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Research Article

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Implications for the Role of Cognate Interactions in In Vitro Human B Cell Activation by *Staphylococcus aureus* Cowan I and Pokeweed Mitogen

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Abstract

Human B cell-triggering mechanisms were investigated using the polyclonal activators *Staphylococcus aureus* Cowan I (SAC) and pokeweed mitogen (PWM). When the cultures of B cells, T cells, and monocytes were stimulated for 5 d by SAC or PWM, B cells could be activated by both mitogens to proliferate and secrete Ig. Even when T cells were substituted by T cell-derived soluble factors, SAC-stimulated B cells could differentiate into Ig-secreting cells. In contrast, interactions of B and T cells for at least the first 6 h of culture were necessary for the B cell triggering by PWM.

Experiments that allow a more precise delineation of the B cell-triggering mechanisms by PWM demonstrated that interactions of B cells with T4⁺ but not T8⁺ cells are required for the B cell triggering; anti-Ia or anti-T4 antibody can block this triggering; in contrast, anti-T3 or anti-T8 antibody do not exert any effects on the B cell triggering. However, all these monoclonal antibodies could not modulate the ability of B cells that had been already activated by PWM to respond to T cell-derived factors. These data suggest that SAC can directly activate B cells, while cognate interactions between Ia-like antigens on B cells and T4⁺ cells are essential for B cell triggering by PWM. Furthermore, once B cells are triggered, they will proliferate, differentiate, and secrete Ig in response to T cell-derived factors; Ia-like antigens or T cell differentiation antigens may not be involved in the processes in this cascade.

Introduction

Recent studies of the events leading to B cell activation, proliferation, and differentiation have documented that occupancy of membrane-bound Ig by antigen and/or cognate T cell interactions involving the Ia antigens of resting B cells results in functional expression of receptors for T cell-derived soluble factors, and that it is these factors that in turn stimulate B cells to proliferate and differentiate into cells secreting Ig (1-5). Many studies regarding human B cells have used polyclonal B cell activators. These polyclonal activators include mitogens such as a T cell-independent pure B cell mitogen *Staphylococcus aureus* Cowan

I (SAC)¹ that directly acts on B cells by cross-linking of the Ig receptors on their surfaces and anti-Ig antibody that interacts with the B cell surface Ig. The results of studies with these polyclonal activators suggest that SAC or a high concentration of anti-Ig antibody directly induces the proliferation of resting human B cells, while a less powerful surface-membrane Ig-mediated signal, such as that delivered by low concentrations of anti-Ig antibody, results in the activation of B cells without subsequent proliferation. However, the induction of proliferation by these activated B cells can occur in the presence of B cell stimulatory factors (BSF) (5-10). Moreover, the proliferating B cells that have been activated by SAC, a high concentration of anti-Ig antibody, or low concentrations of anti-Ig antibody plus BSF, can in turn be induced to differentiate into Ig-secreting cells by B cell differentiation factors (BCDF) (5-10).

Many of the great advances in our understanding of the regulatory mechanisms that control human Ig synthesis have emerged from studies that have used another polyclonal activator, pokeweed mitogen (PWM) (11-15). PWM is a T cell-dependent T cell and B cell mitogen and induces proliferation and differentiation of B cells into Ig-secreting cells under the adequate help of T cells (11, 15). In the present study, we have addressed the question of whether some differences exist in the B cell-triggering mechanisms by SAC and PWM. We have also investigated the role of interactions of Ia-like antigens on B cells with T4 antigens on T4⁺ cells in the B cell-triggering mechanisms by mitogens. Our results clearly delineate that SAC stimulation of resting B cells results by itself in functional expression of receptors for T cell-derived soluble factors and that cognate T cell interactions involving the Ia-like antigens of B cells are required for the induction of B cells to express such functional receptors by PWM.

Methods

Cell separations. Peripheral blood mononuclear cells from healthy volunteer donors were separated into T cells and non-T cells by means of a sheep red blood cell (SRBC)-rosette technique. The rosetted T cells were further fractionated into T cell subsets, T4⁺ and T8⁺, by complement-mediated cell lysis with monoclonal antibodies OKT8 and OKT4 (Ortho Pharmaceutical Corp., Raritan, NJ), respectively. Analysis of the T cell subpopulations showed that the OKT8-treated population yielded >94% T4⁺ cells and <1% T8⁺ cells, whereas the OKT4-treated population contained >96% T8⁺ cells and <2% T4⁺ cells. B cells and monocytes were obtained from the unrosetted non-T cell fraction as follows: B cells were obtained by further depletion of T cells remaining in the non-T cell fraction by complement-mediated cell lysis with OKT3 antibody (Ortho Pharmaceutical Corp.). This was followed by depletion of mono-

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1. **Abbreviations used in this paper:** BCDF, B cell differentiation factors; BSF, B cell stimulatory factors; Con A, concanavalin A; IL-2, interleukin 2; MHC, major histocompatibility complexes; PFC, plaque-forming cell; PHA, phytohemagglutinin; PWM, pokeweed mitogen; SAC, *Staphylococcus aureus* Cowan I; SRBC, sheep red blood cell.

cytes by removal of cells adhering to petri dishes and by complement-mediated cell lysis with OKM1 antibody (Ortho Pharmaceutical Corp.). The resultant B cell population contained no T cells, <0.2% monocytes, and >90% cells bearing surface-membrane Ig. Moreover, the B cells obtained did not respond at all to phytohemagglutinin (PHA; Wellcome Research Laboratories, Beckenham, England) and concanavalin A (Con A; Pharmacia Fine Chemicals, Piscataway, NJ) at their optimal concentrations. Monocytes were obtained by collecting the cells adhering firmly to petri dishes. All these purification procedures were repeated twice and have been described in detail elsewhere (16–18).

Preparation of partially purified T cell-derived soluble factors. Fresh, pooled human peripheral mononuclear cells (2.5×10^6 /ml) were stimulated for 48 h with PHA (1 μ g/ml). The supernatants were collected, purified by several chromatographic steps as described elsewhere (19, 20), and were designated partially purified T cell-derived soluble factors. The characterization of the factors obtained has been described previously (18–20). The factors obtained had neither PHA activity nor interferon γ activity; they contained BSF activity and BCDF activity as well as interleukin 2 (IL-2) activity.

Overall experimental design. Activated B cells were generated by PWM stimulation in a first culture for 6 h. These cells were washed four times to remove PWM, and further incubated in the presence of partially purified T cell-derived soluble factors in a second culture. After incubation of 5 d in the second culture, cells were assayed either for their proliferative responses or for their plaque-forming cell (PFC) responses. In all studies, T cells and monocytes to be used in the first culture were obtained from the same normal donor who provided B cells.

Generation of activated B cells by stimulation with PWM (first culture). 5×10^6 B cells, 5×10^6 T cells, and 5×10^5 monocytes were incubated in 4-ml culture medium, RPMI 1640 (FLow Laboratories Inc., McLean, VA) supplemented with 10% fetal bovine serum (Flow Laboratories Inc.) with 4 μ g PWM (PWM-activated B cells), or without PWM (unactivated control B cells) at 37°C in a humidified 5% CO₂/95% air environment. 6 h later, the cells were harvested, washed four times, and B cells were isolated exactly as described above. These B cells were then incubated in a second culture (see below).

The basic first culture system was then perturbed by a number of different procedures. In some experiments, unfractionated T cells were substituted by T4⁺ or T8⁺ cells. Anti-Ia, anti-T3, anti-T4, or anti-T8 monoclonal antibody (Coulter Diagnostics, Coulter Electronics, Inc., Hialeah, FL) was also added to the first culture at a final concentration of 25 μ g/ml to investigate the influence of the respective monoclonal antibodies on the B cell-triggering mechanisms. Our preliminary experiments revealed that addition of 25 μ g/ml anti-T3 antibody at the initiation of culture resulted in complete abrogation of either PHA responses, Con A responses, or autologous and allogeneic mixed lymphocyte reactions by T cells. In addition, anti-T3 antibody at the above concentration could not induce T cell-proliferative responses at all. In other experiments, 5×10^5 monocytes were added to 1×10^7 B or T cells and each was individually prestimulated for 6 or 24 h with 1 μ g/ml of PWM. Thereafter, the individually prestimulated lymphocytes were washed extensively. The individually prestimulated T cells and individually prestimulated B cells, or individually prestimulated T cells and fresh B cells, or individually prestimulated B cells and fresh T cells, or fresh T cells and fresh B cells were paired and subjected to a first culture for 6 h in the presence or absence of PWM. B cells were then isolated from such first cultures and added to the second assay culture systems. Fresh lymphocytes to be used here were obtained 6 or 24 h later from a new bleeding of the same normal individual who originally provided the individually prestimulated lymphocytes. Purified fresh T cells and B cells that were paired with the individually prestimulated lymphocytes were prepared exactly as described above.

Assay for proliferative responses by PWM-activated B cells in a second culture. In order to examine proliferative responses of PWM-activated B cells in the first culture, a second culture was performed in 96-well, round-bottomed microtiter plates (Costar Data Packaging Corp., Cambridge, MA). Each culture contained 5×10^4 PWM-activated or unactivated control B cells from the first culture and 25% (vol/vol) of partially

purified T cell-derived soluble factors. All cultures were performed in a total volume of 200 μ l and were harvested on day 5. Proliferative responses were measured by the incorporation of [*methyl*-³H]thymidine (5 Ci/mmol; Amersham Corp., Arlington Heights, IL) over the last 20 h of the culture.

Assay for PFC responses by PWM-activated B cells in a second culture. The second culture to induce Ig-secreting cells was established in a 1-ml vol in 12 \times 75 mm plastic tubes (Falcon Labware Div., Becton-Dickinson & Co., Oxnard, CA) with 2×10^5 PWM-activated or unactivated control B cells from the first culture and 400 μ l partially purified T cell-derived soluble factors. After 5 d culture in 5% CO₂ in a humidified air atmosphere, cells were harvested, washed three times in balanced salt solution, and then assayed for their PFC responses.

When B cells were activated by PWM, IgG + IgA + IgM-producing cells were detected in a reverse hemolytic PFC assay (14, 16, 21) in almost all experiments. When B cells were stimulated with SAC, IgM secretion by B cells was measured with the reverse hemolytic PFC assay. This was done because the Ig binding characteristics of Staphylococcal protein A that SAC would display in an insoluble form to act, at least in part, as B cell activators, might interfere with reverse hemolytic PFC assay for IgG-producing cells (22). Hemolytic plaques were enumerated in triplicate and the results were expressed as the mean PFC/ 10^6 B cells in the original culture.

Results

Requirement of T-B cell interactions for Ig production of PWM-but not SAC-stimulated B cells. To determine whether direct T-B cell interactions are needed for human B cells to be activated by SAC or PWM to mature into Ig-secreting cells, or alternatively, to determine whether T cell-derived soluble factors could be substituted for T cells, B cells (2×10^5 /ml) plus monocytes (2×10^4 /ml) were cultured for 5 d with one of the polyclonal B cell activators, SAC (0.002% vol/vol) and PWM (1 μ g/ml), and either T cells (2×10^5 /ml) or T cell-derived soluble factors (40% vol/vol). Ig-secreting cells on day 5 were measured using reverse hemolytic PFC assays for IgM. As shown in Fig. 1, substantial PFC responses were observed by giving a stimulation with either SAC or PWM to a combination of B cells, T cells, and monocytes. Ig production equivalent to that in the presence of T cells was also observed by stimulating B cells and monocytes with SAC in the presence of T cell-derived soluble factors (Fig. 1). On the contrary, when PWM stimulation was given to a B cell plus monocyte combination, Ig production did not occur at all, even if T cell-derived soluble factors were present (Fig. 1). These results indicate that the induction of Ig production by PWM requires direct interactions of B cells with T cells and that T cell-derived soluble factors cannot substitute T cells.

Optimal conditions for B cell triggering by PWM in a first culture. To study whether the direct interactions of B cells with T cells throughout the course of culture are needed to activate B cells by PWM, or alternatively, whether contact of B and T cells for only a certain period of time is sufficient, B cells, T cells, and monocytes were stimulated with PWM in the first culture for various periods of time. PWM-stimulated B cells were then isolated by means of SRBC-rosetting technique, removal of cells adhering to petri dishes, and complement-mediated cell lysis using monoclonal antibodies, OKT3 and OKM1. These isolated B cells were further cultured for 5 d with or without T cell-derived soluble factors in the second culture to induce proliferative responses and PFC responses. When the first culture was performed for <3 h, very few PFC responses were observed, even when T cell-derived soluble factors were added to the second culture (Fig. 2). However, substantial PFC responses were

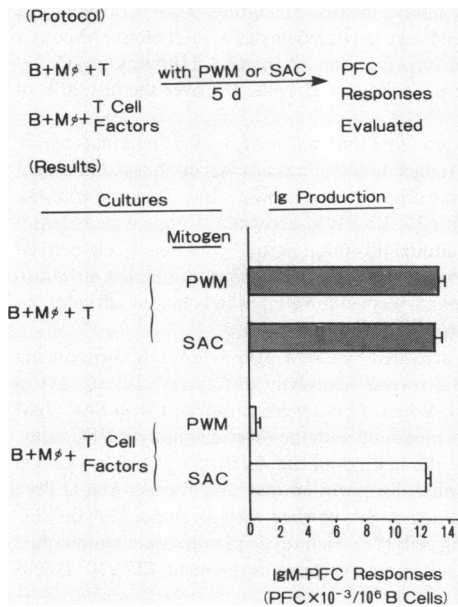


Figure 1. Requirement of T cells for PWM-induced, but not SAC-induced, Ig production: summary of the results from eight different normal donors. B cells (2×10^5 /ml) and monocytes (2×10^4 /ml) were stimulated with SAC (0.002% vol/vol) or PWM (1 μ g/ml) in the presence of the following: ■, T cells (2×10^5 /ml); and □, partially purified T cell-derived soluble factors (40% vol/vol). The number of cells secreting IgM was measured on day 5 using reverse hemolytic PFC assays. The mean PFC responses \pm SEM are shown.

induced when B cells stimulated with PWM in the first culture for more than 6 h were further incubated in the second culture in the presence of T cell-derived factors (Fig. 2). In data not

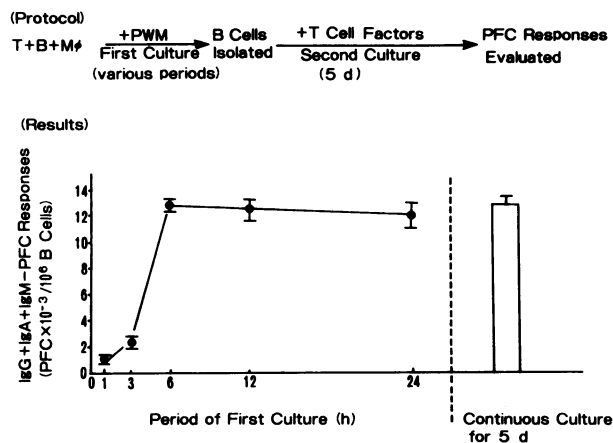


Figure 2. Necessary period of time in the direct contact of B and T cells for B cell triggering by PWM. 5×10^5 B cells, 5×10^6 T cells, and 5×10^5 monocytes were stimulated for various period of time with PWM (1 μ g/ml) in the first culture. B cells were then isolated from the first culture by means of SRBC-rosetting technique, removal of cells adhering to petri dishes, and complement-mediated cell lysis using OKT3 and OKM1 monoclonal antibodies. The isolated B cells (2×10^5 /ml) were further incubated with T cell-derived soluble factors (40% vol/vol) in the second culture. Cells secreting IgG + IgA + IgM were enumerated on day 5 by reverse hemolytic PFC assays. Data represent the mean PFC responses \pm SEM of four separate experiments using cells from four different normal donors.

shown, substantial proliferation in the second culture was also obtained from B cells stimulated with PWM for ≥ 6 h but not for < 3 h in the first culture. The results indicate that only direct contact of B and T cells for a certain period of culture is sufficient for the B cell triggering by PWM.

The next investigation was concerned with whether B cells are activated only in the coexistence of B cells, T cells, and PWM or whether B cells are activated and become sensitive to T cell-derived factors even when the first culture is established using B and T cells individually prestimulated by PWM. To this end, individually prestimulated B and T cells were prepared by stimulating B cells plus monocytes and T cells plus monocytes with PWM separately for 6 h. These individually prestimulated B and T cells, or individually prestimulated B or T cells and autologous fresh T or B cells were combined, incubated for 6 h in the presence or absence of PWM (first culture), and then B cells were isolated for the second culture containing T cell-derived factors. As shown in Fig. 3, only small PFC responses were observed when PWM was not present in the first culture of any of the combinations of individually prestimulated B plus T cells, individually prestimulated B cells plus fresh T cells, individually prestimulated T cells plus fresh B cells, and fresh B and T cells. However, when PWM was present in the first culture, substantial PFC responses could be induced in any of the combinations of individually prestimulated lymphocytes and/or fresh lymphocytes. Similar results were also obtained even when either T or B cells were individually prestimulated for 24 h with PWM. This indicates that B cell triggering by PWM does not occur even if individually prestimulated B and T cells coexist, but does occur only when B and T cells and PWM coexist.

T cell subsets involved in B cell triggering by PWM. To identify T cell subsets involved in the B cell triggering by PWM, B cell plus monocyte combinations added either with T4⁺ or T8⁺ cells were stimulated with PWM for 6 h in the first culture. Thereafter, B cells were isolated from the first culture and further incubated in the second culture for 5 d in the presence of T cell-derived soluble factors. As shown in Fig. 4, B cells cultured with T4⁺ cells in the first culture showed vigorous proliferative responses and PFC responses by reacting to T cell-derived soluble factors in the second culture. However, B cells incubated with T8⁺ cells in the first culture poorly responded in both proliferation and Ig production even if T cell-derived factors were present in the second culture. These results indicate that B cell triggering by PWM occurs only when T4⁺, but not T8⁺, cells are in direct interactions with B cells. The results presented here also suggest the possibility of B cell activation occurring through the interaction of T4 antigens on T cells and Ia-like antigens on B lymphocytes and/or monocytes, since T4 antigens are receptors that recognize class II antigens of the major histocompatibility complexes (MHC) (23).

Modulation by a series of monoclonal antibodies of B cell triggering by PWM. Anti-Ia, anti-T3, anti-T4, and anti-T8 antibodies were added to the first culture of B cells, T cells, monocytes, and PWM to examine the ability of the individual antibodies to modulate the mechanisms of B cell triggering (Fig. 5). When the first culture was performed in the presence of anti-Ia or anti-T4 antibody, proliferative as well as PFC responses induced in the second culture were remarkably decreased. However, when anti-T3 and anti-T8 antibodies were added to the first culture, both of the responses in the second culture were not inhibited at all. These results strongly support the above hypothesis that T4 antigens on T cells and Ia-like antigens on

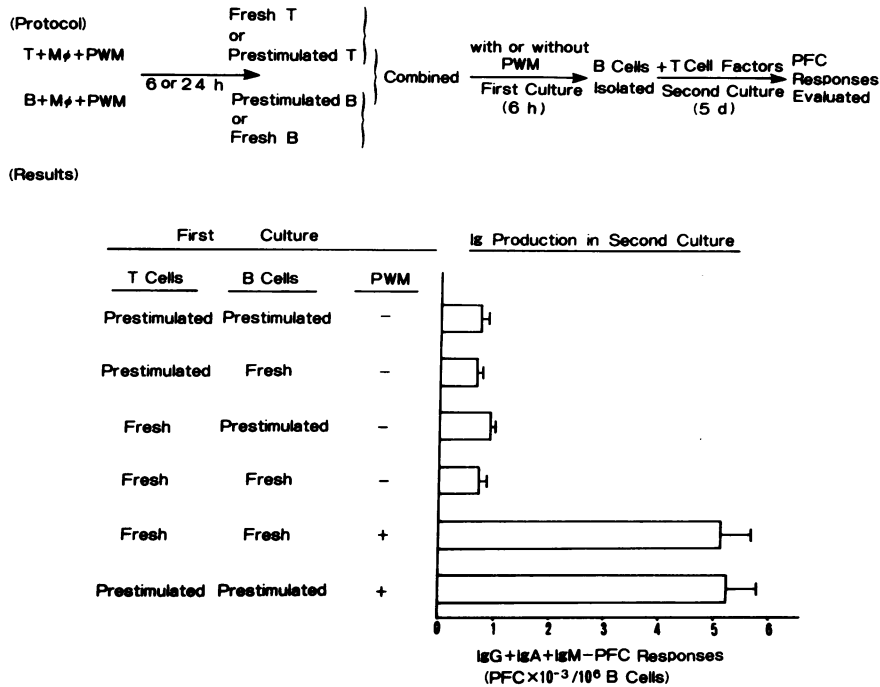


Figure 3. Requirement of the coexistence of B cells, T cells, and PWM for B cell triggering by PWM. 1×10^7 T or B cells together with 5×10^5 monocytes were individually prestimulated for 6 h with $1 \mu\text{g/ml}$ PWM. The cells were then washed four times to remove PWM. Either the individually prestimulated T cells and individually prestimulated B cells, individually prestimulated T cells and fresh B cells, fresh T cells and individually prestimulated B cells, or fresh T cells and fresh B cells were paired and incubated for 6 h with or without PWM in the first culture. Fresh T and B cells to be used in the first culture were obtained 6 h later from a new bleeding from the same normal individuals who originally provided the individually prestimulated lymphocytes. Thereafter the B cells were isolated and further incubated in the second culture in the presence of T cell-derived soluble factors. The number of cells secreting IgG + IgA + IgM was measured on day 5 using reverse hemolytic PFC assays. Data represent the mean PFC responses \pm SEM of four separate experiments using four different normal donors.

non-T cells or possibly B cells are involved in B cell-triggering mechanisms by PWM.

Effect of various monoclonal antibodies in the second culture.

B cells, T cells, and monocytes were incubated with PWM for 6 h in the first culture and B cells were then isolated. Anti-Ia, anti-T3, anti-T4, anti-T8, or anti-Tac antibodies were added to the second culture of the B cells isolated from the first culture plus T cell-derived soluble factors to examine the influence of these individual antibodies in the second culture on the proliferative and PFC responses. As shown in Fig. 6, even if these

various monoclonal antibodies were added to the second culture, once activated, B cells responded to T cell-derived factors sufficiently, and their proliferation and Ig production were not inhibited by the monoclonal antibodies.

Discussion

SAC can directly activate and induce a degree of proliferation of resting human B cells by cross-linking of the Ig receptors on B cell surfaces without requirement for the process of self rec-

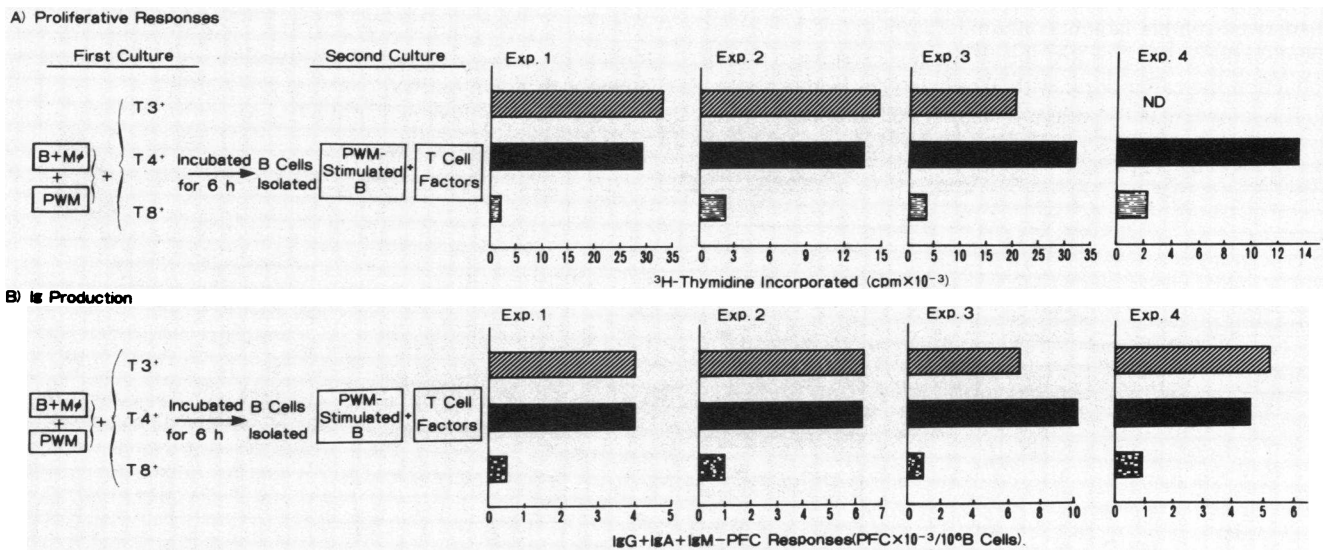


Figure 4. Requirement of T4 $^+$, but not T8 $^+$, cells for B cell triggering by PWM. 5×10^6 B cells and 5×10^5 monocytes were incubated for 6 h with $1 \mu\text{g/ml}$ PWM in the presence of the following: \blacksquare , 5×10^6 T3 $^+$ cells; \blacksquare , 5×10^6 T4 $^+$ cells; and \blacksquare , 5×10^6 T8 $^+$ cells (first culture). The B cells were then isolated from the first culture and further incubated for 5 d with partially purified T cell-derived soluble factors in the sec-

ond culture. Proliferative responses were measured by incorporation of [^3H]thymidine during the last 20 h of the second culture. Cells secreting IgG + IgA + IgM were measured by reverse hemolytic PFC assays. Similar results were obtained from three additional experiments.

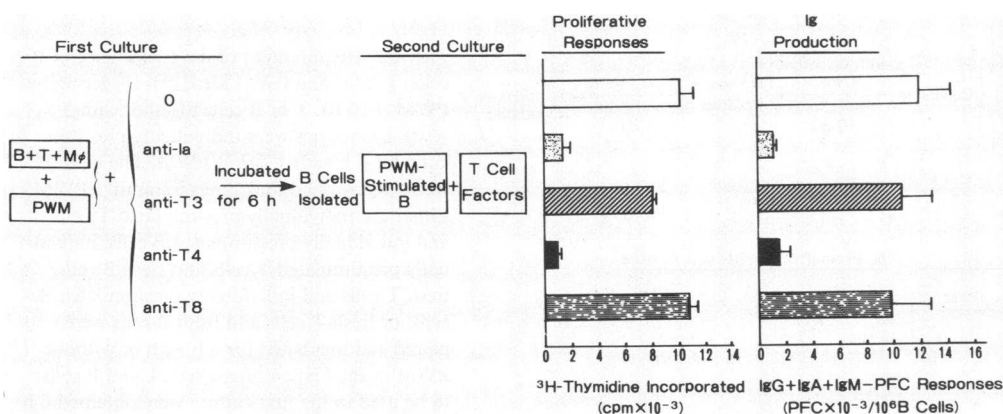


Figure 5. Effect of addition of a series of monoclonal antibodies to the first culture on the B cell triggering by PWM. 5×10^6 B cells, 5×10^6 T cells plus 5×10^5 monocytes were stimulated for 6 h with $1 \mu\text{g/ml}$ PWM in the first culture in the presence of the following: □, medium; ■, $25 \mu\text{g/ml}$ anti-Ia-like antibody; ▨, $25 \mu\text{g/ml}$ anti-T3 antibody; ▩, $25 \mu\text{g/ml}$ anti-T4 antibody; and ▧, $25 \mu\text{g/ml}$ anti-T8 antibody. The B cells were then isolated and further incubated for 5 d in the second culture in the presence of T cell-

derived soluble factors. Proliferative responses were measured by incorporation of [³H]thymidine during the last 20 h of the second culture. Cells producing IgG + IgA + IgM were measured by reverse hemolytic PFC assays. Data represent the mean responses+SEM of 10 separate experiments using cells from 10 different normal donors.

ognition through the interaction of T and B cells (7). Moreover, when T cell-derived soluble factors including IL-2, BSF, and BCDF coexist, a proliferative state of the SAC-stimulated B cells is much enhanced and sustained. In addition, these cells can be further induced to differentiate into Ig-secreting cells (8, 18). On the other hand, PWM activates B cells and induces their proliferation and Ig production if combined with adequate help of T cells (11). Our results also support the above results well. We have further extended those results to elucidate not only the precise B cell-triggering mechanisms but also complete activation mechanisms that drive a cell through proliferation as well as differentiation by PWM. We have demonstrated that the B cell triggering by PWM occurs through the interactions of B cells and T4⁺ cells, and that effective interactions occur only when the three are in direct contact. It was also clarified that the interactions occurring between T and B cells rest upon the recognition by T4⁺ cells of the Ia-like antigens on B cells and/or monocytes, that once B cells are triggered by PWM through such interactions, these activated cells become sensitive to T cell-derived soluble factors containing proliferative as well as differentiation signals, so that the proliferative responses and Ig production thereafter occur with T cell-derived factors alone.

It is known that T4⁺ cells are major T cell subsets that func-

tion as helper cells in the PWM-stimulated Ig production system (12, 13). T4⁺ cell plus B cell combinations induce much more Ig production than do T8⁺ cell plus B cell combinations (12, 13). We have provided evidence that the difference in helper activity between T4⁺ and T8⁺ cells is also observed in the B cell-triggering phase and that only B cells in intimate contact with T4⁺ cells are triggered and reach the proliferation and differentiation phases in the cascade of B cell-activation mechanisms. Suppressor or cytotoxic cells are mainly distributed in T8⁺ cells (13). To mature into effector cells of their precursors, T8⁺ precursors must go through the interactions with T4⁺ cells that bear the function as inducer cells (15, 24). Therefore, the failure of B cell activation in the T8⁺ cell plus B cell combinations could be due to the failure of T8⁺ cells to recognize B cells rather than to the suppressor or cytotoxic effect of T8⁺ cells. Difference in helper activity between T4⁺ and T8⁺ cells would also be expected to exist in the differential ability to produce T cell-derived soluble factors. In this regard, our preliminary experiments revealed that PWM-stimulated T4⁺ and T8⁺ cells produced equally active IL-2, BSF, and BCDF activities, suggesting that the difference in helper activity between the T4⁺ and T8⁺ subsets is mainly dependent on the B cell-triggering levels.

A series of monoclonal antibodies directed at differentiation

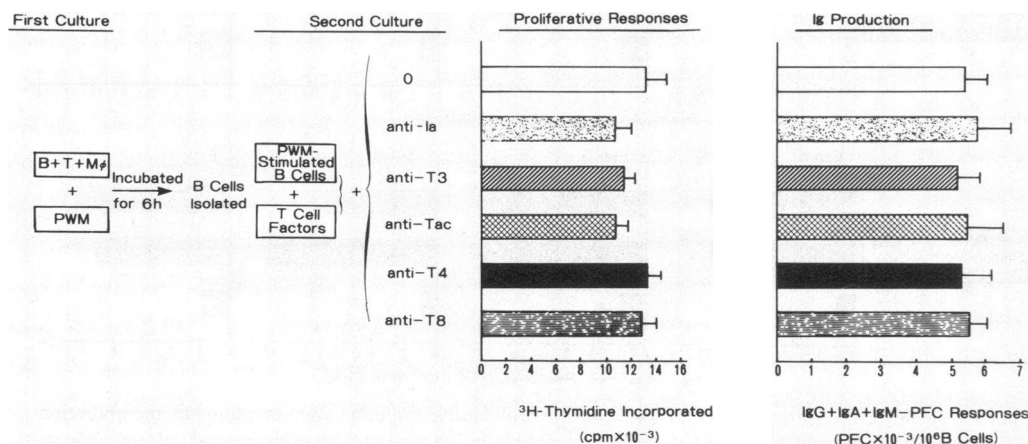


Figure 6. Effect of addition of a series of monoclonal antibodies to the second culture on the proliferation and differentiation of B cells that have been already triggered by PWM. B cells were first activated for 6 h with PWM in the first culture in the presence of T cells and monocytes. B cells isolated from the first culture were further incubated for 5 d with T cell-derived soluble factors in the second culture in the presence of the following: □, medium; ■, $25 \mu\text{g/ml}$ anti-Ia antibody; ▨, $25 \mu\text{g/ml}$ anti-T3 antibody; ▩, anti-Tac ascites at a final dilution of 1:1,000; ▧, $25 \mu\text{g/ml}$ anti-T4 antibody; and ▦, $25 \mu\text{g/ml}$ anti-T8 antibody. Proliferative responses were measured by incorporation of [³H]thymidine during the last 20 h of the second culture. Cells secreting IgG + IgA + IgM were measured by reverse hemolytic PFC assays. Data represent the mean responses+SEM of eight separate experiments using cells from eight different normal donors.

antibody; ▨, $25 \mu\text{g/ml}$ anti-T3 antibody; ▩, anti-Tac ascites at a final dilution of 1:1,000; ▧, $25 \mu\text{g/ml}$ anti-T4 antibody; and ▦, $25 \mu\text{g/ml}$ anti-T8 antibody. Proliferative responses were measured by incorporation of [³H]thymidine during the last 20 h of the second culture. Cells secreting IgG + IgA + IgM were measured by reverse hemolytic PFC assays. Data represent the mean responses+SEM of eight separate experiments using cells from eight different normal donors.

antigens on T or B lymphocyte surfaces have been used in recent years in the analysis of the role of these antigens in a complex series of immune responses. Monoclonal antibodies against T4 antigens, for instance, are known to selectively inhibit the helper activity or cytotoxic activity of T4⁺ cells (25, 26). It is also known that T4 antigens play an important role in lymphokine production, helper activity, or cellular interactions by which the proliferation and Ig production can be induced either in autologous mixed lymphocyte reactions or antigen-specific systems (27–29). Monoclonal antibodies against T8 antigens selectively interfere with the suppressor activity or cytotoxic activity of T8⁺ cells (25, 26). More recently, it has been further found that T4 and T8 antigens do recognize and respond to MHC antigens, T4 molecules bind to a constant region of a class II MHC gene product, and T8 antigens represent receptors for a constant region of a class I MHC gene product (23). On the other hand, the antibodies against T3 antigens not only act on T cells and induce their proliferation (30, 31) but also act to suppress functional maturation of T cells after antigenic stimulation (25, 32) as well as to inhibit the development of effector functions of cloned T cells (33). The antibodies against Ia-like antigens are also known to suppress both cellular immunity and humoral immunity well (4, 34, 35).

Our results also revealed suppressive effects by anti-Ia-like antibodies and anti-T4 antibodies when added to the first culture. Probably the occurrence of direct interaction of Ia-like antigens on B cells and T4 antigens on T4⁺ cells is essential to B cell triggering and both of the antibodies might have blocked the interaction and consequently the activation of B cells. It is not clear from the results presented in this paper whether the interactions between T4 antigens and Ia-like antigens occurred between T and B cells, or whether anti-Ia antibodies acted on Ia-like antigens only on macrophages, but not on B cells, resulting in blocking the function of macrophages and subsequent activation of T cells (36), whereby B cells failed to evolve into Ig-producing cells. If the latter is the case, T cells that have been individually prestimulated with PWM in the presence of monocytes could be activated, such that if individually prestimulated T and B cells are combined without PWM (first culture), the B cells should be rendered sensitive to T cell-derived soluble factors during the first culture. However, such B cells from the first culture did not respond to exogenous T cell-derived soluble factors, so that these cells neither proliferated nor differentiated in the second culture (Fig. 3). Moreover, it has in general been known that T cell activation by mitogens, particularly PHA and Con A, requires the recognition of mitogens by the T3-Tin complexes as well as the recognition of Ia-like antigens on macrophages by the T4 antigens (30, 31). The results that anti-T3 antibodies did not block the B cell triggering by PWM (Fig. 5) indicate that the T cells that contact with PWM-stimulated B cells are not necessarily limited in activated cells but all that is necessary is for T4⁺ cells, either activated or unactivated, to recognize Ia-like antigens on B cells. Thus it is likely that the Ia-like antigens that T4 antigens recognize are, at least in part, those on B cells. It is known that anti-Ia antibodies can induce suppressor T cells (37) and that some activated T4⁺ cells function in the role of suppressor cells (14, 16, 38). However, it is very unlikely that suppressor function by either T4⁺ or T8⁺ cells that might be generated in the first culture might be mediated by anti-Ia antibodies, because anti-Ia antibodies should have first activated either T4⁺ or T8⁺ cells and rendered the profound suppressor activity to these cells in the course of 6 h in the first

culture. Moreover, these suppressor effector cells should have also inhibited the B cell triggering in the very same course of 6 h in the first culture.

The effect of a series of monoclonal antibodies in the second culture was also investigated. Once B cells were triggered in the first culture, they proliferated and went on to differentiate into Ig-secreting cells in the second culture, regardless of the presence of anti-Ia, anti-T3, anti-T4, anti-T8, or anti-Tac antibody. IL-2 receptors, Tac antigens, have been found to be expressed on B lymphocytes as well in recent years (39–41) and it has been documented that IL-2 exerts a direct effect on B cell function (41). As far as our results are concerned, Ig production was not inhibited even if anti-Tac antibodies were added to the second culture of PWM-activated B cells. Conceivable reasons for this phenomenon would be (a) the possibility that BSF and BCDF as well as IL-2 would be enriched in our preparations of T cell-derived soluble factors and that complementary proliferation and differentiation by BSF and BCDF would occur even though the proliferation through interactions of IL-2 and IL-2 receptors is impeded by anti-Tac antibodies and (b) the possibility that B cell subsets activated by PWM would mature in an IL-2-independent manner. Further study is now underway on this point. On the other hand, the fact that anti-Ia-like antibodies have no inhibitory effect in the second culture indicates that the responsiveness to T cell-derived factors of B cells that once have been activated and become sensitive to the factors could no longer be mediated by Ia-like antigens. Thus, the *in vivo* use of monoclonal antibodies to regulate immune responses, particularly activation steps of B cells, should require a careful evaluation to determine in which phase(s) of the activation cascade abnormal B cells reside and the selection of proper monoclonal antibodies, depending on the phase in this cascade. We hope that our results will provide new and potentially important avenues of approach to the clinical use of monoclonal antibodies in the future.

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