Scaffold Protein JLP Is Critical for CD40 Signaling in B Lymphocytes*

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Background: The roles of JLP in CD40 trafficking and signaling in B lymphocytes remain elusive.

Results: JLP deficiency in B lymphocytes resulted in selective diminished CD40 internalization and activation of MAPK and JNK.

Conclusion: JLP-mediated CD40 internalization is essential for downstream JNK and MAPK signaling in B lymphocytes.

Significance: We report a novel role of JLP in the bridging of CD40 internalization and CD40-dependent signaling in B lymphocytes.

CD40 expression on the surface of B lymphocytes is essential for their biological function and fate decision. The engagement of CD40 with its cognate ligand, CD154, leads to a sequence of cellular events in B lymphocytes, including CD40 cytoplasmic translocation, a temporal and spatial organization of effector molecules, and a cascade of CD40-induced signal transduction. The JLP scaffold protein was expressed in murine B lymphocytes. Using B lymphocytes from jlp-deficient mice, we observed that JLP deficiency resulted in defective CD40 internalization upon CD154/CD40 engagement. Examination of interactions and co-localization among CD40, JLP, dynein, and Rab5 in B lymphocytes suggested that CD40 internalization is a process of JLP-mediated vesicle transportation that depends on Rab5 and dynein. JLP deficiency also diminished CD40-dependent activation of MAPK and JNK, but not NF-κB. Inhibiting vesicle transportation from the direction of cell periphery to the cell center by a dynin inhibitor (ciliobrevin D) impaired both CD154-induced CD40 internalization and CD40-dependent MAPK activities in B lymphocytes. Collectively, our data demonstrate a novel role of the JLP scaffold protein in the bridging of CD154-triggered CD40 internalization and CD40-dependent signaling in splenic B lymphocytes.

CD40 is a member of the TNF receptor superfamily. It is expressed in various immune cells, such as B lymphocytes, dendritic cells, and macrophages. CD154 (a ligand of CD40) is transiently expressed on the surface of activated CD4+ T cells. The CD40-CD154 interaction results in the binding of TNF receptor-associated factors (TRAFs) to the CD40 cytoplasmic tail and subsequent activation of MAPK cascades. MAPK signaling triggered by CD40 engagement is pivotal for shaping B lymphocyte function and fate, such as immunoglobulin class switching, memory B lymphocyte development, affinity maturation and germinal center formation, cytokine production, and plasma cell survival (1).

In MAPK signal transduction, scaffold proteins are needed to coordinate the signaling network by organizing discrete components in space and time. JNK-interacting proteins (JIPs) are defined as scaffold proteins in the JNK and p38 signaling pathways. Several members of the JIP family have been discovered: JIP1–4 and JNK-associated leucine zipper protein (JLP), which assemble various components in the JNK and p38 signaling pathways (2). Aside from roles in the control of signaling flow, the JLP scaffold protein is also involved in intracellular molecular trafficking or vesicle transportation (3). JLP has been documented, along with other JIPs, to be important in vesicle or protein transport via its association with the motor protein kinesin or dynein, which is generally required for anterograde or retrograde transport of cargo between the cell center and periphery along microtubules (4, 5).

Upon receptor-ligand engagement, receptor internalization is a common mechanism through which the surface receptor is transported to the cytoplasm to down-regulate receptor density on the cell surface and to initiate downstream signal transduc-

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4 The abbreviations used are: TRAF, TNF receptor-associated factor; JIP, JNK-interacting protein; JLP, JNK-associated leucine zipper protein; rCD154, recombinant CD154; DIC, dynein intermediate chain; MFI, mean fluorescence index.
tion. Ligand engagement leads to the formation of CD40 lipid rafts on the cell membrane, followed by CD40 internalization and downstream signaling (6–10). Using a gene-silencing approach, we reported previously that the JLP scaffold protein regulated cytoplasmic trafficking of CD40 receptors in dendritic cells (11). However, whether JLP plays a role in regulating CD154-initiated CD40 receptor internalization and downstream signaling in B lymphocytes is unknown. Here, we examined, for the first time, the role of JLP and its underlying mechanism in the regulation of CD154-induced CD40 internalization and MAPK signaling in B lymphocytes.

**EXPERIMENTAL PROCEDURES**

**Animals**—Wild-type (Jlp<sup>+/+</sup>) and homozygous (Jlp<sup>−/−</sup>) mice were generated by inbreeding Jlp<sup>+/+</sup> mice, and they were genotyped by RT-PCR and Western blotting as described previously (12). 8–10-week-old weight- and sex-matched Jlp<sup>+/+</sup> and Jlp<sup>−/−</sup> littermates were used in these experiments. Mice were housed under specific pathogen-free conditions at the Center for Animal Experiments at Wuhan University. All animal experiments were approved by the Animal Ethics Review Board of Wuhan University and performed in accordance with the guidelines of the National Health and Medical Research Council of China.

**Reagents**—Recombinant mouse CD40 ligand (rCD154) was from R&D Systems (Minneapolis, MN). FITC-conjugated anti-CD19, phycoerythrin-conjugated anti-CD80, allophycocyanin-conjugated anti-CD40, allophycocyanin-conjugated anti-CD86, allophycocyanin-conjugated anti-MHC-II, and rat anti-CD40 (1C10) antibodies and 7-aminomycin D were purchased from BioLegend (San Diego, CA). LPS (Escherichia coli O111:B4) and anisomycin were from Sigma-Aldrich. The dynein inhibitor ciliobrevin D and mouse anti-DIC antibody were from Merck Millipore (Billerica, MA). Anti-ERK, anti-JNK, anti-p38, anti-phospho-ERK (Thr-202/Tyr-204), anti-phospho-JNK (Thr-183/Tyr-185), anti-phospho-p38 (Thr-180/Tyr-182), anti-phospho-IL-6 receptor (Ser-32), and anti-phospho-c-Jun (Ser-73) antibodies were purchased from Cell Signaling Technology (Danvers, MA). Rabbit anti-JLP, rabbit anti-CD40, and rabbit anti-dynein intermediate chain (DIC) were purchased from Abcam (Cambridge, MA).

**Cell Culture**—B lymphocytes (CD19<sup>+CD5</sup><sup>−</sup>) were purified from splenocytes with BD IMag mouse B lymphocyte enrichment set-DM and BD IMag cell separation magnet (BD Biosciences) for negative selection. The purity of the BD IMag-sorted cell populations was ~88–95%. Sorted B lymphocytes were cultured in complete medium (RPMI 1640 medium supplemented with 10% FCS (Invitrogen) and 1% penicillin/streptomycin) for negative selection. HEK293 cells were maintained in complete medium with a <0.1% Tween 20 and 5% skim milk powder for 1 h at room temperature and cultured with the indicated primary antibodies at 4 °C overnight. The next day, membranes were prewashed three times for 5 min with TBST before incubation with IRDye 800CW secondary antibodies (LI-COR Biotechnology, Lincoln, NE) for 1 h at room temperature. After washing three times for 5 min with TBST, membranes were scanned using an Odyssey CLx infrared imaging system (LI-COR Biotechnology), and data were acquired using Odyssey application software. Some protein bands were quantified with Quantity One analysis software (Bio-Rad).

**Immunoprecipitation**—Cells were lysed, and cell lysates were precleared by adding an appropriate control IgG and Protein G PLUS/Protein A-agarose (Merck Millipore). Precleared lysates were immunoprecipitated by overnight incubation with primary antibodies and subsequent incubation with Protein G PLUS/Protein A-agarose for 2 h. Samples were then separated by SDS-PAGE and analyzed by Western blotting as described above. An isotype-specific IgG-agarose was applied as a negative control in these experiments.

**Confocal Microscopy**—Purified splenic B lymphocytes were harvested, washed three times with ice-cold PBS, and fixed with 2% paraformaldehyde for 20 min at room temperature. After blocking with blocking buffer (PBS containing 10% donkey serum), samples were incubated with primary antibodies (from rabbit, mouse, or rat) at 4 °C overnight. Alexa Fluor 488- or Alexa Fluor 594-conjugated donkey anti-rabbit, anti-mouse, and anti-rat IgG antibodies (Jackson ImmunoResearch Laboratories, West Grove, PA) were added and incubated for 1 h. After three washings with PBS, cells were visualized using a PerkinElmer UltraVIEW VoX spinning disk confocal microscope system with a ×100 oil immersion objective lens.

**Statistical Analysis**—SPSS software was used to analyze data, and a two-tailed Student’s t test was applied to determine dif-
JLP Deficiency Has No Effect on Splenic Lymphocyte Numbers and B Cell Phenotypes—Previous studies suggest that the JLP protein is expressed at moderate levels in the brain, lung, spleen, and ovary and at low levels in the heart, liver, kidney, epididymis, and uterus (12). Thus, JLP deficiency may cause abnormalities in the spleen. To test this, we first examined the spleen volume of both $jlp^{-/-}$ and $jlp^{+/+}$ mice and observed no significant differences in spleen size and absolute numbers of splenic T and B lymphocytes (Fig. 1, A and B). JLP proteins are expressed in some immortal cell lines and primary dendritic cells (13–15), but its expression in primary B lymphocytes has not yet been reported. Here, we found that JLP proteins are expressed in B lymphocytes (Fig. 1C). To examine whether JLP deficiency affects development of fully differentiated B lymphocytes and LPS-induced activation, we compared the surface expression of CD19, CD40, CD80, CD86, and MHC-II in resting splenic B lymphocytes from $jlp^{-/-}$ and $jlp^{+/+}$ mice and found the markers to be comparable. Upon LPS stimulation, we observed similar up-regulation of the surface expression of CD19, CD40, CD80, CD86, and MHC-II in the activated $jlp^{+/+}$ and $jlp^{-/-}$ B lymphocytes (Fig. 1D). Of note, no significant difference in surface CD40 expression was observed between groups.

RESULTS

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FIGURE 1. JLP deficiency has no effect on splenic lymphocyte numbers and B cell phenotypes. A, $jlp^{-/-}$ and $jlp^{+/+}$ mice had similar spleen volumes. Spleens were obtained from 8-week-old mice, and one representative result is shown. B, analysis of absolute numbers of splenic B lymphocytes (CD19$^+$) and T cells (CD3$^+$) from four pairs of age- and weight-matched $jlp^{+/+}$ and $jlp^{-/-}$ mice. Horizontal bars are mean values. C, expression of JLP in splenic B lymphocytes from $jlp^{-/-}$ and $jlp^{+/+}$ mice. Purified splenic B lymphocyte lysates were subjected to SDS-PAGE and analyzed by Western blotting. HeLa cell extract (Hela CE) and actin were used as positive and loading controls, respectively. D, phenotype analysis of splenic B lymphocytes from two mouse genotypes with or without LPS (1 µg/ml) stimulation for 24 h. B lymphocytes were stained with antibodies specific to CD19, CD40, CD80, CD86, and MHC-II. Cells were counted with a flow cytometer, and the histograms represent surface expression of the indicated markers on gated CD19$^+$ B lymphocytes from $jlp^{-/-}$ (solid black lines) and $jlp^{+/+}$ (dashed gray lines) mice. The filled curves are negative controls. The results from statistical analysis of the mean fluorescence index (MFI) of each marker from three independent experiments are shown. Error bars indicate S.D.
observed in the LPS-matured jlp<sup>++/+</sup> and jlp<sup>−/−</sup> B lymphocytes (Fig. 1D).

**Scaffold Protein JLP Is Required for CD154-induced CD40 Internalization—**Engagement of the surface CD40 receptor by its cognate ligand CD154 induces CD40 internalization in B lymphocytes and vascular endothelial cells (10, 16–19). The molecular mechanism underlying CD154-induced CD40 internalization has not yet been fully investigated. Using a gene-silencing approach, we demonstrated previously that LPS-stimulated JLP-knockdown dendritic cells were impaired in regulating intracellular CD40 trafficking upon CD40 activation (11). The availability of the JLP-deficient mice allowed us to further examine whether JLP regulation of CD40 internalization is a common mechanism observed in B lymphocytes. As freshly isolated splenic B lymphocytes express low levels of surface CD40 molecules (Fig. 1D), LPS (1 µg/ml) was used to up-regulate surface CD40 expression on B lymphocytes in all of the CD154 stimulation experiments described in the work. We confirmed that addition of CD154 did not interfere with the binding of anti-CD40 antibody to B lymphocytes (data not shown). Upon stimulation of rCD154, surface CD40 expression on the wild-type (jlp<sup>++/+</sup>) B lymphocytes was rapidly down-regulated within 15 min and recovered at 30 min. In contrast, the surface CD40 down-regulation induced by rCD154 was mitigated in the jlp<sup>−/−</sup> B lymphocytes (Fig. 2A).

We confirmed this finding further using a jlp gene-silencing approach and CD40L-HEK293 transfectants (which express membrane-bound CD154, a physiologic form of CD154 in CD154 stimulations) in the activation of surface CD40 on B lymphocytes. JLP-specific shRNA was delivered into a mature B cell line (WEHI-279) to knock down endogenous expression of JLP. Impaired CD40 internalization was observed in the JLP-knockdown WEHI-279 B cells upon CD154 engagement (Fig. 2C). To examine the role of JLP in the membrane-bound CD154-induced CD40 internalization in B lymphocytes, purified jlp<sup>++/+</sup> and jlp<sup>−/−</sup> B cells were co-cultured with mock-transfected HEK293 cells or with HEK293 cells transfected with a CD154-expressing vector. We observed strong surface CD40 down-regulation in jlp<sup>++/+</sup> B cells after co-culture with the CD154-transfected HEK293 cells. In contrast, only a slight decrease in surface CD40 was detected in jlp<sup>−/−</sup> B cells (Fig. 2B). Down-regulation of surface CD40 was a result of CD40 internalization because total CD40 expression (both surface and intracellular, determined by flow cytometry analysis and Western blotting) remained stable during CD40 ligation in both jlp<sup>++/+</sup> and jlp<sup>−/−</sup> mice (Fig. 2, D and E).

**JLP Interacts with CD40 during CD40 Internalization—**We next examined whether JLP proteins associate with CD40 in the internalization process. Using anti-CD40 and anti-JLP antibodies in co-immunoprecipitation experiments, we observed that there was a basal level of JLP and CD40 association before rCD154 stimulation. Upon addition of rCD154, the association of JLP and CD40 was increased by 2.2-fold at 5 min and by 2.7-fold at 15 min as determined by immunoprecipitation of JLP (Fig. 3A, left panel) or by 4.2-fold at 5 min and by 4.3-fold at 15 min as determined by immunoprecipitation of CD40 (Fig. 3A, right panel). Confocal microscopy also confirmed the co-localization of CD40 with JLP in the intracellular vesicles after B lymphocytes were stimulated with rCD154 for 15 min (Fig. 3B). These results demonstrate that CD40-CD154 engagement promotes the formation of CD40-JLP complexes in stimulated B lymphocytes.

**CD40 Internalization Depends on Both JLP and Dynein—**Vesicle transportation is involved in various cell biological activities, including receptor internalization and endocytosis. Motor proteins are needed to drive vesicle complexes to move along cellular microtubules. Kinesin and dynein are two major superfamilies of motor proteins identified to date and are required for anterograde or retrograde transport of cargos between the cell center and periphery along these microtubules (20). Dynein transports cargos from the cell periphery to the cell center with the help of an accessory complex, dynactin (5, 21). To investigate whether dynein is required for CD40 internalization in B lymphocytes, we studied the effects of ciliobrevin D, a recently described dynein-specific inhibitor (22–24), on the CD40 internalization induced by CD154 engagement. A previous study reported that dynein inhibition in B lymphocytes resulted in severe impairment of the net movement of antigen (anti-IgM) toward the center (25). We observed a similar effect in the ciliobrevin D-treated B lymphocytes. As shown in Fig. 4A, anti-IgM monoclonal antibody stimulation induced significant antigen intracellular gathering in a dimethyl sulfoxide control but not in the presence of ciliobrevin D, thus confirming the efficiency of ciliobrevin D in dynein inhibition. No cytotoxicity effects was observed in jlp<sup>++/+</sup> and jlp<sup>−/−</sup> cells upon treatment with ciliobrevin D at 50 µM for 60 min (Fig. 4B). Surface CD40 down-regulation caused by rCD154 engagement was largely decreased in the ciliobrevin D-treated jlp<sup>−/−</sup> splenic B lymphocytes. In contrast, ciliobrevin D pretreatment did not affect surface CD40 expression upon rCD154 stimulation in the jlp<sup>−/−</sup> splenic B lymphocytes (Fig. 4C). These data demonstrate that dynein is involved in CD40 internalization by CD154 ligation in B lymphocytes.

**Association among CD40, Rab5, JLP, and Dynein during CD40 Internalization—**Rab proteins are important in recruiting specific motors to organelar membranes or adaptor/scaffold proteins. Among these Rab molecules, Rab5 is reported to be responsible for receptor transportation from the plasma membrane to early endosomes (26, 27). To further identify the molecular mechanism of CD40 internalization in B lymphocytes, we investigated interactions and co-localizations among CD40, Rab5, JLP, and dynein in jlp<sup>++/+</sup> and jlp<sup>−/−</sup> B lymphocytes.

We first examined the localization of CD40 and Rab5 in B lymphocytes during CD154-induced CD40 internalization. Under normal conditions, no significant co-localization of CD40 and Rab5 can be observed because CD40 and Rab5 are located in different cellular compartments. However, 15 min after CD40-CD154 engagement, substantial co-localization of CD40 and Rab5 in the same subcellular compartment was observed in the cytoplasm of jlp<sup>−/−</sup> B lymphocytes. In contrast, CD40 ligation in jlp<sup>−/−</sup> B lymphocytes did not lead to detectable CD40 and Rab5 co-localization (Fig. 5A).

Immunoprecipitation of proteins from lysates of jlp<sup>++/+</sup> B lymphocytes revealed a constitutive association between CD40

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and DIC without CD40 engagement. This association was strongly enhanced at 15 min after rCD154 addition. In contrast, a weak association between CD40 and DIC was detected in jlp<sup>−/−</sup> B lymphocytes despite CD40 ligation (Fig. 5B). Confocal microscopy confirmed a JLP-dependent CD40-DIC association because co-localization between CD40 and DIC could be initiated by rCD154 only in jlp<sup>+/+</sup> B lymphocytes, but not in jlp<sup>−/−</sup> B lymphocytes (Fig. 5C). Also, we confirmed an association between dynein and JLP in B lymphocytes by immunoprecipitation and confocal microscopy, and this association was not altered by CD40 ligation (Fig. 5, D and E). We found that upon CD154 engagement, internalized CD40 was recruited together with JLP, dynein, and Rab5 in the same subcellular compartment.
Effects of JLP Deficiency on the CD40-initiated Signaling Pathway in B Lymphocytes—In B lymphocytes, CD40 engagement leads to activation of the ERK, JNK, p38, and NF-κB signaling pathways (28–30), which are coordinated by several scaffold proteins (31, 32). JLP has been reported to potentially interact with diverse effector components involved in JNK and p38 MAPK signaling (2). We analyzed the effects of JLP deficiency on these signaling pathways in B lymphocytes. We found that CD154-induced phosphorylation of JNK and c-Jun was impaired in jlp−/− B lymphocytes compared with jlp+/+ B lymphocytes. There was slightly less CD154-induced ERK activation and moderately less p38 MAPK activation in JLP-defective B lymphocytes compared with jlp+/+ B lymphocytes. However, no substantial difference in phosphorylated IκBα was observed between the two genotypic B lymphocytes (Fig. 6A).

Because JLP functions as a scaffold protein that may coordinate interactions between MAPK modules, weakened JNK and p38 MAPK signaling in jlp-deficient B lymphocytes may not be

FIGURE 3. JLP interacts with CD40 in B lymphocytes. A, cell extracts of splenic B lymphocytes from wild-type mice treated with rCD154 at the indicated time points were first incubated overnight with specific primary antibodies and then analyzed with the indicated antibody by co-immunoprecipitation. Input was 10% of total cell extracts, and immunoglobulin G was used as a negative control. Data are representative of three independent experiments. The indicated band intensity of immunoprecipitation (IP) was normalized to the relevant band intensity of the input. Means ± S.D. were calculated from three independent experiments. Error bars indicate S.D. *, p < 0.05; **, p < 0.01. B, fluorescence microscopy of B lymphocytes from wild-type mice. Cells were untreated (0 min) or were treated with rCD154 for 15 min and then labeled with the indicated primary antibody followed by a fluorescence-conjugated secondary antibody. The white arrow indicates areas of overlap. Scale bar = 5 μm. Experiments were performed in triplicate. The co-localization correlation coefficients of CD40 and JLP were collected from three separate experiments and statistically analyzed using UltraVIEW software. *, p < 0.05 (Student’s t test).

FIGURE 2. CD40 internalization after CD154 ligation is mediated by the scaffold protein JLP. A, CD40 expression on splenic B lymphocytes with rCD154 stimulation for different times. B lymphocytes from jlp+/+ or jlp−/− mice pre-activated with LPS (1 μg/ml) were stimulated with rCD154 (1 μg/ml) for the indicated times and subsequently subjected to CD19 and CD40 staining. CD40 expression on gated CD19− B lymphocytes is shown in the histograms. The percentage MFI reduction of CD40 was calculated by dividing the CD40 MFI at the indicated time points by that at 0 min. B, CD40 expression on splenic B lymphocytes co-cultured with CD154-transfected HEK293 cells. Splenic B lymphocytes from jlp+/+ or jlp−/− mice were co-cultured with mock- or CD154-transfected HEK293 cells (HEK293/B cell ratio of 1:50) for 15 min. Cells were harvested and subjected to flow cytometry analysis of CD40 expression. C, CD40 expression on siRNA-transfected WEHI-279 B cells upon rCD154 stimulation. JLP knockdown in WEHI-279 cells with siRNA transfection was confirmed by immunoblot analysis. The percentage MFI reduction of CD40 was calculated by dividing the CD40 MFI after CD154 treatment by that at baseline. Ctrl, control. Total CD40 expression in splenic B lymphocytes at the indicated time points upon soluble CD154 treatment was examined by flow cytometry (D) and immunoblotting (E). For flow cytometry, cells were permeabilized for CD40 staining. The percentage MFI reduction of CD40 expression at the indicated time points is compared with that at 0 min. For immunoblotting, whole-cell lysates were prepared as described above. In the flow cytometry analysis, data from three different experiments were collected and statistically analyzed. In the immunoblot analysis, the indicated band intensity of JLP was normalized by the relevant band intensity of β-actin. Means ± S.D. were collected from at least three independent experiments. Error bars indicate S.D. *, p < 0.05 (Student’s t test).
CD40-specific. To confirm its specific role in CD40 signaling, we cultured B lymphocytes in the presence of anisomycin, a specific agonist targeting the p38 and JNK signaling pathways. We observed that phosphorylation of JNK and p38 in anisomycin-stimulated \( jlp^-/- \) B lymphocytes was comparable with that in the \( jlp^+/+ \) B lymphocytes (Fig. 6B). Furthermore, blocking CD154-induced CD40 cytoplasmic movement by treatment with ciliobrevin D prevented CD154-induced JNK and p38 activation in B lymphocytes (Fig. 6C). There was no difference in the activation of JNK and p38 signaling in the anisomycin-stimulated B lymphocytes pretreated with or without ciliobrevin D (Fig. 6D), indicating a specific upstream signaling flow of MAPK activation triggered by CD40 engagement and anisomycin may exist.

**DISCUSSION**

The engagement of CD40 by its ligand (CD154) leads to CD40 clustering in membrane rafts (7, 8), TRAF recruitment to lipid rafts (16, 17), CD40 cytoplasmic translocation (16, 17), and subsequent signal initiation and transduction (33) in B lymphocytes. Upon ligation of CD40-CD154, effector molecules are finely organized both temporally and spatially via scaffold proteins. However, to our knowledge, the details of scaffold protein-mediated CD40 trafficking and signaling are still poorly understood. Here, we investigated the role of the scaffold protein JLP in CD154-triggered CD40 internalization and subsequent CD40-dependent signaling pathways, including MAPK and NF-κB, in primary murine B lymphocytes. We observed that JLP deficiency causes defective CD40 internalization upon CD154 engagement. Through analysis of interactions and colocalizations among CD40, JLP, dynein, and Rab5 in B lymphocytes, we found that CD40 internalization is a form of JLP-mediated vesicle transportation that depends on Rab5 and dynein. JLP deficiency diminishes CD40-dependent activation of the MAPK (but not NF-κB) signaling pathway, and JNK activity is the most severely affected. The effect of JLP deficiency in signaling pathways is CD40-specific because the p38 and JNK pathways can be activated by a p38- and JNK-specific agonist (anisomycin) in \( jlp^- \)-deficient B lymphocytes.

Intracellular synthesized CD40 proteins are separated from the Golgi complex and transported to the cell membrane as a surface receptor. There, surface CD40 can translocate to the cytoplasm upon engagement with its ligand and initiate various cellular responses. We observed that CD40 endocytosis is halted more obviously than its recycling to the surface in the \( jlp^- \)-deficient B lymphocytes. However, we cannot rule out the possibility that the impaired CD40 recycling in the JLP-deficient cells is due to less surface CD40 being internalized into the cytoplasm. We also demonstrated that JLP deficiency cannot completely abolish CD40 internalization, indicating that other JIP molecules may be also involved in CD40 internalization (34–38). In fact, JIP1 and JIP3 have been confirmed to bind dynactin to regulate cargo or organellar retrograde trafficking...
FIGURE 5. JLP deficiency impairs association among CD40, Rab5, and dynein during CD40 engagement. A, confocal microscopy was used to visualize co-localization of CD40 and Rab5 in splenic B lymphocytes from *jp**/** and *jp**/** mice. B lymphocytes were left untreated (0 min) or were treated with rCD154 (1 µg/ml) for 15 min. Cells were then stained with antibody specific to the appropriate molecules. The co-localization correlation coefficients of CD40 and Rab5 from *jp**/** (black bars) and *jp**/** (white bars) B lymphocytes were collected from three separate experiments and statistically analyzed using UltraVIEW software. **, *p < 0.01 (Student’s t test). CD40 immunoprecipitated with anti-dynein antibody in *jp**/** and *jp**/** mouse splenic B lymphocytes (B), and JLP immunoprecipitated with anti-dynein antibody in cell extracts of splenic B lymphocytes from wild-type mice (D). Cells were treated as described above and then immunoprecipitated. Input was 10% of total cell extracts. IgG served as a negative control. The indicated band intensity of immunoprecipitation (IP) was normalized to the relevant band intensity of the input. Means ± S.D. were calculated from three independent experiments. Error bars indicate S.D. **, *p < 0.05; **, *p < 0.01 (Student’s t test). IB, immunoblot. The co-localization of CD40 (red) and dynein (green) in *jp**/** and *jp**/** splenic B lymphocytes (C) or JLP (red) and dynein (green) in wild-type splenic B lymphocytes (E) was visualized using a confocal microscope. B lymphocytes were left untreated (0 min) or were treated with rCD154 (1 µg/ml) for 15 min. Cells were then stained with a primary antibody, followed by a fluorescence-conjugated secondary antibody. The white arrows indicate areas of overlap. Scale bar = 5 µm. Experiments were performed three times, and similar images were obtained. The co-localization correlation coefficients of the indicated molecules were collected from three separate experiments and statistically analyzed. **, *p < 0.01 (Student’s t test).
We speculate that CD40 cytoplasmic translocation in B lymphocytes occurs through multiple JIP-dependent mechanisms. Several signaling pathways can be activated in response to CD40 engagement in B lymphocytes (33). Activation of the Akt (but not MAPK or NF-κB) signaling pathway has been documented to be dependent on CD40-lipid raft association, which is independent of any signaling events (42). The formation of a signaling complex containing CD40, TRAFs, and other components is required but not sufficient in CD40-dependent MAPK.

FIGURE 6. JLP deficiency selectively attenuates CD154-triggered activation of the MAPK (but not NF-κB) signaling pathway in B lymphocytes. A, purified splenic B lymphocytes from jlp+/− and jlp−/− mice were left untreated (0 min) or were treated with rCD154 (1 μg/ml) at the indicated time points. Cells were harvested, and whole-cell lysates were electrophoresed and analyzed for total ERK, p38, JNK, and relevant phosphorylated (p) forms of ERK, p38, JNK, c-Jun, and IκBα by immunoblotting. Equal protein loading was controlled by actin staining. B, purified splenic B lymphocytes from jlp+/− and jlp−/− mice were stimulated with anisomycin (250 ng/ml) at the indicated time points. Whole-cell extracts were analyzed by immunoblotting as described above. Wild-type B lymphocytes were pretreated with or without ciliobrevin D (50 μM) for 30 min and left unstimulated (0 min) or stimulated with rCD154 (C) or anisomycin (D) at 15 min and 30 min. Cells were then harvested, and the extracts were analyzed by immunoblotting as described above. Expression of the phosphorylated molecules was normalized to the indicated total molecule expression or β-actin. Means ± S.D. were collected from at least three independent experiments. Error bars indicate S.D. *, p < 0.05 (two-tailed Student’s t test).
and NF-κB activation, emphasizing the importance of cytoplasmic translocation of the signaling complex (43). This two-stage signaling mechanism can explain the spatial and temporal separation of MAPKs and NF-κB, but the molecular details regarding how these two stages are linked is unclear. In this study, we elucidated an essential role of the scaffold protein JLP in bridging signaling complex translocation and downstream signal transduction. Previous studies show that TRAF2/3 is critical for CD40-induced JNK activation, but p38 MAPK and NF-κB are activated via multiple TRAF-dependent processes (17, 44). Also, TRAF2 or TRAF3 may have a structural function in mediating CD40 receptor internalization (17). In this study, we found that CD40-induced activation of ERK, p38, and JNK, but not NF-κB, is impaired in JLP-deficient B cells, which is consistent with previous findings showing that CD40 triggers different signal pathways via distinct molecular mechanisms. In addition, we have provided further details regarding JLP-mediated CD40 trafficking and JNK activation. With JLP deficiency, CD154-triggered CD40 internalization is severely limited, and CD40-induced activation of ERK, p38, and JNK, but not NF-κB, is impaired to various extents. Thus, different scaffold proteins are responsible for tethering CD40, TRAFs, and relevant downstream signaling kinases together to trigger different signaling events. Among these scaffold proteins, JLP may be critical for trafficking of a signaling complex containing CD40 and TRAF2/3. Whether JLP associates with TRAF2/3 directly or indirectly and whether other JIPs interact with other TRAFs to facilitate CD40-triggered signaling responses coordinated in B lymphocytes need to be examined further in future investigations.

In conclusion, we analyzed the role of the JLP scaffold protein in CD40 receptor internalization and subsequent signaling pathway activation in B lymphocytes. We described, for the first time, a putative molecular mechanism of CD40 receptor translocation and CD40-dependent signal transduction coordinated by the JLP scaffold protein. This study highlights further the exciting roles of scaffold proteins in the immune regulation of B lymphocytes.

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