DOCK2 and phosphoinositide-3 kinase δ mediate two complementary signaling pathways for CXCR5-dependent B cell migration

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Naive B cells use the chemokine receptor CXCR5 to enter B cell follicles, where they scan CXCL13-expressing ICAM-1⁺ VCAM-1⁺ follicular dendritic cells (FDCs) for the presence of antigen. CXCL13-CXCR5-mediated motility is mainly driven by the Rac guanine exchange factor DOCK2, which contains a binding domain for phosphoinositide-3,4,5-triphosphate (PIP3) and other phospholipids. While p110δ, the catalytic subunit of the class IA phosphoinositide-3-kinase (PI3K) δ, contributes to CXCR5-mediated B cell migration, the precise interdependency of DOCK2, p110δ, or other PI3K family members during this process remains incompletely understood. Here, we combined in vitro chemotaxis assays and in vivo imaging to examine the contribution of these two factors during murine naïve B cell migration to CXCL13. Our data confirm that p110δ is the main catalytic subunit mediating PI3K-dependent migration downstream CXCR5, whereas it does not contribute to chemotaxis triggered by CXCR4 or CCR7, two other chemokine receptors expressed on naive B cells. The contribution of p110δ activity to CXCR5-driven migration was complementary to that of DOCK2, and pharmacological or genetic interference with both pathways completely abrogated B cell chemotaxis to CXCL13. Our data confirm that p110δ is the main catalytic subunit mediating PI3K-dependent migration downstream CXCR5, whereas it does not contribute to chemotaxis triggered by CXCR4 or CCR7, two other chemokine receptors expressed on naive B cells. The contribution of p110δ activity to CXCR5-driven migration was complementary to that of DOCK2, and pharmacological or genetic interference with both pathways completely abrogated B cell chemotaxis to CXCL13. Intravital microscopy of control and gene-deficient B cells migrating on FDCs confirmed that lack of DOCK2 caused a profound migration defect, whereas p110δ contributed to cell speed and directionality. B cells lacking active p110δ also displayed defective adhesion to ICAM-1; yet, their migration impairment was maintained on ICAM-1-deficient FDCs. In sum, our data uncover two complementary signaling pathways mediated by DOCK2 and p110δ, which enable CXCR5-driven naive B cell examination of FDCs.

**KEYWORDS**

B cell migration, CXCR5 (C-X-C motif chemokine receptor 5), intravitral 2-photon microscopy, phosphoinoside-3-kinase, DOCK2
Introduction

Naïve follicular B cells are highly motile cells, which scan ICAM-1+ VCAM-1+ follicular dendritic cells (FDCs) for the presence of microbial antigen and the initiation of humoral responses. The chemokine receptor CXCR5 is critical for naïve B cell access to follicles, where FDCs, together with other stromal cells such as marginal reticular cells, produce its only ligand CXCL13 (1, 2). Furthermore, CXCR5 promotes together with the ICAM-1 receptor LFA-1 dynamic B cell surveillance of FDCs (3, 4). The lymphocyte-expressed guanine exchange factor (GEF) DOCK2 is a key signaling molecule for Rac activation and F-actin polymerization downstream of chemokine receptors in lymphocytes. In the absence of DOCK2, in vitro T and B cell migration towards homeostatic chemokines is strongly compromised, although residual migration persists (5). Accordingly, direct observation of peripheral lymph nodes (PLN) using intravital twophoton microscopy (2PM) uncovered that follicular accumulation and interstitial motility are substantially reduced but not completely abolished in DOCK2−/− deficient B cells (6).

DOCK family proteins contain two DOCK homology regions (DHR), of which DH1R is involved in phospholipid binding for membrane localization and DH2R mediates the GEF activity (7–9). The DH1R domain of DOCK2 binds the phosphoinositide-3-kinase (PI3K) product phosphoinositide-3,4,5-triphosphate (PIP3) as well as phosphatidic acid (PA). In B cells, the relation between DOCK2 and PI3K activity remains unclear to date. Whereas DOCK2 activity is not required for PI3K activation (5) and PI3K inhibition does not affect DOCK2-mediated migration in T cells (10), neutrophil-expressed DOCK2 regulates migration through PIP3-dependent membrane translocation and Rac activation (11). Along the same line, the class IA p110δ catalytic subunit is involved in B cell chemotaxis towards CXCL13 not but CCL19, CCL21 and CXCL12 (12), and regulatory subunits of class IA are required for basal B cell motility in vitro (13). A potential participation of class I catalytic subunits besides p110δ during CXCR5-mediated B cell chemotaxis has not been examined yet, despite evidence for activation of additional class I PI3K family members downstream of G-protein coupled receptors (14).

Here, we examined the migratory behavior of B cells carrying mutations in DOCK2 and the catalytic site of p110δ (p110δD910A/D910A), in combination with PI3K-specific pharmacological inhibitors, to dissect their contribution for CXCL13-elicited motility. Among class I PI3K catalytic subunits, we confirm a key contribution of p110δ to CXCR5-mediated migration, but not CXCR4 and CCR7-dependent migration. DOCK2 and p110δ activity comprised two complementary pathways for CXCR5-triggered B cell migration, and inhibition of both factors completely abolished chemotaxis. We corroborated our in vitro findings using intravital imaging of interstitial B cell scanning of FDCs. Finally, we found that while LFA-1 activity is reduced in the absence of catalytically active p110δ, the interstitial migration defect of p110δD910A/D910A B cells is maintained on ICAM-1-deficient FDCs. In sum, our study sheds light on intracellular signaling pathways governing CXCR5-driven follicular B cell motility, a prerequisite for the unfolding of humoral immune responses.

Results

p110δ is the dominant class I PI3K mediating B cell chemotaxis to CXCL13

The class I PI3K family member p110δ contributes to directed B cell migration towards CXCL13 (12). Using Transwell assays, we confirmed a role for the catalytic activity of p110δ for in vitro chemotaxis of primary murine B cells towards CXCL13, which was particularly evident at lower chemokine concentrations (reduction of 48% at 100 nM and 33% at 250 nM CXCL13 for p110δD910A/D910A B cells as compared to WT B cells, respectively; Figure 1A). To address whether additional catalytic subunits might contribute to WT and p110δD910A/D910A B cell migration, we performed chemotaxis assays in presence of the p110δ inhibitor TGX221, the p110γ inhibitor AS604850, the p110α/β/δ/γ inhibitor PI-103 and, as control, the p110δ inhibitor IC-87114. These data uncovered a decrease of WT B cell chemotaxis towards 100 nM CXCL13 only with PI-103 and IC-87114 (52% and 54% inhibition, respectively), while none of the inhibitors had a significant effect on p110δD910A/D910A B cell migration (Figure 1B). These findings suggest that other class IA and IB subunits do not substantially contribute to primary B cell migration towards CXCL13. The promigratory signaling function of p110δ was restricted to CXCR5, since B cell migration to CCR7 and CXCR4 ligands remained unchanged by genetic or pharmacological inhibition of its activity (Figures 1C, D), as reported (12). Similarly, CCR7-mediated primary T cell chemotaxis was not reduced by genetic or pharmacological inhibition of the catalytic activity of p110δ (Figure 1E).

DOCK2 and p110δ comprise two complementary pathways for CXCR5-mediated B cell migration

We next examined the potential relationship of DOCK2 and p110δ during in vitro B cell chemotaxis towards CXCL13, given that DOCK2 contains a PIP3 binding domain. In a first set of experiments, we treated WT B cells separately or in combination with IC-87114 and CPYPP, which blocks the GEF activity of DOCK2 by binding to its catalytic DHR2 domain (15). These data showed that DOCK2 and p110δ comprised two complementary pathways for
CXCR5-mediated chemotaxis, since only simultaneous treatment with both inhibitors completely abolished migration (Figure 2A). A blocking effect of CPYPP and IC-87114 was also observed for CXCL13-induced migration of p110\textsuperscript{dD910A/D910A} and DOCK2\textsuperscript{-/-} B cells, respectively (Figure 2A).

Since inhibitors are often not entirely specific, we generated p110\textsuperscript{dD910A/D910A} x DOCK2\textsuperscript{-/-} mice to corroborate our findings in a genetic model. Double-deficient mice were born at sub-mendelian ratios and showed growth retardation (not shown). Owing to the difficult breeding, we could isolate cells from these mice for only limited amounts of chemotaxis assays. In these experiments, residual migration of DOCK2\textsuperscript{-/-} B cells to 250 nM CXCL13 was abolished when p110\textsuperscript{d} activity was additionally compromised (Figure 2B). Taken together, these data suggest that DOCK2 and p110\textsuperscript{d} act in largely non-overlapping pathways downstream of CXCR5 signaling.
p110δ activity contributes to B cells speed and directionality during follicular migration

CXCR5 is required for B cell entry to B cell follicles (1), where it contributes to fast motility (4). This motility is in large part driven by DOCK2-mediated Rac activation, since DOCK2−/− B cells show substantially reduced interstitial movement (6). Using 2PM of popliteal PLN containing adoptively transferred B cells (4), we confirmed a substantial drop in mean speeds in DOCK2-deficient B cells (from 7.9 ± 4.7 to 4.0 ± 2.9 µm/min for WT and DOCK2−/− B cells, respectively; Figure 3A). This decline in speed was accompanied by broader turning angles and a low motility coefficient (MC), a proxy for a cell’s ability to scan an area (20.7 and 3.6 µm²/min for WT and DOCK2−/− B cells, respectively; Figures 3B, C), in line with our previous observations (6).

We then examined whether p110δ contributed to B cell scanning of B cell follicles in vivo. In contrast to DOCK2−/− B cells, WT and p110δD910A/D910A B cells accumulated efficiently in B cell follicles (Figure 3D, Supplemental Movie 1). However, p110δD910A/D910A B cells moved with decreased speeds and less directionality compared to WT B cells, as measured by meandering index and turning angle distribution (Figures 3E-G). As a result, p110δD910A/D910A B cells had an approximately 50% reduction of their MC compared to WT B cells (Figure 3H). In contrast, interstitial p110δD910A/D910A T cell migration speeds were similar to those of WT T cells (Figure 3I). These data support a contribution of p110δ activity to B cell motility along the FDC network inside B cell follicles.

![Figure 3](image-url)

**FIGURE 3** p110δ contributes to B cell scanning in follicles. (A, B) Average track speeds and relative frequency of WT and DOCK2−/− B cells. (C) Mean displacement versus time of WT and DOCK2−/− B cells. The motility coefficient (MC) is indicated. (D) 2PM image from the FDC (white)-positive B cell follicle containing WT (green) and p110δD910A/D910A (blue) B cells after adoptive transfer in WT recipients. The boxed area is shown magnified for the 0 and 2 min time points, with WT B cell tracks shown in white and p110δD910A/D910A cell tracks shown in red. Number of tracks analyzed: WT, n = 708; p110δD910A/D910A, n = 835; DOCK2−/−, n = 104. Scale bar, 40 µm. (E-G) Average track speeds (E), meandering index (F) and turning angle distribution (G) of WT and p110δD910A/D910A B cells. (H) Mean displacement versus time of WT and p110δD910A/D910A B cells. (I) Average track speeds of WT and p110δD910A/D910A T cells (n = 298 and 370 tracks, respectively) in intravital imaging of lymphoid tissue. Data from one (A, B), two (I) or five (E-H) independent experiments (= recipient mice) and analyzed by an unpaired t-test (A, E, I) or Mann-Whitney test (B, F, G). ***p < 0.001.
Reduced speed and directionality of p110<sup>d</sup>D<sup>910A/D910A</sup> B cells are maintained in the absence of stromal ICAM-1

In addition to CXCR5, LFA-1 contributes to B cell motility on ICAM-1<sup>+</sup> VCAM-1<sup>+</sup> FDCs, whereas β4 integrins play no detectable role (16). In line with this, β2 integrin-dependent <em>in vitro</em> leukocyte migration requires Syk-mediated p110<sup>d</sup> translocation to the leading edge (17). Given the comparable impact of defective LFA-1 and p110<sup>d</sup> activity on dynamic B cell motility parameters, we examined whether p110<sup>d</sup> activity mediated its promigratory effect via LFA-1 activation. In support of this, an analysis of CXCL13-triggered <em>in vitro</em> adhesion to FDC-expressed adhesion molecules uncovered a reduction in p110<sup>d</sup>D<sup>910A/D910A</sup> B cell binding to ICAM-1 but not VCAM-1 (Figures 4A, B). Again, this adhesion defect was restricted to B cells, since p110<sup>d</sup>D<sup>910A/D910A</sup> T cell adhesion to ICAM-1 was not impaired (Figure 4C). We transfected WT and p110<sup>d</sup>D<sup>910A/D910A</sup> B cells into ICAM-1<sup>−/−</sup> recipients, the main stromal LFA-1 ligand used by B cells in lymphoid tissue (16). We hypothesized that WT and p110<sup>d</sup>D<sup>910A/D910A</sup> B cells would show similar migration speeds if p110<sup>d</sup> exerted its promigratory effect via LFA-1. However, we still observed reduced migration speeds, meandering index and increased turning angles in p110<sup>d</sup>D<sup>910A/D910A</sup> B cells compared to WT B cells (Figures 4D–F). As a consequence, their MC remained lower than the one of WT B cells (Figure 4G). These data suggest that the migration defect of p110<sup>d</sup>D<sup>910A/D910A</sup> B cells is largely independent of LFA-1-mediated adhesion to the FDC network. In sum, our data uncover a role for p110<sup>d</sup> activity during B cell migration in lymphoid tissue, which is less pronounced than the effect caused by absence of DOCK2.

Discussion

CXCR5-driven B cell chemotaxis to CXCL13 is critical for the development of humoral immune responses, as it enables efficient surveillance of FDCs and the proper formation of germinal centers (1, 18, 19). Here, we examined the intracellular wiring of CXCR5 that transmits biochemical input into a promigratory response. Our <em>in vitro</em> chemotaxis assays confirmed a critical role for the Rac GEF DOCK2 in mediating robust B cell chemotaxis to CXCL13, while p110<sup>d</sup> participates in a complementary signaling module. These observations were recapitulated <em>in vivo</em>, suggesting the existence of two signaling pathways underlying CXCL13-mediated motility. The requirement of a PI3Kδ-dependent signaling module appears restricted to CXCR5, since migration to CXCR4 and CCR7 ligands was not impaired.

Intravital imaging has uncovered that B cell adhesion in PLN high endothelial venules (HEV) is more strongly attenuated by
the absence of DOCK2 as compared to adhesion in Peyer’s patch (PP) HEV, although in both cases there is a significant reduction in B cell attachment (10). In contrast, lack of PI3Kγ activity mainly affects B cell homing to mesenteric lymph nodes (MLN) and PP, while these cells show normal homing to PLN (12). This may be due to the fact that CXCR5 plays a more prominent role for B cell homing to MLN and PP as compared to PLN, where CCR7 and CXCR4 play compensating roles (20, 21). Thus, the cooperative action of DOCK2 and PI3Kγ activity appears to extend to CXCR5-driven B cell entry into secondary lymphoid organs.

The parallel occurrence of a major, DOCK2-dependent pathway and a minor PI3K-dependent pathway in B cells mirrors observations made in naïve T cells. In T cells, the class IB p110γ isoform mediates DOCK2-independent migration via a pathway involving the PI3K-binding pleckstrin homology (PH)-domain containing Tec family kinase Itk (6, 10, 12, 22). Accordingly, DOCK2-/- x p110γ-/- T cells show no residual migration to CCL21 (10). In combination with the lack of p110δ involvement during naïve T cell migration in vitro and in vivo, our data support a model where p110γ and p110δ catalytic subunits contribute to T and B cell motility in a subset-specific manner. Of note, during CD4+ T cell differentiation to follicular helper T cells (T FH ), p110δ signals downstream ICOSL induce T FH precursor migration into the B cell follicles (23), suggesting context-specific roles for PI3K family members during lymphocyte positioning within lymphoid organs.

It remains incompletely understood how p110δ signaling contributes mechanistically to B cell migration downstream CXCR5, although Rac activation is likely to be required (24). A conceivable scenario is that PI3Kα activates the B cell homologue of Itk, the PH-domain-containing Btk (25). In chronic lymphocytic leukemia (CLL) cell lines, pharmacological blockade of either p110δ or Btk reduces migration to CXCL13 (26, 27). Btk is linked to Vav phosphorylation, leading to downstream WASP activation and F-actin remodeling (28).

Unexpectedly, we found that the defect of p110δ (D910A/D910A) B cell was maintained in lymphoid microenvironment lacking stromal ICAM-1, despite the known involvement of Syk-p110δ signaling during β2-integrin-mediated migration on 2D surfaces (17). A plausible explanation is that akin to naïve T cell migration within lymph node parenchyma, the main role for LFA-1 might be for generation of traction forces without inducing substantial adhesion (29). In the 3D confined environment of lymphoid tissue, substrate adhesion is externally enforced by juxtaposed cells, thus compensating for reduced LFA-1 activity.

The robust DOCK2-driven migration of p110δ (D910A/D910A) B cells to CXCL13 raises the question whether PI3K-mediated signaling has additional roles beyond promoting cell motility. Another open point is whether PI3Kδ signaling might be involved in signal transduction downstream GPR183, although this receptor appears to have an inhibitory effect on CXCR5-mediated migration (4). In T cells, Itk contributes to homeostasis, suggesting a role for PI3K-dependent signaling in maintaining peripheral T cell numbers (22). Similarly, it is conceivable that CXCR5-mediated PI3K activation contributes to B cell homeostasis, in line with the well-documented role of this pathway for survival (30). In addition, the selective integration of p110δ signaling downstream CXCR5, but not other receptors for homeostatic chemokines, might facilitate B cell activation by feeding into the BCR-triggered PI3K-Btk signaling axis. A similar costimulatory signaling pathway was reported for CCL21 during T cell activation (31).

In sum, our data uncover dual signaling pathways mediating physiological CXCR5-triggered B cell motility that underpins rapid detection of cognate antigens presented on FDCs. Given that small tyrosine kinase inhibitors targeting p110δ and Btk are widely used in the treatment of leukemias (32–34), it is of clinical interest to understand potential implications on the patients’ immune system.

Materials and methods

**Mice**

Six to 12-week-old male and female DOCK2-/- (5), p110δ (D910A/D910A) (35), DOCK2-/- x p110δ (D910A/D910A) and ICAM-1-/- (36) mice on the C57BL/6 background were bred at the University of Bern and Fribourg. Sex-and age-matched C57BL/6 mice (Harlan, The Netherlands) were used as WT lymphocyte donors or recipient mice. All experiments were performed in accordance with federal animal experimentation regulations and approved by the corresponding cantonal committee.

**Isolation and labeling of primary lymphocytes**

B or T cells from PLN, MLN and spleen were purified by negative immunomagnetic cell sorting according to manufacturer’s instructions (Dynal or Stemcell technologies; purity of >95%). For intravital imaging experiments, purified B or T cells (5 x 10⁶) from C57BL/6 or genetically modified mice were fluorescently labeled for 15 min at 37°C with Cell Tracker blue (20 µM final concentration), Cell Tracker orange (5 µM), Cell Tracker green (3 µM) or CFSE (2.5 µM), washed and injected intravenously into sex-matched C57BL/6 recipient mice, together with 10-15 µg Alexa633-conjugated MECA-79 to label high endothelial venules. Dyes were switched between experiments to control for non-specific effects.
Chemotaxis

CCL21 and CXCL12 were from Peprotech, and CXCL13 was purchased from R&D systems. Chemotaxis assays were carried out using Transwell chambers (5 µm pore size; CoStar) adding 100 µl cell suspension (5 x 10⁶ cells/ml) in complete medium (RPMI/10% FCS/standard supplements) to the top chamber and indicated amounts of chemokine in the bottom chamber. After 2 h at 37°C, 7% CO₂, the percentage of migrated cells was calculated by flow cytometry after comparing with a precalibrated bead standard (Sigma-Aldrich) and correcting for variations in input concentrations. The DOCK2 inhibitor CPYPP (Selleck) was used at 40 µM throughout the chemotaxis assay (15). The isoform-specific PI3K inhibitors TGX221 (0.1 µM final conc.; Tocris), PI-103 (1 µM; Tocris), AS604850 (1 µM; Selleck), and IC-87114 (0.5 µM; Selleck) were present throughout the chemotaxis assay.

Adhesion assay

Adhesion assays were performed as described (10). In brief, purified B or T cells were allowed to settle on 8-well-slides coated with 1.5 µg/ml murine ICAM-1 or 2.5 µg/ml VCAM-1 (R&D Systems). Chemokine was added at a final concentration of 1 µM for 3 min. Slides were rinsed with PBS to wash off unbound cells, fixed in glutaraldehyde, and the number of adherent cells was determined at the site of chemokine addition.

Two-photon intravital microscopy

Fluorescently labeled WT and genetically modified B cells were adoptively transferred into WT or ICAM-1−/− recipients 12-48 h before 2PM recording. In some experiments, PE-conjugated anti-CD35 mAb (0.5 µg in 10 µl PBS/mouse) was injected into the footpad 12 h before 2PM to label the FDC network of the draining popliteal PLN. Recipient mice were surgically prepared to expose the right popliteal PLN, which was kept at 36-38°C. Mice were then transferred to an Olympus BX50WI fluorescence microscope attached to a 2PM scanner (TrimScope system, LaVision Biotec, Bielefeld, Germany) equipped with a 20X objective (Olympus, NA 0.95). For four-dimensional analysis of cell migration, 8-16 z-stacks (spacing 4 µm) of 200-300 µm x-y sections were acquired every 20 s for 20 to 30 min, with typically 3-4 distinct areas recorded per preparation. Image sequences were transformed into volume-rendered four-dimensional movies using Volocity (Perkin Elmer) or Imaris (Bitplane), which was also used for semi-automated tracking of cell motility in three dimensions. From x, y, and z coordinates of cell centroids, parameters of cellular motility were calculated as described previously. In brief, the track speed is depicted as average speed, with each dot representing one track. Owing to the large number of tracks, they are shown as box and whisker plots with whiskers covering 1-99% of data points. For turning angles and motility coefficients, we used MatLab scripts kindly provided by Dr. Sarah Henrickson and Prof. Ulrich H. von Andrian (Harvard University, Boston, USA). In some experiments, purified WT and p110δD910A/D910A T cells were transferred into WT recipients and their migratory behavior was analyzed in the T cell area as above.

Statistical analysis

The student’s t-test or ANOVA were used to determine statistical significance (Prism, GraphPad). Statistical significance was set at p < 0.05.

Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Ethics statement

The animal study was reviewed and approved by Canton of Fribourg and the Canton of Bern.

Author contributions

SW, BS, and AM performed experiments. JS supervised the work and wrote the manuscript with input from all coauthors. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2022.982383/full#supplementary-material

SupPLEMENTAl MOVIE 1

2PM image sequence showing WT (green) and p110

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