readings from three mice per group.

Ab ELISA

Anti-OVA, anti-HEL, and anti-collagen Abs were detected as previously described (20, 21).

Multiphoton microscopy

To image cellular behavior in LN s, excised LN s were transferred into CO2-independent media at room temperature. The LN was bound with veterinary glue (Vetbond, 3M, St. Paul, MN) onto a plastic coverslip that was then adhered with glue to the bottom of the imaging chamber continuously supplied with warmed (36.5°C) and gassed (95% O2 and 5% CO2. RPMI 1640 before and throughout the period of microscopy and was imaged, as previously described (22).

Statistical analysis

Results are shown as mean ± 1 SD, unless stated otherwise, and groups were compared using the Student two-tailed unpaired t test. Two-way ANOVAs were used to compare Ab levels.

Results

Abatacept modulated phenotypic and functional aspects of Ag-specific T cell activation

Abatacept inhibits T cell proliferation in vitro in human MLRs and in vivo in alloresponses (12, 18). We first assessed whether this could be reproduced in our Ag-specific model system in vivo. We measured division of CFSE-labeled, DO11.10 OVA-specific T cells transferred into naive BALB/c recipients that were then immunized s.c. with 100 μg CD44+KJ1.26+ cells that had undergone fewer than four divisions (Fig. 1B, 1C).

Next, we assessed the ability of abatacept to modulate the activation of OVA-specific T cells in vivo. As before, naive BALB/c

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mice received $1 \times 10^6$ DO11.10 T cells i.v. and were immunized with OVA/CFA in the footpad 24 h later. Three days later, the draining popliteal LN was harvested, and OVA-specific T cells were assessed for the expression of activation markers and markers associated with tolerance and/or regulatory T cells. Abatacept significantly reduced Ag-induced downregulation of CD62L on OVA-specific T cells (Fig. 2A), and the upregulation of OX40 by Ag-specific T cells (Fig. 2B). Strikingly, abatacept prevented the Ag-induced expression of ICOS and programmed death-1 (PD-1) (Fig. 2C, 2D), which are coexpressed by CXCR5+ TFH cells (23). The expression of CD28 and CD25 3 d postimmunization was not altered by treatment with abatacept (Fig. 2E, data not shown). Finally, CD69 expression by Tg T cells 1 d postimmunization was also significantly reduced by treatment with abatacept (data not shown).

The ability of abatacept to alter Ag-specific T cell functionality was assessed following transfer of DO11.10 T cells and OVA/CFA immunization. On day 3 postimmunization, total dLN cells were cultured in vitro with or without OVA peptide (323–339) for 48 h, and the supernatants were analyzed by Luminex. Production of inflammatory mediators was only detectable when OVA peptide was present in vitro and then only from cultures derived from animals receiving OVA/CFA in vivo (Fig. 2F–I). Abatacept significantly decreased synthesis of the inflammatory mediators IFN-γ, IL-17, MIP-1α, and IL-13 compared with control IgG-treated animals.

Abatacept failed to alter T cell velocity or meandering in dLNs

We and other investigators demonstrated previously that formation of dendritic cell (DC)–T cell clusters of smaller sizes and interactions of shorter duration fail to drive fully productive T cell responses (22, 24). Together with the postulated mode of action of abatacept, this prompted us to use multiphoton microscopy to examine whether abatacept affected the migration of specific T cells in intact LNs ex vivo. CMTPX-labeled DO11.10 T cells were adoptively transferred into BALB/c mice that were subsequently immunized s.c. with OVA/CFA and treated with control IgG or abatacept. To directly compare T cell behavior in immunized mice following treatment with control IgG or abatacept, the volume and location of each cell were measured in 15 planes per three-dimensional stack collected at 1.95-μm intervals between each plane, progressing deeper into the LN from an initial depth of 100 μm below the surface. Twenty-four hours after immunization, Ag-specific T cells had similar velocities and meandering indices (a measurement of directionality of cell movement) in the presence of abatacept compared with control IgG (Fig. 3A, 3B, Supplemental Videos 1, 2). Previous studies by ourselves and other investigators showed that DCs migrate to the LN within 24 h of immunization; this is when the interaction with DCs that determines the outcome for the T cell is taking place (22). To determine whether abatacept affected T cell interactions with APCs, we analyzed the proportion of T cells in clusters, the number of cells in clusters, and the number of cells per cluster, as previously described (22). Although a trend for a reduced proportion of T cells in clusters and the number of cells in clusters was apparent in the presence of abatacept, it was not significant (Fig. 3C, 3D). In addition, there was no difference in the number of T cells per cluster after abatacept treatment (Fig. 3E).

Abatacept suppressed acquisition of TFH cell phenotype and follicular migration of Ag-specific T cells

Although abatacept did not alter T cell velocity within the dLNs, the reduced expression of markers associated with a TFH cell phenotype prompted us to investigate whether abatacept affected the localization of Ag-specific T cells within LNs in situ. BALB/c mice were adoptively transferred with DO11.10 T cells and immunized with OVA/CFA, as before. Three days later, the dLN was assessed for...
the localization of Ag-specific T cells, as shown previously (25). Following immunization, Ag-specific T cells increased in number, and a large proportion of these could be seen entering the B cell area (Fig. 4A). In contrast, few Ag-specific T cells were observed in the presence of abatacept, and strikingly, the majority of these were restricted to the paracortical T cell zone. The total number of Ag-

FIGURE 2. Abatacept reduces phenotypic and functional aspects of Ag-specific T cell activation. LNs from mice that had received 1 × 10^6 DO11.10 T cells before immunization with OVA/CFA in the footpad and treated with PBS, control IgG, or abatacept i.p. immediately after immunization were harvested 3 d later. Unimmunized animals received PBS. Tg T cells (CD4*KJ1.26*) were assessed for the expression of CD62L (A), OX40 (B), ICOS (C), PD-1 (D), and CD28 (E). Some LN cells were also cultured with or without OVA peptide (323–339; 1 μg/ml) for 48 h in vitro, and supernatants were analyzed by Luminex. The mean ± SD production of IFN-γ (F), IL-17 (G), IL-13 (H), and MIP-1α (I) is shown. Data are representative of at least two independent experiments. ∗p < 0.05, ∗∗p < 0.01, ∗∗∗p < 0.001; Student two-tailed unpaired t test. ns, not significant.
Abatacept reduced Ab production independently of B cell clonal expansion

We demonstrated previously in infectious diseases, such as malaria, that reduced migration of T cells into B cell follicles results in suppressed B cell responses to heterologous Ags (27). Moreover, CTLA-4Ig was reported to suppress Ab production following immunization with sheep erythrocytes and keyhole limpet hemocyanin (19). However, the mechanism by which CTLA-4Ig exerts these effects, and its potential effects on other aspects of B cell responses, remains unclear. Therefore, we assessed the impact of the impaired ability of Tg T cells to migrate into B cell follicles in the presence of abatacept on Ag-specific B cell responses. We approached this using a Tg system that allows measurement of systemic anti-HEL B cell responses, which are dependent upon help from Tg T cells (25). We cotransferred OVA-specific Tg T cells and HEL-specific Tg B cells into naive IgHb recipients and then immunized them with an OVA-HEL conjugate prepared in CFA. Following immunization, anti-HEL IgM° serum titers were significantly reduced by a single treatment with abatacept at the time of immunization (Fig. 5A). The endogenous anti-HEL response was negligible by day 5 postimmunization, because serum anti-HEL IgM° was undetectable (data not shown). However, the endogenous B cell response to OVA was detectable by day 5 postimmunization; although anti-OVA IgG2a responses were undetectable, increased levels of anti-OVA IgG1 responses were observed in immunized animals (Fig. 5B, data not shown). Similarly, the endogenous anti-OVA IgG1 response was significantly reduced after abatacept treatment (Fig. 5B). Interestingly, although abatacept reduced the number of CD4°KJ1.26° Tg T cells in the dLN as expected, it did not significantly alter the clonal expansion of IgMa° B cells 5 d after immunization (Fig. 5C, 5D). Because the effect of abatacept on Ab production seemed to be independent of clonal expansion, we next assessed whether the formation of germinal centers (GCs) was affected. Somewhat surprisingly, the percentage of Ag-specific B cells that expressed the GC B cell markers GL7 and Fas was not significantly reduced by abatacept, as assessed by flow cytometry (Fig. 5E). Together, these results suggest that abatacept modulates the function of B cells rather than reduces B cell numbers.

Abatacept prevented breach of B cell self-tolerance in arthritis

In light of the effects of abatacept on T cells, TFH cells, and B cells, as well as its therapeutic application, we next assessed its impact on these parameters in a novel model of RA. In this model, breach of self-tolerance is characterized by the spontaneous development of an autoimmune response against articular Ags, such as type II collagen (CH) (21). Following challenge with HAO, abatacept treatment reduced footpad swelling and clinical scores compared with control IgG (Supplemental Fig. 1). Popliteal LNs were...
harvested, and T cell recall responses to OVA were assessed. In vitro recall anti-OVA responses by CD4+ T cells were greater in mice that had been challenged with HAO compared with PBS, indicating increased expansion and/or recruitment following secondary Ag exposure (Fig. 6A). Animals treated with abatacept manifested significantly reduced T cell proliferative responses to OVA (Fig. 6A). In addition, the proportion of total CD4+ T cells expressing CXCR5 was significantly reduced by treatment with abatacept in vivo (Fig. 6B).

We measured autoreactive Ab levels to assess the impact of abatacept on breach of B cell self-tolerance. Following HAO challenge, anti-CII IgG responses increased compared with PBS challenge, indicating increased expansion and/or recruitment following secondary Ag exposure (Fig. 6C). Animals treated with abatacept manifested significantly reduced T cell proliferative responses to OVA (Fig. 6A). In addition, the proportion of total CD4+ T cells expressing CXCR5 was significantly reduced by treatment with abatacept in vivo (Fig. 6B).

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**Discussion**

Abatacept has proven efficacious in the treatment of RA, but its mode of action in vivo remains to be fully elucidated. Indeed, its effects on specific T cell responses and the development of specific T-dependent B cell responses in vivo are unclear, as are its effects on memory cell responses and breach of tolerance to self, joint-associated Ags. In this article, we provide evidence that, in addition to inhibiting T cell activation and proliferation, abatacept reduces the migration of T cells into B cell follicles. Importantly, this results in reduced Ab responses, independently of B cell clonal expansion. This failure of T cell help for B cells and the subsequent Ab response are recapitulated in a novel model of RA in which abatacept prevents breach of self-tolerance, as well as reduces anti-self Ab responses and associated pathology.

Abatacept was demonstrated to inhibit T cell proliferation in in vitro human MLRs and in alloresponses in vivo (12, 18). In this study, we examined its effects on Ag-specific T cell responses in vivo; abatacept dramatically reduced proliferation after challenge with Ag in adjuvant. Furthermore, abatacept reduced the downregulation of CD62L expression, a characteristic of activated T cells that is needed for cells to exit the LN to migrate to tissue sites of inflammation (28). The modulation of CD62L expression suggests abatacept could inhibit the migration of T cells into inflamed tissues. As expected (29), Ag-specific T cells increased their expression of OX40, which is downstream of CD28, after challenge with Ag in adjuvant. However, treatment with abatacept suppressed the expression of OX40, which could culminate in reduced survival of specific T cells as a result of the reduced expression of the antiapoptotic factors Bcl-2 and Bcl-xL (30). Interestingly, OX40 also controls the expression of survivin, which maintains T cell division over time (31). These findings are consistent with a previous report demonstrating reduced CD28-
dependent OX40 expression in CTLA-4Ig Tg mice, which correlated with reduced expression of CXCR5 on CD4+ T cells (32). The most striking effect of abatacept on activation marker expression was on the upregulation of ICOS, which was completely abrogated, consistent with previous reports on T cell populations in systemic lupus erythematosus (33). ICOS plays a critical role in T cell proliferation, the development of TFH cells, the formation of GC, and the generation of T-dependent Ab responses (34–36). Abatacept also significantly reduced the upregulation of PD-1, a negative regulator of T cell activation (37), by activated Ag-specific T cells. Interestingly, ICOS and PD-1 are also coexpressed by CXCR5+ TFH cells (23, 38, 39). Abatacept had no effect on the upregulation of CD28 by activated T cells. We also demonstrated that abatacept reduced the Ag-driven production of the inflammatory mediators IFN-γ, IL-17, IL-13, and MIP-1α ex vivo. These cytokines are known to play important roles in several inflammatory diseases, including RA (40–42). Although CTLA-4Ig was demonstrated to reduce serum IL-17 during collagen-induced arthritis (43), this is the first study to show a direct inhibition of Ag-specific T cell IL-17 production by abatacept. Interestingly, IL-13 was shown to be produced by TFH cells in human tonsils (44), and this was significantly reduced by treatment with abatacept.

We and other investigators demonstrated previously that formation of DC–T cell clusters of smaller sizes and interactions of shorter duration fail to drive fully productive T cell responses (22, 24). This, together with the proposed mode of action of abatacept, prompted us to examine the migration of specific T cells in intact LNs ex vivo using multiphoton microscopy. In this study, abatacept did not significantly alter Ag-specific T cell velocity or meandering following immunization with Ag in adjuvant. However, when we analyzed T cell clustering, there was a trend toward a reduced proportion of cells in clusters, as well as in the number of cells per cluster, but this was not significant. However, as we showed previously (45), further studies using fluorescently labeled DCs and T cells will allow more detailed analyses of the effects of abatacept on cellular interactions.

However, when we examined the localization of Ag-specific T cells in situ, we observed that in abatacept-treated animals, the majority of cells were restricted to the paracortical T cell zone even following immunization. The total number of Ag-specific T cells entering B cell follicles was clearly reduced after abatacept treatment, in part because of its dramatic effects on proliferation; however, the proportion of T cells in follicles was also reduced. Interestingly, this lack of TFH cells could also be related to the inhibition of proliferation, because far fewer cells had undergone four or more divisions, which is a characteristic of this more-differentiated subset (46). The lack of follicular migration was also associated with suppression of the acquisition of a CXCR5+ ICOS+ TFH phenotype among the Tg population. Failed migration likely reflects the inability of Ag-specific T cells to migrate to CXCL13 via CXCR5.

We demonstrated previously in infectious diseases, such as malaria, that reduced migration of T cells into B cell follicles results in suppressed B cell responses to heterologous Ags (27). Therefore, we assessed the impact of the impaired ability of Tg T cells to migrate into B cell follicles in the presence of abatacept on Ag-specific B cell responses. CTLA-4Ig was reported to suppress Ab production following immunization with sheep erythrocytes and keyhole limpet hemocyanin (19). Although abatacept suppressed the proliferation of Tg T cells in these experiments, we also demonstrated that it reduced the systemic Ab response without affecting the clonal expansion of Tg B cells. Together, these results suggest that abatacept acts to modulate the function of B cells rather than reduce B cell numbers. It remains possible that in the T and B cell cotransfer experiments, T cell migration into follicles was increased as a result of the effects of chemotactic factors, such as CXCL13, which can be expressed by activated...
B cells (47). It is also interesting to speculate that because B cells express CD80/CD86, abatacept could prevent phenotypic maturation of B cells, which may relate to the reduced levels of Ab production observed. We then investigated whether this failure of T cell help for B cells and the subsequent Ab response were recapitulated in a novel OVA-driven model of RA. We previously described the suggested mode of action of this model in detail (21, 48–50). Briefly, we think that by transferring a large number of activated Th1 cells of an irrelevant specificity and inducing a local inflammatory stimulus in the joint, the normal regulatory mechanisms are overwhelmed and breach of self-tolerance occurs. We suggested previously that the inflammation initiated by OVA-specific T cells causes normally quiescent, tolerogenic DCs carrying self-Ag (collagen) from the joint to the local LN to present this in an immunogenic manner to self-reactive T cells, thereby breaking tolerance and initiating autoimmunity. In this study, we demonstrated that abatacept reduced the clinical scores following challenge. Abatacept also prevented breach of B cell tolerance, inhibiting the generation of anti-CII IgG Abs. Strikingly, abatacept also reduced anti-OVA IgG2a recall responses. In addition,

![FIGURE 6. Abatacept prevents the breach of self-tolerance in experimental arthritis. A, pLNs from unchallenged, challenged/PBS, challenged/control IgG, and challenged/abatacept mice were harvested 7 d after footpad challenge. Mice received PBS/control IgG/abatacept i.p. on days −2, 0, 2, 4, and 6, with HAO injection on day 0. HAO was given alone by footpad injection. Total LN cells were restimulated in vitro with medium or OVA. Results shown are the mean percentage ± SD of total CD4+ cells that had incorporated EdU (5-ethynyl-2’-deoxyuridine) for five mice per group. B, Percentage of total CD4+ cells that was CXCR5+ in each condition. Serum samples were assayed for the presence of total anti-CII IgG (C) and anti-OVA IgG2a (D) by ELISA. E and F, The percentage of B220+ cells that was Fas+GL7+ was assessed; results are shown as representative staining of B220+ gated cells (E) and the mean percentage ± SD of B220+ cells that were Fas+GL7+ for five mice per group (F). Results are representative of two independent experiments. **p < 0.01; ***p < 0.001; Bonferroni-corrected two-way ANOVA (C, D) or a Student two-tailed unpaired t test (A, B, F). ns, not significant.](http://www.jimmunol.org/)

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abactept dramatically reduced the number of GC B cells in the LN draining the affected joint, providing a reason for the diminished anti-OVA and anti-CD11b Ab levels.

Abactept has proven efficacious in the treatment of RA, but its mode of action in vivo remains to be fully elucidated. Its effects on specific T cell responses and the development of specific T-dependent B cell responses in vivo remained unclear, as were its effects on memory cell responses and the breach of tolerance to self, joint-associated Ags. We provide novel data that directly inform these questions. Important effects of abactept on the development of TFH cells and the development of B cell responses were observed. Abactept prevents full T cell proliferation and activation; importantly, it also blocks follicular migration of T cells. In turn, cognate B cells, although able to undergo normal clonal expansion, have a reduced capacity to produce Abs. The inhibition of B cell responses also occurs during experimental arthritis, in which the endogenous breach of B cell self-tolerance is prevented, as reflected by reduced GC formation and reduced levels of autoantibodies. Together, these experiments provide, for the first time, a mechanistic insight into how abactept suppresses T cell responses, as well as T-dependent B cell responses, in vivo. This, in turn, provides strong rationale for early application of abactept in the disease course of RA.

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Disclosures

The authors have no financial conflicts of interest.

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