Qualitative Regulation of B Cell Antigen Receptor Signaling by CD19: Selective Requirement for PI3-Kinase Activation, Inositol-1,4,5-Trisphosphate Production and Ca^{2+} Mobilization

By Anne Mette Buhl,* Christopher M. Pleiman,§ Robert C. Rickert,* and John C. Cambier‡

From the *Division of Basic Sciences, Department of Pediatrics, National Jewish Medical and Research Center, 1400 Jackson Street, Denver, Colorado 80206; ‡Department of Immunology, University of Colorado Health Sciences Center, Denver, Colorado 80206; §Cadus Pharmaceuticals, 1601 Pierce St., Suite 110, Lakewood, Colorado 80214; and *Department of Biology, University of California San Diego, 9500 Gilman Drive, La Jolla, California 92093-0322

Summary

Genetic ablation of the B cell surface glycoprotein CD19 severely impairs the humoral immune response. This requirement is thought to reflect a critical role of CD19 in signal transduction that occurs upon antigen C3dg coligation of antigen receptors with CD19 containing type 2 complement receptors (CR2). Here we show that CD19 plays a key accessory role in B cell antigen receptor signaling independent of CR2 coligation and define molecular circuitry by which this function is mediated. While CD19 is not required for antigen-mediated activation of receptor proximal tyrosine kinases, it is critical for activation of phosphatidylinositol 3-kinase (PI3-kinase). PI3-Kinase activation is dependent on phosphorylation of CD19 Y484 and Y515. Antigen-induced CD19-dependent PI3-kinase activation is required for normal phosphoinositide hydrolysis and Ca^{2+} mobilization responses. Thus, CD19 functions as a B cell antigen receptor accessory molecule that modifies antigen receptor signaling in a qualitative manner.

CD19 is a B-lymphocyte–specific transmembrane glycoprotein that is expressed from the pro-B cell until the plasma cell stage. CD19 appears to function in multiple capacities. It is a component of the type 2 complement receptor (CR2) complex that contains CD19, CD21 (CR2), CD81, (TAPA-1), and Leu-13. CR2 is expressed by mature B cells and is activated upon binding of C3dg (reviewed in reference 1). Evidence indicates that CD19 may also be a constitutive component of the B cell antigen receptor (BCR) complex and is phosphorylated upon BCR crosslinking (2). Finally, although its ligand has not been identified, CD19 appears to function as a transducer in pro- and pre-B cells where it is expressed in the absence of CD21 and the BCR. The function of CD19 in these various contexts is poorly understood.

CD19 is a highly conserved protein containing two immunoglobulin-like domains in its extracellular aspect and nine potential tyrosine phosphorylation sites in its cytoplasmic tail (3). It is tyrosine phosphorylated upon cell activation in all three of the signaling contexts described above, presumably by Lyn, Fyn, or Lck associated with unligated CD19 (4–8). After phosphorylation, tyrosines Y^{484} and Y^{515} motifs have been shown to mediate the association of the phosphoinositide hydrolysis and Ca^{2+} mobilization responses. Thus, CD19 functions as a B cell antigen receptor accessory molecule that modifies antigen receptor signaling in a qualitative manner.

Abbreviations used in this paper: BCR, B cell antigen receptor; IP_{3}, inositol-1,4,5-trisphosphate; ITAM, immunoreceptor tyrosine based activation motif; NP, nitrophenol; PH, pleckstrin homology; PI, phosphatidylinositol; PI3-kinase, phosphatidylinositol 3-kinase.
phorylation (17, 18), PLC-activation and calcium mobilization (19). Colligation also lowers the threshold for antigen stimulation of B cell proliferation (20) and enhances by \(\geq 10,000\)-fold the immunogenicity of antigen–complement complexes (HEL–C3d) (21).

Antibody crosslinking of CD19 on certain immature B cell lines that do not express IgM has been shown to induce activation of Src-family kinases and tyrosine phosphorylation of CD19 and Vav (5, 13, 22). In this context CD19 crosslinking also results in PLC-activation, calcium mobilization and activation of the MAP kinase cascade (10, 19, 23).

Finally, anti-Ig stimulation of the BCR without CD19 crosslinking results in tyrosine phosphorylation of CD19, indicating that CD19 is a target of BCR-activated kinases (4, 5, 13). Also, anti-IgM stimulation of Daudi B cells leads to an increase in CD19 association with PI3-kinase (9). Consistent with this finding, several studies indicate that CD19 may be physically associated with the BCR. Incubation of B cells with Ig-specific mAb leads to Ig and CD19 comodulation and cocapping (24, 25). In Daudi B cells CD19 can be immunoprecipitated with IgM (2). Whether CD21, Leu-13 and CD81 are required for the constitutive association of CD19 with the BCR is unknown; however CD19 expression levels in mature B cells appear to be about fivefold higher than CD21 expression levels (14) suggesting that CD19 occurs in both CD21-associated and -nonassociated forms.

Recent production of CD19 knockout mice have provided an opportunity to analyze B cell function in the absence of CD19 (26, 27). B cell development appears normal in these mice, but humoral immune responses are defective and B cells are unresponsive to antigen receptor stimulation as indicated by proliferation.

In this study we have investigated the constitutive role of CD19 in BCR-mediated signal transduction using CD19-deficient and -sufficient J558L mice. Our results demonstrate a critical role for CD19 in antigen-mediated BCR activation of multiple signaling events. CD19 expression is required for detectable BCR-mediated PI3-kinase activation and for maximal phosphoinositide hydrolysis and calcium mobilization. The latter effects appear causally linked to failed PI3-kinase activation. Other more subtle effects of CD19 on BCR signaling include increased antigen-mediated Lyn and Syk tyrosine phosphorylation, but these are not attributable to enhanced PI3-kinase activation.

Materials and Methods

Reagents and Antibodies. The murine B cell plasmacytoma J558L was provided by M. Reth (Max-Planck Institute for Immunology, Freiburg, Germany). We transfected this cell line previously to obtain a mouse full-length (B220) CD45 + variants and primary B lymphocytes from CD19 deficient mice. The human CD19 cDNA employed was provided by T. Tedder (Duke University Medical Center, Durham, NC). Rabbit polyclonal antibodies against CD19, Ig \(\alpha\), Syk, and Lyn, were prepared using GST fusion proteins produced in bac-
trifuged for 5 min at 14,000 rpm. Cleared lysates were incubated with antibodies to signaling molecules and protein A-Sepharose or protein G-Sepharose at 4°C, washed four times in lysis buffer, boiled with reducing SDS sample buffer, and then fractionated on 10% SDS-PAGE. SDS-PAGE–fractionated proteins were subjected to electrophoretic transfer to PVDF membranes. Transfers were probed with antibodies and developed using the ECL detection system. Sequential immunoblotting for phosphotyrosine and effector was performed. For this purpose PVDF membranes were stripped in stripping buffer (100 mM 2-mercaptoethanol, 2% SDS, 62.5 mM Tris-HCl, pH 6.7) for 30 min at 54°C, washed and blocked before immunoblotting.

Flow Cytometric Analysis of [Ca^2+]i Mobilization. Cells were loaded with indo-1 AM and the intracellular [Ca^2+]i was monitored by flow cytometry (model 50H; Ortho Diagnostic Systems, Westwood, MA) as previously described (28). The mean [Ca^2+]i and percent of cells responding was determined with an appended data acquisition system and the MultiTIME software (Phoenix Flow Systems, San Diego, CA).

Measurement of Inositol Phosphate Generation. Inositol-1,4,5-trisphosphate generation was measured using a [3H] radioreceptor assay kit (DuPont-NEN) according to the manufacturer's instructions. Generation of total inositol phosphates was measured as follows: cells were labeled for 18 h in 2 μCi/ml of [2-3H(N)]-inositol in inositol-free medium containing dialyzed 5% FCS before being washed in inositol-free media and incubated with inositol-free media containing 20 mM LiCl for 5 min. Cells were then stimulated for 25 min, lysed in 2 ml of methanol/HCl (99:1), and then 2 ml of chloroform was added. The aqueous phase was extracted, applied to a formate anion-exchange column, and then inositol phosphates were eluted with elution buffer (0.1 M formic acid, 1.0 M ammonium formate). Total [3H] inositol incorporation was determined by measuring the radioactivity in the organic phase.

Assay for PI3-Kinase Activity. Cells were stimulated, lysed, and then immunoprecipitated for 1 h at 4°C using anti-p85 antibodies and protein A-Sepharose. PI3-kinase activity was measured as previously described (30). Samples were dried and reaction products resolved by thin-layer chromatography on precoated TLC Silica Gel 60 plates in CHCl3/MeOH/H2O (9:7:2). The incorporation of 32PO4 into phosphatidylinositol 3-phosphate was determined using a PhosphorImager.

Isolation of Splenic B Cells. Splenic B cells were prepared as previously described (31). In brief, spleens were excised from mice and cells were dispersed through a 100-μm mesh in IMDM. Red blood cells were lysed using Gey's solution. Total nucleated spleen preparations were depleted of T lymphocytes by complement-mediated lysis using HO13.4 and T24 antibodies and dense cells (ρ > 1.066) isolated by discontinuous Percoll density gradient centrifugation.

Results

Although tyrosine phosphorylation of CD19 after BCR ligation is well documented, the contribution of CD19 to BCR signaling is not defined. To study the role of CD19 in antigen-induced signal transduction we generated a wild-type CD19-positive and Y484/Y515 to F CD19-positive variant of the J558L/mm3CD451 plasmacytoma. The parent clone expresses a NP-specific BCR and CD45, but does not express CD19 (28, 32). Human CD19 (hCD19) was introduced into this clone by retroviral infection, and lines (polyclonal populations) were isolated by G418 selection and fluorescence activated cell sorting of cells. Flow cytometric analysis revealed similar expression levels of sIgM,
CD45 and CD19 in the plasmacytoma cell lines (Fig. 1). Human CD19 was utilized since antibodies to mouse CD19 were not available at the time the studies were undertaken, and human and mouse CD19 are highly homologous (79% in nucleotide sequence, 75% in amino acid sequence) particularly in the cytoplasmic domain (3, 33). Recent studies indicate that human CD19 complements B cell function in CD19\(^{−/−}\) mice (34). In addition, using a recently generated rat mAb, 1D3, specific for murine CD19 (35), Krop et al. (22) have demonstrated that murine CD19 shares with human CD19 an association with complement receptor CD21 and CD81, tyrosine phosphorylation, binding of PI3-kinase, and synergistic signaling with membrane IgM.

Antigen-stimulated Intracellular Ca\(^{2+}\) Release and Extracellular Ca\(^{2+}\) Influx Is Increased in Cells Expressing CD19. To investigate the differences in BCR signaling in the CD19-negative and -positive contexts, we loaded the J558Lμm3CD45\(^{−}\)CD19\(^{−}\) and J558Lμm3CD45\(^{−}\)CD19\(^{+}\) cells with indo-1 AM, and analyzed changes in intracellular free calcium concentration ([Ca\(^{2+}\)]\(_{i}\)) in response to antigen stimulation. Initially, this was done under conditions where the free calcium concentration in the medium ([Ca\(^{2+}\)]\(_{o}\)) was maintained at physiological levels of 1.3 mM. As shown in Fig. 2 (solid lines), antigen stimulation induced a rise in [Ca\(^{2+}\)]\(_{i}\) in both cell lines; in the CD19-positive cell line the maximal mean [Ca\(^{2+}\)] reached 430 nM while it was 260 nM in the CD19-negative cells. In the CD19-positive cell line, antigen stimulation also induced a much more prolonged increase in [Ca\(^{2+}\)]\(_{i}\) than in the CD19-negative clone. To determine whether intracellular or extracellular calcium was responsible for the respective [Ca\(^{2+}\)] increases, we added 3 mM EGTA at the time of initiation of analysis of [Ca\(^{2+}\)]\(_{i}\). This results in buffering of [Ca\(^{2+}\)]\(_{o}\) to a concentration of 65 nM (which corresponds to the [Ca\(^{2+}\)] of resting cells) and prevents influx of extracellular calcium into the cell. Comparison of antigen-induced responses of CD19-negative and CD19-positive cells in low [Ca\(^{2+}\)]\(_{o}\) revealed that influx is also dependent on the expression of CD19 (Fig. 2, dotted lines) indicated that optimal receptor-mediated release of calcium from intracellular stores requires CD19. Comparison of responses in high [Ca\(^{2+}\)]\(_{o}\) revealed that influx is also dependent on the expression of CD19 (Fig. 2, dotted lines). Antigen-stimulated Inositol-1,4,5-trisphosphate (IP\(_3\)) production is also dependent on CD19 expression. To study CD19 dependence of signaling events upstream from calcium mobilization, we first compared antigen stimulation of IP\(_3\) production in CD19-positive and -negative cells. IP\(_3\) binds to its receptors on the ER leading to release of Ca\(^{2+}\) into the cytoplasm. Emptying of the ER stores triggers entry of calcium from the extracellular space by a mechanism known as capacitative Ca\(^{2+}\) entry (36). Thus, CD19 depen-
Antigen stimulation of PI3-kinase activation is severely impaired in the CD19-negative J558L μm3CD45+ plasmacytoma (41). As shown in Fig. 4, CD19 expression led to modest increases in antigen-induced tyrosine phosphorylation of Ig-α, Lyn, Syk, PLCγ1, and PLCγ2. Quantitation of tyrosine phosphorylation by scanning densitometry (at least three independent experiments for each effector enzyme) and normalization to effector levels revealed increased phosphorylation of Ig-α, Lyn, and Syk by 1.5–2-fold in the CD19-positive line. Induced PLCγ1 and PLCγ2 tyrosine phosphorylation was 2.5–3-fold higher in the CD19-positive cell line than in the CD19-negative cell line. These results suggest that CD19 functions in part by increasing the magnitude of certain proximal BCR coupled tyrosine phosphorylation events.

Antigen stimulation of PI3-Kinase Activation Is Severely Impaired in the CD19-negative J558L μm3CD45+ Plasmacytoma. CD19 contains two cytoplasmic YXXM motifs that mediate its association with the p85 subunit of PI3-kinase after phosphorylation of the tyrosines (9). The phosphorylation of these tyrosines may contribute to BCR-mediated PI3-kinase activation. We determined the activity of PI3-kinase in the cell lines by measuring the PI3-kinase activity associated with anti-p85 immunoprecipitates after stimulation with NP6BSA for 2 min. The results shown in Fig. 5 demonstrate a 250% increase in PI3-kinase activity in the CD19-positive cell line after antigen stimulation, but only a 40% increase in the CD19-negative variant. The residual...
activation of PI3-kinase in the absence of CD19 expression is probably a result of PI3-kinase activation through interaction with SH3 domains of Src-family kinases (42). The results described above are summarized in the two first columns of Table 1. It is clear that CD19 influences multiple intracellular signaling events. Some signaling events are only slightly increased in the CD19-positive cell line, (i.e., Ig-α, Lyn, and Syk tyrosine phosphorylation) while others are almost totally dependent on expression of CD19 (i.e., IP₃ generation, calcium influx, and PI3-kinase activation).

Wortmannin Inhibits Antigen-induced [Ca²⁺]ᵢ Mobilization and IP₃ Generation in the J558Lm3CD45⁺CD19⁺ Plasma­cytoma. To begin to dissect the relationship among the observed CD19 effects, we assessed the effects of the PI3-kinase inhibitor wortmannin on signaling events. Wortmannin has been shown to inhibit PI3-kinase activity by covalent modification of a lysine residue in the PI3-kinase catalytic subunit, p110 (43). We also used the competitive PI3-kinase inhibitor Ly294002 (44) and obtained similar results as in our studies with wortmannin (results not shown). The effect of wortmannin on antigen-induced activation of PI3-kinase was determined. The results shown in Fig. 6 A demonstrate the inhibitory effect of wortmannin on antigen-mediated PI3-kinase activation in the J558Lμm3CD45⁺CD19⁺ plasmacytoma. Cells were pretreated for 30 min with DMSO (vehicle) or increasing concentrations of wortmannin, stimulated with antigen for 2 min, washed, and then lysed before PI3-kinase was immunoprecipitated and activity measured in an in vitro kinase assay with phosphatidylinositol (P1) containing vesicles as substrate. Consistent with its previously established IC5₀, 10 nM wortmannin inhibited the antigen-mediated PI3-kinase activation by 54%, and 25 nM wortmannin inhibited 75% of antigen-induced PI3-kinase activity. Wortmannin added to the cells immediately before washing and lysis did not effect the PI3-kinase activity.
Table 1. Phenotypic Differences Between J558L \( \mu m3 \)CD 45- CD19- and J558L \( \mu m3 \)CD 45+ CD19+ Cell Lines

<table>
<thead>
<tr>
<th>Signal transduction event</th>
<th>CD19+</th>
<th>CD19-</th>
<th>25 nM wortmannin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ig-( \alpha ) Tyr Phos</td>
<td>++++</td>
<td>++</td>
<td>+</td>
</tr>
<tr>
<td>Lyn Tyr Phos</td>
<td>++++</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Syk Tyr Phos</td>
<td>++++</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>PLC( \gamma )/PLC( \gamma 2 ) Tyr Phos</td>
<td>++++</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>IP3 production</td>
<td>++++</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>[Ca( ^{2+} )] mobilization</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Intracellular stores</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Influx</td>
<td>+++</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>PI3-Kinase activation</td>
<td>++++</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

(data not shown) indicating that the inhibitor was acting intracellularly.

To investigate the influence of PI3-kinase on the BCR-mediated calcium mobilization seen in the CD19-positive cell line, we preincubated the cells with increasing concentrations of wortmannin and measured the antigen-induced increase in [Ca\( ^{2+} \)]. As shown in Fig. 6B, wortmannin strongly inhibited Ca\( ^{2+} \) mobilization in the J558L \( \mu m3 \)CD 45- CD19- cells. Sensitivity to wortmannin was also evident in the CD19-positive response. A wortmannin concentration of 25 nM strongly inhibited IP3 production in the J558L \( \mu m3 \)CD 45+ CD19+ cell line after 1 minute of antigen stimulation (Fig. 6C). In both cases wortmannin at a concentration of 25 nM converted the phenotype of the CD19-positive cells essentially to that of the CD19-negative cells. This result was consistent with the 75% inhibition of PI3-kinase activation by 25 nM wortmannin in Fig. 6A and indicates that the CD19 requirement for IP3 production and calcium mobilization may be attributable to PI3-kinase activation. Interestingly, wortmannin inhibited the residual antigen-stimulated PI3-kinase activation, IP3 production, and Ca\( ^{2+} \) mobilization of the CD19-negative cell line (data not shown) consistent with the possible absolute dependence of these responses on PI3-kinase activation.

Induced Tyrrosine Phosphorylation of Intracellular Effector Enzymes Is Insensitive to Wortmannin. To investigate the potential role of CD19-activated PI3-kinase on early BCR-mediated protein tyrosine phosphorylation events, CD19-positive cells were preincubated for 30 min with DMSO (vehicle) or increasing concentrations of wortmannin and stimulated for 1 min with antigen before effectors were immunoprecipitated and analyzed. As shown in Fig. 7, antigen-induced tyrosine phosphorylation of Ig-\( \alpha \), Lyn, and Syk was insensitive to wortmannin. PLC\( \gamma 1 \) and PLC\( \gamma 2 \) tyrosine phosphorylation was somewhat sensitive to wortmannin preincubation although the decrease in induced tyrosine phosphorylation was modest (35% inhibition at 25 nM wortmannin, Table 2) compared with the degree of wortmannin inhibition of IP3 generation and calcium mobilization (91.5% inhibition, Table 2). It seems very unlikely that the >90% inhibition of PLC\( \gamma \) activity could be caused by a 35% inhibition of PLC\( \gamma \) phosphorylation. Thus, as summarized in Table 1, CD19 expression by the J558L \( \mu m3 \)CD 45+ plasmacytoma is required for optimal antigen stimulation of multiple intracellular signaling events, but only some of these are secondary to the nearly absolute dependence of PI3-kinase activation on CD19.

Mutation of p85 Binding Sites Decreases the Ability of CD19 to Mediate an Increase in BCR-mediated Ca\( ^{2+} \) Mobilization. To further explore the requirement of CD19 for PI3-kinase activation and optimal antigen-mediated Ca\( ^{2+} \) mobilization, we introduced a double-point mutant of CD19, CD19Y484F,Y515F, into the J558L \( \mu m3 \)CD 45+ plasmacytoma. The expression levels of sIgM, CD45 and CD19 (Y484F,Y515F) in this cell line is shown in Fig. 1. It has previously been demonstrated that the mutations Y484F and Y515F abrogates the ability of CD19 to interact with the p85-subunit of PI3-kinase (9). Like the CD19-negative cell line, the CD19 mutant cell line, showed no significant anti-

---

Table 2. Anti-CD19 and Wortmannin Sensitivity of PI3-Kinase and PIP2 Synthase Activity

<table>
<thead>
<tr>
<th>dition of IP3 generation</th>
<th>25 nM wortmannin</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD19+</td>
<td>CD19-</td>
</tr>
<tr>
<td>p85-subunit of PI3-kinase</td>
<td>9</td>
</tr>
<tr>
<td>Wortmannin inhibition</td>
<td>91.5%</td>
</tr>
<tr>
<td>CD19 requirement for IP3 production</td>
<td>75%</td>
</tr>
<tr>
<td>Ca( ^{2+} ) mobilization</td>
<td>91.5%</td>
</tr>
</tbody>
</table>

---

Figure 6. Wortmannin inhibits the antigen-induced increase in PI3-kinase activity, [Ca\( ^{2+} \)], mobilization and IP3 generation in the J558L \( \mu m3 \)CD 45+ plasmacytoma. (A) Cells were preincubated with DMSO (vehicle) or wortmannin at various concentrations (in 0.2% final DMSO concentration) for 25 min at 25\( ^\circ \), prewarmed for 4 min at 37\( ^\circ \), and then stimulated for 2 min with 1 \( \mu g \) NP-BSA/4 \( \times 10^6 \) cells/ml before analysis of PI3-kinase activation as described in Fig. 4. Shown is the mean fold increase in activity over basal from four independent experiments ± standard error of the mean. Asterisk indicates a statistical significance of P < 0.05 determined by JMP 3.16 statistical software. (B) To study the effects of wortmannin on the calcium response, J558L \( \mu m3 \)CD 45+ CD19- cells were Indo-1 AM loaded and preincubated with vehicle or wortmannin at various concentrations (resulting in 0.2% final DMSO) for 25 min at 25\( ^\circ \), prewarmed for 4 min at 37\( ^\circ \), and stimulated at arrow with 250 ng NP-BSA/10\( ^6 \) cells/ml. Shown is the mean [Ca\( ^{2+} \)] over time with analysis of ~600 cells/s. [Ca\( ^{2+} \)] in resting cells is 100 nM. (C) J558L \( \mu m3 \)CD 45+ CD19+ cells were preincubated with DMSO or increasing concentrations of wortmannin, prewarmed, and stimulated with 2.5 \( \mu g \) NP-BSA/10^7/ml for 1 min before IP3 production was measured by the IP3 receptor binding inhibition assay. Shown is the mean fold increase over basal in three independent experiments ± standard error of the mean. Asterisk indicates a statistical significance of P < 0.05 determined by JMP 3.16 statistical software.
gen-mediated activation of PI3-kinase (results not shown). When we assayed the ability of the cell line expressing the mutant CD19 to mobilize $[Ca^{2+}]_i$ in response to antigen stimulation, we found that the mutant CD19 cell line behaved equivalently to the J558L $\mu m3CD45^+$ cell line (Fig. 8). This suggests that both CD19-mediated increase in BCR-induced PI3-kinase activation and $Ca^{2+}$ mobilization require phosphorylation of CD19 Y484 and Y515.

Wortmannin Inhibits Antigen-stimulated $[Ca^{2+}]_i$ Mobilization in Splenic B Cells. To investigate the role of PI3-kinase in antigen-induced signaling in a more physiological system, we purified splenic B cells ($r > 1.066$) from spleens of normal mice and assessed the effects of wortmannin on BCR signaling. After preincubation for 30 min with DMSO or wortmannin, splenic B cells were stimulated with 13.2 $\mu g/10^6$ cells $F(ab')_2RAMIG$. As observed in the J558L $\mu m3CD45^+$ cell line, wortmannin inhibited, as a function of its concentration, the antigen-mediated calcium mobilization in these primary B cells (Fig. 9). Wortmannin also inhibited the residual antibody-mediated $Ca^{2+}$ mobilization in CD19 $^{-/-}$ splenic B cells (data not shown). Thus, BCR signaling in J558L $\mu m3CD45^+CD19^+$ and normal B cells is similarly sensitive to PI3-kinase inhibition and CD19 function (see below).

CD19 Is Required for Anti-BCR Stimulation of PI3-Kinase Activation and $Ca^{2+}$ Mobilization in Splenic B Cells. To extend our observations regarding CD19 dependence of BCR-mediated signaling in B lymphocytes we compared BCR-mediated PI3-kinase activation and calcium mobilization responses of B cells from CD19 knockout mice and normal mice. It has previously been reported that no distinguishable difference can be seen between CD19-deficient and normal B cells in the pattern or the degree of substrate phosphorylation after surface immunoglobulin stimulation (27). We obtained the same results when comparing whole cell lysates from CD19 $^{+/+}$ and CD19 $^{-/-}$ cells (results not shown). However, careful investigation of several signaling events revealed that $r > 1.066$ resting splenic B lymphocytes from CD19 $^{-/-}$ mice are impaired in their response to surface immunoglobulin crosslinking when compared with CD19 $^{+/+}$ littermates. $F(ab')_2RAMIG$-mediated PI3-kinase activation was severely diminished in the CD19 $^{-/-}$ B cells (Fig. 10), reminiscent of the situation in the plasmacytoma cell lines. As shown in Fig. 11, absence of CD19 also clearly affects

Table 2. Percent Inhibition of Maximal Responses by Wortmannin Preincubation

<table>
<thead>
<tr>
<th>Wortmannin(nM)</th>
<th>$IP_3$ production</th>
<th>PLC-$\gamma_1$ tyrosine phosphorylation</th>
<th>PLC-$\gamma_2$ tyrosine phosphorylation</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 nM</td>
<td>41.7 ± 10.8% ($n = 2$)</td>
<td>18.2 ± 5.6% ($n = 2$)</td>
<td>16.3 ± 4.4% ($n = 6$)</td>
</tr>
<tr>
<td>25 nM</td>
<td>91.5 ± 8.6% ($n = 2$)</td>
<td>34.6 ± 11.4% ($n = 3$)</td>
<td>34.9 ± 7.4% ($n = 5$)</td>
</tr>
<tr>
<td>50 nM</td>
<td>97.6 ± 2.5% ($n = 2$)</td>
<td>44.7 ± 10.2% ($n = 3$)</td>
<td>58.4 ± 3.8% ($n = 6$)</td>
</tr>
</tbody>
</table>
BCR-mediated calcium mobilization. A decrease in calcium mobilization was seen in B lymphocytes from the CD19<sup>2/2</sup> mice in response to stimulation with an optimal dose of antibody (Fig. 11A). At suboptimal concentrations of antibody (1.3 μg F(ab')<sub>2</sub>RAMIG) almost no BCR-mediated calcium mobilization was seen in cells from CD19<sup>2/2</sup> mice, while cells from the CD19<sup>1/1</sup> littermates still responded well (Fig. 11B). We obtained equivalent results in 10 independent experiments. Our results appear to conflict with those of Sato et al. (34), who observed enhancement of later phase Ca<sup>2+</sup> mobilization in CD19<sup>2/2</sup> cells. These contrasting findings may result from technical differences in the respective studies.

Discussion

Recent studies of CD19<sup>2/2</sup> mice and CD19 transgenic mice have demonstrated the importance of CD19 in B cell development, but particularly in the immune response of mature B cells. In human CD19 transgenic mice B cell development is impaired (45) as indicated by the fact that these mice have dramatically fewer B cells in the periphery. In these transgenic animals proper B cell development is very sensitive to even small increases in CD19 expression levels (46). CD19<sup>2/2</sup> mice also have lower levels of B cells in the periphery, and mature B cells from these animals mount a poor proliferative response to anti-BCR antibodies. These animals respond only weakly to T cell-dependent antigens, germinal center formation is lacking or diminished, and the serum immunoglobulin levels are significantly reduced (26, 27, 47). Based on these findings, it has been suggested that CD19 functions by enhancing the cellular sensitivity to antigen, setting the threshold for antigen-mediated signaling (46). This may reflect a CR2-dependent coreceptor function and/or an accessory function of CD19 in BCR signal transduction.

Here we have defined an important accessory function for CD19 in antigen-induced BCR signal transduction and described a molecular basis for this accessory function. CD19 was found to influence the magnitude of activation...
of multiple intracellular signaling pathways after antigen stimulation of the BCR. While expression of CD19 leads to modestly enhanced antigen-induced Ig-α, Lyn, Syk, PLC-γ1, and PLC-γ2-inductive tyrosine phosphorylation, CD19 is absolutely required in order to obtain substantial PI3-kinase activation, IP3 generation, and maximal calcium mobilization. Thus, CD19 modifies BCR signaling in a qualitative manner. Using the PI3-kinase inhibitor wortmannin we further determined that while the increased antigen-induced tyrosine phosphorylation of Ig-α, Lyn, and Syk is independent of PI3-kinase activation, differences in IP3 turnover and calcium mobilization are a function of PI3-kinase activation. Results obtained in CD19-positive and -negative variants of the J558Lµm3CD45+ plasmacytoma cells were confirmed using primary B lymphocytes from CD19−/− mice and wild-type littermates. Thus CD19 functions in promoting BCR-mediated phosphoinositide hydrolysis via activation of PI3-kinase, and also supports antigen-induced phosphorylation of BCR receptor components and receptor activated kinases by an unknown mechanism. Since the src-family kinases Lyn and Lck are found associated with CD19 in unstimulated cells (5, 6), CD19 may mediate the latter effect via an increase in Ig-α phosphorylation by these kinases.

It has been demonstrated previously that wortmannin can inhibit angiotensin II-mediated IP3 production in adrenal glomerulosa cells (48) as well as anti-DNP-IgE-mediated IP3 production in BL-2H3 cells (49). The mechanisms by which PI3-kinase affects inositol lipid hydrolysis is unclear. PLC-γ1 and PLC-γ2 are highly homologous enzymes, both contain two SH2 domains and one SH3 domain. In vitro studies suggest that tyrosine phosphorylation of the enzymes is required to increase the catalytic activity, but phosphatase treatment of phosphorylated PLC-γ1 only partially inhibits PLC-γ1-mediated inositol trisphosphate production (50), indicating that mechanisms other than tyrosine phosphorylation may regulate PLC-γ1 activation. Our studies using the PI3-kinase inhibitor, wortmannin, indicate that antigen-stimulated inositol lipid turnover and calcium mobilization are dependent on PI3-kinase activation in B cells. Interestingly, induction of PLC-γ1 and PLC-γ2 tyrosine phosphorylation is relatively insensitive to wortmannin. As shown in Table II, IP3 production is substantially inhibited by 10 nM wortmannin, while this concentration of wortmannin has a negligible effect on PLC-γ1 and PLC-γ2 tyrosine phosphorylation. PI3-kinase phosphorylates the phosphoinositol containing lipids PI, PI(4)P and PI(4,5)P2 at the D3 position of the inositol ring (30). Two of the products of this phosphorylation, PI(3,4)P2 and PI(3,4,5)P3, are important regulators of cell proliferation (51). Recently it was suggested that PI(3,4,5)P3 may play a role in signaling by associating with SH2 domains, and thereby modulating PI3-kinase association with tyrosine phosphorylated proteins (52). PI(3,4,5)P3 could thereby regulate the activity of SH2 containing effector enzymes either directly or indirectly by relocating them. We suggest that this may reflect PI(3,4,5)P3, the product of PI3-kinase, association with the SH2 domain of PLC-γ1 and translocation of the enzyme to the plasma membrane where its substrate resides. Indeed addition of PI(3,4,5)P3 to substrate vesicles has been shown to increase PLC-γ hydrolysis of PI(4,5)P2 (53). It has been demonstrated in other cell types that cell stimulation results in translocation of PLC-γ1 and PLC-γ2 from the cytosol to the membrane (54, 55). This suggests that activation of phosphatidylinositol hydrolysis involves both a PLC-γ tyrosine phosphorylation event and a translocation event. Alternatively, or additionally, interactions between PI(3,4,5)P3 and pleckstrin homology (PH) domains could be involved in translocation of effector proteins to the membrane to obtain optimal phosphoinositide hydrolysis and Ca2+ mobilization. A likely candidate is Bruton’s tyrosine kinase, Btk, which has been shown to selectively interact with PI(3,4,5)P3 vesicles via its PH domain (56). Indeed recent findings suggest that Btk is required for BCR-mediated Ca2+ mobilization; in particular for extracellular influx (R awlings, D., personal communication). In this regard we do see a higher antigen-mediated inductive tyrosine phosphorylation of Btk in our CD19-positive plasmacytoma cell
line than in the CD19-negative line, suggesting that CD19 may influence the activation of Btk (results not shown). Recently it was shown in both T cells and B cells that under certain circumstances cell stimulation can lead to tyrosine phosphorylation of PLCγ, but no IP3 production or calcium mobilization (41, 57–59). These examples further underscores the concept that tyrosine phosphorylation of PLCγ is not in itself sufficient to induce maximal IP3 generation.

A critical role for CD19 phosphorylation in BCR activation of PI3-kinase, IP3 production, and calcium mobilization is also supported by studies of FcγRIIB1 signal transduction. We and others have recently shown that coligation of BCR with FcγRIIB1 leads to rapid dephosphorylation of CD19 and failed PI3-kinase activation (60, 61). Therefore, FcγRIIB1-mediated CD19 dephosphorylation appears to prevent the normal contribution of CD19 to BCR signaling. Importantly, FcγRIIB1 cocrusslinking with BCR also results in aborted IP3 production and reduced calcium mobilization. Thus, the calcium mobilization and IP3 production phenotypes of the CD19-positive and -negative cell lines are clearly reminiscent of the signaling phenotypes of B cells stimulated with F(ab′)2, or intact antibodies, respectively (62, 63). F(ab′)2 stimulation leads to a strong increase in both intracellular and extracellular [Ca2+] mobilization, comparable to the antigen-mediated calcium mobilization in the CD19-positive cells, and a strong and sustained IP3 production. In contrast coengagement of the FcγRIIB1 receptor by stimulation with intact antibodies only induces a transient production of IP3, and calcium mobilization as seen in the CD19-negative B cell line (Figs. 2 and 3). These data support a role for CD19 and its effector PI3-kinase in BCR-mediated PLCγ activation and calcium mobilization.

Since our studies indicate that CD19 is required for maximal (i.e., both primary and secondary phase) Ca2+ mobilization, modulation of CD19 tyrosine phosphorylation, and resulting modulation of [Ca2+] mobilization pattern may be important in differentially regulating downstream signaling pathways such as activation of MAP kinase pathways, transcription factors, and proliferation/apoptosis responses (64, 65). Recently it was demonstrated that the amplitude and duration of Ca2+ signals contribute to transcriptional specificity (64). Such effects could at least in part be a consequence of the contribution of CD19, i.e., stimulation of B lymphocytes with F(ab′)2RAMIG, in which case CD19 is highly tyrosine phosphorylated, leads to a Ca2+ spike and a constant elevated second phase Ca2+ response. In contrast binding of immune complexes to FcγRIIB1 and the BCR, in which case CD19 tyrosine phosphorylation is decreased, results in a Ca2+ spike, but no prolonged second phase Ca2+ response triggering an effector output different from the F(ab′)2RAMIG stimulation. These modulations of the Ca2+ response may result in modulation of cell proliferation and apoptosis. Splenic B cells proliferate in response to B cell mitogens such as Staphylococcus aureus (SA) and anti-IgM. The SA-proliferation response is partially inhibited by the PI3-kinase inhibitor wortmannin (66), consistent with a role of PI3-kinase in BCR-mediated proliferation responses. CD19 activation may also be important for the regulation of apoptosis. When immune complexes are bound to the FcγRIIB1 and the BCR, in which case CD19 tyrosine phosphorylation is decreased, B cell proliferation is inhibited and cells undergo apoptosis (67). We hypothesize that this inhibition of proliferation and induction of apoptosis is a consequence of inhibition of PI3-kinase activity associated with CD19, leading to inhibition of downstream signaling pathways. Potentially FcγRIIB1-mediated inhibition of CD19 tyrosine phosphorylation and PI3-kinase activation could result in a failure to activate the serine/threonine protein kinase Akt/PKB, which has been shown to be a target of PI3-kinase (68–70), and is known to protect cells against apoptosis (71–73).

Here we have demonstrated a potential molecular basis for the essential role played by CD19 in antigen-mediated B cell activation, germinal center formation and generation of immune responses. CD19 functions as a constitutive component of the BCR signaling apparatus and is required for antigen-mediated BCR stimulation of PI3-kinase activation and, via PI3-kinase activation, phosphoinositide hydrolysis, and calcium mobilization. In fact, this is the first formal demonstration that CD19 modifies BCR signaling directly. The findings have important biological implications since CD19 and BCR expression are differentially regulated during B cell development. Variation of the ratio of expression of these molecules may provide a mechanism to vary the output of the receptor as a function of cell differentiation stage. Of potential importance in this context is the fact that the contribution of CD19 to BCR signaling is qualitative; it does not simply change the magnitude of the signal.

We thank Michael Reth for provision of the J558Lμm3 cells, Thomas Tedder for the hCD19 cDNA, and Marc Hertz for help with use of the JMP statistical analysis program, comments and reading of the manuscript.

A.M. Buhl is a Leukemia Society Research Fellow. J.C. Cambier is an Ida and Cecil Green Professor of Cell Biology. These studies were supported by the US Public Health Service.

Address correspondence to Dr. John C. Cambier, Division of Basic Sciences, Department of Pediatrics, National Jewish Research and Medical Center, 1400 Jackson St., Denver, CO 80206. Tel.: (303) 398-1325; FAX: (303) 398-1225; E-mail: cambier@nje.org

Received for publication 10 July 1997 and in revised form 19 September 1997.
References


30. Whitman, M., C.P. Downes, M. Keeler, T. Keller, and L.


74. CalCalc—A Program to Calculate Calcium Concentrations in the Presence of EGTA. Department of Biochemistry, Colorado State University, Fort Collins, Colorado.