FGD2, a CDC42-specific Exchange Factor Expressed by Antigen-presenting Cells, Localizes to Early Endosomes and Active Membrane Ruffles

Members of the Fgd (faciogenital dysplasia) gene family encode a group of critical guanine nucleotide exchange factors (GEFs), which, by specifically activating Cdc42, control cytoskeleton-dependent membrane rearrangements. In its first characterization, we find that FGD2 is expressed in antigen-presenting cells, including B lymphocytes, macrophages, and dendritic cells. In the B lymphocyte lineage, FGD2 levels change with developmental stage. In both mature splenic B cells and immature bone marrow B cells, FGD2 expression is suppressed with developmental stage. In mature splenic B cells, FGD2 expression is suppressed with developmental stage. In both mature splenic B cells and immature bone marrow B cells, FGD2 expression is suppressed with developmental stage. In both mature splenic B cells and immature bone marrow B cells, FGD2 expression is suppressed with developmental stage.

**Fgd2** (faciogenital dysplasia 2) is a novel gene discovered by PCR amplification using primer sequences based on the Fgd1 gene (1) and is a member of a small subfamily of predicted RhoGEFs (including Fgd1, Fgd3, Fgd4 (Frabin), Fgd5, and Fgd6) with C-terminal amino acid homology but highly divergent N-terminal extensions. The Fgd subfamily is part of a larger family of 69 genes with homology to Dbl, a transforming gene identified in diffuse B cell lymphomas (2). RhoGEFs are activators of the Rho (Ras homology) subfamily of GTPases, such as Rac1 to -3, RhoA to -G, and Cdc42, because they are GDP/GTP exchange factors (GEFs). As such, they catalyze the release of bound GDP from the GTPases, resulting in the formation of GTP-bound active proteins able to interact with downstream effectors (2). Depending on their subcellular localization and interaction with specific GTPases, they can thus control disparate biological activities, including adhesion, mitosis, cellular polarization, and vesicle trafficking (3).

All Dbl RhoGEF family members carry a signature active site motif composed of tandem Dbl homology (DH) and pleckstrin homology (PH) domains. Substrate specificity is dictated by a “specificity patch” within the DH domain (4), with some RhoGEFs demonstrating specificity for particular GTPases (e.g. TRIO (Rac1) (5) and intersectin (Cdc42) (4)), whereas others, such as the VAV proteins and Dbl, have broad activity for multiple GTPases (6–8). Further domains that influence function, molecular interactions, and cellular localization distinguish RhoGEFs from one another. Among Dbl family proteins, Fgd family members are unique in carrying a FYVE motif (shared in Fab1, YotB, Vac1p, and EEA1) and an additional PH domain at the C terminus.

FYVE domains are zinc finger lipid binding motifs that often target proteins to vesicles by interaction with phosphatidylinositol 3-phosphate (9–13), a lipid that is enriched on endosomal membranes (12). PH domain-containing proteins often bind to phosphatidylinositol 3,4,5-trisphosphate (PI(3,4,5)P3), leading to phosphatidylinositol 3-phosphate (9–13), a lipid that is enriched on endosomal membranes (12). PH domain-containing proteins often bind to phosphatidylinositol 3,4,5-trisphosphate (PI(3,4,5)P3), leading to

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**Abbreviations:**

- GEF, guanine nucleotide exchange factor
- BCR, B cell antigen receptor
- PH, pleckstrin homology
- DH, Dbl homology
- PI(3,4,5)P3, phosphatidylinositol 3,4,5-trisphosphate
- TRX, thioredoxin
- TRIO (Rac1) (5) and intersectin (Cdc42) (4)
- TRIP (RhoA) (3)
- PI(3,4,5)P3, phosphatidylinositol 3,4,5-trisphosphate
- TRX, thioredoxin
- GM-CSF, granulocyte-monocyte colony-stimulating factor
- LPS, lipopolysaccharide
- GST, glutathione S-transferase
- EGFP, enhanced green fluorescence protein
function of FGD2. A recent study indicates that a mutant form of Fgd2 contributes to t-complex ratio distortion, a phenomenon found in some wild strains of mice where decreased motility of sperm is caused by the actions of several genes located in a variant region of chromosome 17 (22). The Fgd2 locus also maps to Idd23, a region of chromosome 17 where the C57BL/6 allele is implicated in the protection from diabetes in the NOD mouse model of diabetes (23).

In this paper, we present the first biochemical characterization of FGD2, including an assessment of its predominant expression in leukocyte subsets, regulation in B cells by B cell receptor (BCR) signaling, enzymatic activity, and subcellular localization. We show that FGD2 is the sole member of the characterized Fgd family that can localize to early endocytic vesicles.

**EXPERIMENTAL PROCEDURES**

**Cloning Mouse Fgd2**—Total RNA was harvested from mouse spleen (strain B10.D2/SnJ) using TRIzol (Life Technologies), and cDNA was generated using the SuperScript First Strand synthesis system for reverse transcription-polymerase chain reaction (Invitrogen) according to the manufacturer’s instructions. Fgd2 cDNA was amplified using PLATINUM Pfx DNA polymerase enzyme (Invitrogen) according to the manufacturer’s instructions with the following primers: 5’-TACTCAAG-CTTAGGATGGAGCGCTGTGAG and 5’-TGTACAC-TCGAGATTTCATGATCCAGGGATA. The sequence for TCGAGATTTCATGATCCAGGGATA. The sequence for the mouse Fgd2 from B10.D2 mice had several changes from the published sequence (1) and concurred with the C57BL/6 genomic sequence and cDNA sequences AK042260 and AY301264 (NCBI GenBankTM) with two exceptions. The predicted amino acids Ser39 and Glu543 of C57BL/6J origin were found in some wild strains of mice where decreased motility of sperm is caused by the actions of several genes located in a variant region of chromosome 17 (22). The Fgd2 locus also maps to Idd23, a region of chromosome 17 where the C57BL/6 allele is implicated in the protection from diabetes in the NOD mouse model of diabetes (23).

Following 10 days of culture in 200 units/ml GM-CSF, nonadherent bone marrow cells were then cultured a further 2 days in GM-CSF (100 units/ml) and 1 μg/ml lipopolysaccharide (055: B5) (LPS; Sigma).

**Western Analysis of FGD2 Expression**—Whole cell lysates from thymocytes, pooled lymph nodes, total bone marrow, bone marrow-derived macrophages, bone marrow-derived dendritic cells, splenocytes, and splenocytes either B220-enriched or -depleted using magnetic beads (Miltenyi Biotec) were generated from B10.D2 mice using 0.5% Nonidet P-40 lysis buffer containing 10 mM Tris, 150 mM NaCl, 0.1 mM EDTA, and 1 mM phenylmethylsulfonyl fluoride and Complete™ protease inhibitors (Roche Applied Science). Insoluble nuclei were removed by centrifugation, and lysates were subjected to SDS-PAGE under reducing conditions, followed by Western blotting and detection using a polyclonal rabbit anti-FGD2 antiserum or glyceraldehyde-3-phosphate dehydrogenase antibody. A mouse tissue INSTA-Blot (Imgenex) containing immobilized brain, heart, small intestine, kidney, liver, lung, muscle, stomach, spleen, ovary, and testis lysates was also probed with the polyclonal rabbit anti-FGD2 antiserum, according to the manufacturer’s instructions.

**Expression with Stimulation**—Immature B cells from 3-83 BCR Tg mice were expanded from bone marrow using interleukin-7 cultures as described (27). One day after interleukin-7 removal, 20 μg/ml either anti-BCR antibody (S23) or control (Y3) antibody was added to the cells for the indicated times. Mature B cells were purified from spleen and lymph nodes from B10.D2 mice using negative selection with CD43 MACS beads and columns (Miltenyi Biotec). CD43-B cells were cultured at 1 × 10⁷ cells/ml with either 10 μg/ml goat F(ab)2 anti-mouse IgM (Jackson Immunochemicals) or LPS (1 μg/ml) and incubated at 37 °C for various time points prior to lysis and analysis by Western blotting.

**Real Time PCR Analyses of mRNA Expression**—Pro-B cells were isolated by cell sorting from bone marrow cells of

5 D. Nemaze, unpublished results.
RAG1−/− mice, where B220+CD43+ were positively selected. Pre-B cells from bone marrow of B10.D2SnJ mice were sorted using the markers B220+ CD43− IgM− IgD−. Immature B cells were isolated from B10.D2 mice carrying the 3-83 BCR transgene (28) and were sorted for B220+ IgM+ IgD− CD43− marked cells. Immature B cells undergoing receptor editing were isolated from B10.Br 3-83 transgenic bone marrow, where B10.Br mice express the BCR antigen H-2k in the bone marrow. Cells were sorted for B220+ IgD− CD43− IgM−. B220+ cells from spleen and pooled lymph nodes were isolated using B220 magnetic bead separation (Miltenyi). Total RNA was purified from sorted B cell subsets using TRIzol (Invitrogen), and contaminating genomic DNA was removed by DNase I digestion. First strand cDNA was generated using oligo(dT) and the Superscript III first strand synthesis system (Invitrogen) before Fgd2 and β-actin were amplified. A 158-bp ampiclon for actin was amplified using primers 5′-gcatgctgctaggaagc and 5′-cctgctcatgctgcatac annealing at 57 °C. Fgd2 was amplified using primers 5′-AGGAACCTAGGAGAAGGGGTC and 5′-gctgctaggaagcctgccg annealing at 57 °C, yielding a 189-bp product. The PCRs were performed in triplicate using the 7900 HT Applied Biosystems machine and quantified using the Sybr green PCR kit (Qiagen). Ct values for each reaction were generated using the SDS software (Applied Biosystems). Fgd2 Ct values were subtracted from actin, and mean results were expressed as the exponential product ratio.

Northern Analysis of Fgd2 Expression—20 μg of total RNA isolated from immature B cells cultured with or without anti-BCR antibody for 24 h was electrophoresed under denaturing conditions. After photographing the RNA gel stained with ethidium bromide, the RNA was transferred to a nylon Zeta probe membrane (Bio-Rad) before UV cross-linking and probing with an Fgd2 cDNA probe generated using the primers 5′-CTCAGGAAGGTGGTACCCCG and 5′-AACCAGGTAGCGTTCACTTG.

Cdc42 Activation Assay—COS-7 cells were transiently transfected with Lipofectamine2000 (Invitrogen) mixed with cDNA encoding Myc-tagged Cdc42, EGFP, EGFP-tagged Fgd2, or EGFP-tagged Fgd2 mutants. The dominant active mutant EGFP-Cdc42Q61L and the dominant negative mutant, EGFP-Cdc42T17N, were used as positive and negative controls, respectively. Active Cdc42 was assessed by a pull-down assay, as described (29). Briefly, 24 h post-transfection, cells were lysed with a buffer containing 50 mM Tris, pH 7.8, 200 mM NaCl, 5 mM MgCl2, 1 mM dithiorthreitol, 1% Nonide P-40, 10% glycerol, and protease inhibitors (Roche Applied Science). Lysates were precleared with GST bound to glutathione beads for 10 min, prior to incubation for 1 h with ACK1 (activated Cdc42-associated kinase 1)-Crib (Cdc42- and Rac1-interacting binding) (amino acids 439–502)-GST fusion protein bound to glutathione beads. After washing three times with wash buffer (25 mM Tris, 40 mM NaCl, 30 mM MgCl2, 1 mM dithiorthreitol, 1% Nonide P-40), beads were subjected to SDS-PAGE and Western blotting. Bead-associated active Cdc42 was detected using anti-Myc and anti-EGFP antibodies. Rac1 and RhoA activity assay kits containing PAK-Crib GST and Rhotekin-RBD GST beads, respectively, were purchased from Upstate Bio-technology (now Millipore) and used according to the manufacturer’s instructions.

JNK1 Activity Assays—HEK293 or COS7 cells were transiently transfected with FLAG-JNK1 together with EGFP or various EGFP-Fgd2 mutants in the presence of either Cdc42 or Rac1. JNK1 activity assays were performed as described (30) by immunoprecipitation of JNK1 with anti-FLAG beads (Sigma) and incubation of [γ-32P]ATP with a GST-ATF-2 fusion protein as substrate. Phosphorylated ATF-2 was determined by autoradiography, and the amounts of expressed proteins were determined by Western blotting of total cellular lysates.

Confocal Microscopy Imaging of Fgd2 Localization—Transiently transfected HeLa cells that had adhered to polylysine- or fibronectin-coated glass coverslips were fixed with 3.4% paraformaldehyde. In some cases, transfected HeLa cells were incubated with either 50 mM wortmannin (Calbiochem) in DMSO or DMSO alone for 30 min prior to fixation and staining. For actin stains, the cells were permeabilized with 0.2% Triton X-100 (phalloidin and cortactin stains). For intracellular compartment stains, the cells were permeabilized with 0.05% saponin prior to staining. After the samples were mounted with Immuno-Fluo mounting medium (MP Biomedicals Inc.), coverslips were visualized by fluorescence microscopy with an MC1024 (Bio-Rad) confocal laser microscope. The EGFP-Fgd2 colocalization with intracellular stained compartments was determined using LSM examiner software (Zeiss).

Phospholipid Binding Membranes—Recombinant His-tagged thioredoxin-Fgd2 (pET32 expression vector (Novagen)) or fusion partner thioredoxin alone were expressed in Escherichia coli BL21 (DE3) and purified using nickel affinity chromatography. The recombinant Fgd2 protein was insoluble and was solubilized with 8 M urea and gradually dialyzed into a final buffer of 25 mM Tris, pH 8, 150 mM NaCl, 10 mM 2-mercaptoethanol, and 0.1% Triton X-100 over 3 days. Purified proteins were incubated with PIP Array membranes (Molecular Probes and Invitrogen) at 0.5 μg/ml for 2 h according to the manufacturer’s instructions. After washing, bound proteins were detected with an anti-His epitope antibody (Sigma) and chemiluminescence.

RESULTS

Fgd2 Is Expressed in Primary Lymphoid Cells—When immature B cells encounter self-antigen, they can undergo secondary recombination of the κ light chain locus, to produce an innocuous surface antigen receptor, a tolerance process known as receptor editing (31). To generate large numbers of immature B cells with uniform receptors, we cultured bone marrow from 3-83 BCR transgenic mice in which immature B cells express a prerearranged receptor against H-2Kk antigen (32). In order to identify genes that were differentially expressed between immature B cells and immature B cells forced to undergo receptor editing by exposure to anti-BCR antibodies in vitro, we performed a PCR-based total gene expression analysis screen (33). In this screen, we identified Fgd2 as a candidate RNA that was expressed in immature B cells but was poorly expressed in editing B cells. These findings were also confirmed by Affymetrix chip analysis of U94B arrays with the probe set 115754_at (data not shown). Perusal of the Symatlas data base (34) indicated
that Fgd2 mRNA expression was concentrated in lymphoid tissues rich in B cells.

We therefore wanted to assess FGD2 protein expression in lymphoid tissues or isolated leukocyte subsets. To this end, an FGD2-specific rabbit antiserum was prepared. The antiserum’s specificity for FGD2 was clear, because, although it could bind to recombinant epitope-tagged FGD2 expressed in HEK293 cells in Western blots, it failed to detect the related FGD1 protein (Fig. 1A, lanes 5 and 6) and FGD3 (not shown). By contrast, both FGD1 and FGD2 could be detected when probed with antibodies directed to their N-terminal FLAG epitope tag (Fig. 1A, lanes 8 and 9). Furthermore, FGD2 has a distinctly lower predicted molecular weight (Mr, 74,600) than all other Fgd family members, including FGD1 (Mr, 105,000) (Fig. 1A, lanes 8 and 9). In addition, the preimmune sera did not detect any proteins in the HEK293 lysates (Fig. 1A, lanes 1–3).

FGD2 Protein Function and Cellular Localization

This rabbit-derived antiserum was used in Western blotting of whole cell lysates prepared from mouse thymus, pooled lymph nodes, bone marrow, and spleen and magnetically separated CD45R+ (B220+) and B220− splenic cells. No reactivity was detected in lysates of unfractonated bone marrow or thymus (Fig. 1B, lanes 1 and 3), whereas high level expression was evident in lymph nodes and spleen (Fig. 1B, lanes 2 and 6), consistent with the high levels of mRNA expressed in these latter tissues. B220+ splenic cells were enriched, whereas B220− splenic cells were depleted of immunoreactive FGD2 protein (Fig. 1B, lanes 7 and 8), indicating that B cells are the major source of FGD2 in the spleen. In addition, in lysates prepared from macrophages and dendritic cells that were expanded from bone marrow by stimulation with granulocyte colony-stimulating factor or GM-CSF, respectively, FGD2 expression was detectable (Fig. 1B, lanes 4 and 5). The expression of FGD2 in nonleukocyte tissues was tested by probing an INSTA-Blot mouse tissue membrane (Imgenex). We were unable to detect FGD2 protein in brain, liver, kidney, testis, muscle, ovary, or small intestine but did confirm splenic expression in this assay (data not shown). These experiments indicate that leukocytes express FGD2, and that in lymphoid tissues, B lymphocytes and myeloid cells, in particular macrophages, also express the protein.

Fgd2 mRNA Expression and Regulation in B Lymphocyte Development—Because we were initially interested in identifying genes that were regulated during B lymphocyte development and selection, and Western blotting and array analysis indicated that Fgd2 expression was particularly high in B cells, we carried out a more detailed analysis of the developmental regulation and response to antigen receptor signaling of its RNA expression. A Northern blot assay confirmed that Fgd2 is differentially expressed in interleukin-7-expanded primary immature B lymphocytes. Fig. 2A shows that immature B cells treated with anti-BCR antibodies (+) for 24 h in vitro had low level expression of Fgd2 mRNA relative to control IgG-treated cells (−). To determine if BCR signaling affected Fgd2 expression in vivo, immature bone marrow B cells were directly sorted from 3-83 BCR transgenic mice that expressed or lacked cognate autoantigen (32) (Fig. 2B). Immature bone marrow B cells differed in their levels of Fgd2 mRNA depending on whether cognate self-antigen was present (Fig. 2B, Imm or present (Edit)). These data confirmed in vivo studies that BCR ligation by self-antigen suppressed Fgd2 expression in immature B cells. We also compared Fgd2 levels in other B lineage subsets. Sorted pre-B cells (CD43− B220− slg−) expressed low levels of Fgd2, whereas pro-B cells (CD43− B220+ slg−) expressed high levels. Mature B lymphocytes from spleen and lymph nodes expressed high levels of Fgd2 mRNA (Fig. 2B). We conclude that Fgd2 expression changes strikingly with preimmune developmental progression and that BCR ligation can alter Fgd2 expression in the bone marrow, probably by inducing developmental arrest.

FGD2 Protein Expression in B Cells Is Regulated by a BCR-specific Signal—Since we found Fgd2 mRNA to be repressed in immature B cells undergoing tolerance-induced receptor editing, we were interested in whether this was also true for the protein level in both immature and mature B cells. We therefore used Western blotting with FGD2 antiserum to assess

**FIGURE 1.** FGD2 protein expression in lymphoid organs and leukocyte subsets. A, specificity of anti-FGD2 serum. FGD2-FLAG (lane 2) or FGD1-FLAG (lane 3) was expressed in HEK293 cells, cell lysates were prepared, and expression was detected by immunoblot using either preimmune serum (left), anti-FGD2 polyclonal serum (center), or FLAG antibodies (right). As a negative control, empty vector was also transiently transfected and cell lysate prepared (Mock lane 1). B, lymphoid tissues and cell lysates were electrophoresed, and Western blotted membranes were probed with FGD2 antiserum (top) or glyceraldehyde-3-phosphate dehydrogenase antibody (bottom). Macrophages (BMMφ) were generated in vitro by expansion from bone marrow in macrophage colony-stimulating factor (GM-CSF); lane 4). Dendritic cells (BMDC) were similarly generated by expansion with GM-CSF and matured with LPS (lane 5). Splenocytes were fractionated into B220-enriched (B220+) or B220-depleted (B220−) samples by magnetic beads prior to lysis.
FGD2 Protein Function and Cellular Localization

FGD2 protein expression and regulation in primary mouse B cells stimulated with anti-BCR antibody. Immunoblotting analysis carried out with lysates from both immature (Fig. 3A) and mature splenic B cells (Fig. 3B) cultured up to 24 h with or without anti-BCR antibodies revealed that anti-BCR-treated cells had strikingly lower FGD2 protein expression (Fig. 3, A and B), which correlated with the changes observed in RNA expression (Fig. 2A). FGD2 protein expression in immature B cells was suppressed after 24 h of BCR stimulation (Fig. 3A) but was unaffected by the presence of a control antibody (Fig. 3A, right panels) or glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (Fig. 3B). Real-time PCR was performed, and the mean and S.D. of Fgd2 and actin amplicons were calculated from triplicate samples.

To test if the repression of FGD2 was specific to a BCR-mediated signal and is suppressed by BCR ligation in vitro and in vivo, immature B cells were stimulated with (+) or without (−) anti-BCR antibodies for 24 h. Left, ethidium bromide-stained gel revealing 18 and 28 S ribosomal RNA bands; right, Northern blot detection of Fgd2 message in total RNA of immature bone marrow B cells cultured up (+) or without (−) anti-BCR antibodies for 24 h. Both immortalized and primary cultures of immature B cells were challenged with 20 μg/ml idiotypic anti-BCR or isotype control antibody for the indicated times, and protein expression was analyzed by Western blotting. Immature splenic B cells were stimulated with anti-BCR antibody (goat F(ab)_2, anti-IgM 10 μg/ml) or LPS (1 μg/ml) for the times indicated, and FGD2 (top) or glyceraldehyde-3-phosphate dehydrogenase (GAPDH; bottom) protein levels were determined by Western blotting.

**Figure 3.** FGD2 is regulated by a BCR-specific signal. A, immature B cells generated in interleukin-7 culture of 3-83 BCR transgenic bone marrow were challenged with 20 μg/ml idiotypic anti-BCR or isotype control antibody for the indicated times, and protein expression was analyzed by Western blotting. Immature splenic B cells were stimulated with anti-BCR antibody (goat F(ab)_2, anti-IgM 10 μg/ml) or LPS (1 μg/ml) for the times indicated, and FGD2 (top) or glyceraldehyde-3-phosphate dehydrogenase (GAPDH; bottom) protein levels were determined by Western blotting.

FGD2 contains a FYVE domain, and the binding of phospholipids via FYVE domains is associated with the endosomal localization of some proteins, we compared the intracellular localization of FGD2 with the co-staining of intracellular compartment markers. Imaging experiments in transfected HeLa cells indicated that about half of EGFP-FGD2 present on vesicular structures associated with early endosomes, as judged by EEA1 colocalization (Fig. 4, upper right panel). In contrast, little colocalization of FGD2 with either LAMP2 or GM130 staining was observed, indicating poor colocalization of FGD2 with lysosomes or the cis-Golgi, respectively (Fig. 4, middle and lower panels). The same intracellular localization of FGD2 was observed when linked to a different tag (FLAG epitope) and stained for FLAG and EEA1 (data not shown).

We also observed concentration of FGD2 on membrane areas that appeared to be ruffles. We co-stained EGFP-FGD2-expressing cells with phalloidin to highlight the areas of actin-rich membrane ruffles. Strong co-localization of EGFP-FGD2 with phalloidin was observed (Fig. 4, bottom right panel). In HeLa cells overexpressing FGD2, we also observed increases in the numbers of membrane ruffles at steady state, but this was not consistent across many experiments.
Localization of FGD2 to Early Endosomes Is FYVE Domain-dependent—To determine if the conserved FYVE domain directed the vesicular staining pattern of FGD2, we assessed the endosomal association of FGD2 and mutant molecules in which the FYVE domain had been mutated (FGD2FYVEKT; Q454K/W455T). Although wild type EGFP-FGD2 still demonstrated a vesicular pattern co-staining with EEA1 (Fig. 5A, upper panel), mutation of the FYVE domain abrogated the co-localization with this early endosomal marker (Fig. 5A, middle panel). However, a functional DH domain is not required for endosomal localization as the FGD2<sub>GEEFAA</sub> DH domain mutant co-localized with EEA1 (Fig. 5A, lower panel). Mutants used in this study are depicted in a diagram (Fig. 5C).

The affinity of FYVE domains for endosomal vesicles requires phosphorylated phosphatidylinositol (phosphatidylinositol 3-phosphate) and is regulated by phosphatidylinositol 3-kinase activity (11). We therefore assessed the FGD2 endosomal localization in the presence and absence of the broad phosphatidylinositol 3-kinase inhibitor, wortmannin. As shown in Fig. 5B, the vesicular pattern of staining of EGFP-FGD2 was largely blocked when transfected HeLa cells were cultured for 30 min in the presence of wortmannin (lower panels) but was unchanged by incubation of the cells with the solvent DMSO alone (upper panels). The treatment of cells with wortmannin and other more specific class III phosphatidylinositol 3-kinase inhibitors like 3-methyladenine has been shown to affect the localization of FYVE domain-containing proteins like EEA1, leading to alterations in the morphology of early endosomes but not their disappearance (38, 39). In our cells, the wortmannin treatment had the expected effects on EEA1 localization (Fig. 5B, lower panel).

We conclude that a proportion of FGD2 is recruited to endosomal membranes via the FYVE domain and by the presence of phosphatidylinositol 3-phosphate or other phosphatidylinositides generated by phosphatidylinositol 3-kinase activity.

FGD2 Recruitment to Membrane Ruffles is PH Domain-dependent—Both FGD1 and FGD4 (Frabin) are recruited to the actin cytoskeleton within membrane ruffles either directly, by binding actin (40), or indirectly, through interactions with proteins like cortactin (16).

Since FGD2 lacks an identifiable actin or cortactin binding domain we speculated that the C-terminal PH domain might mediate its recruitment to membrane ruffles. To test this hypothesis, we expressed an EGFP-FGD2 mutant lacking the C-terminal PH domain (FGD2<sub>ΔPH2</sub>) in HeLa cells and co-stained these cells for cortactin, a protein enriched in areas of newly polymerized actin filaments. In contrast to wild type FGD2, the FGD2<sub>ΔPH2</sub> mutant remained predominantly cytoplasmic and was not concentrated in membrane ruffles (Fig. 6).

Its localization to early endosomes was not affected by the deletion of this second PH domain. On the contrary, it appeared as if the vesicular staining pattern was increased in FGD2<sub>ΔPH2</sub> expressing cells, and approximately half of these vesicles co-stained with the endosomal marker EEA1. Given that PH domains are known to specifically bind to phosphoinositides, our experiments are consistent with a role of the C-terminal PH 3-kinase activity.
FGD2 has RhoGEF activity for Cdc42—Other Fgd family members have been shown to have exchange activity toward Cdc42 directly (FGD1 (11) and FGD4 (17, 18)) and Rac1 indirectly (FGD4 (18)), suggesting that FGD2 might have similar specificity. Alignment of the “specificity patch” region of the DH domain sequences of FGD2 with other Cdc42-specific RhoGEFs (ASEF, h-PEM2, intersectin) further supported this prediction (Fig. 8A); a conserved leucine, which permits effective binding of RhoGEFs to Cdc42 (4), was also conserved in FGD2 and all other Fgd family members (Fig. 8A). To experimentally test this potential regulation of Cdc42 by FGD2, we performed pull-down experiments using lysates of COS-7 cells transiently transfected with Myc-tagged Cdc42 and either EGFP, EGFP-tagged FGD2, or EGFP-tagged FGD2 mutants (Fig. 8B). The Myc-Cdc42 levels were monitored by Western blot in whole cell lysates (WCL) or after pull-down with ACK1-GST beads (IP; ACK). GTP-bound active Cdc42 is specifically pulled down by the ACK1-GST beads in this assay, since an EGFP-tagged constitutively active GTP-bound mutant of Cdc42 (Cdc42DAH11002) was brought down by the beads, but the GDP-bound mutant of Cdc42 (Cdc42DN) was not (Fig. 8B, left panels). Compared with the EGFP protein control, EGFP-FGD2 appeared to promote association of the ACK1-GST with Cdc42 (Fig. 8B, right panels), suggesting that it elevates the level of active Cdc42 in the cell lysate. This elevation of Cdc42 GTP levels was also observed when cDNA of an EGFP-tagged FYVE domain mutant of FGD2 (FGD2FYVEKT) was overexpressed (Fig. 8B, right panels). We therefore believe that a functional FYVE domain is not required for the activation of Cdc42 by FGD2. However, an EGFP-tagged truncation mutant of FGD2, consisting of the DH-PH domains only (DHP), did not appear to activate Cdc42 in this assay (Fig. 8B, right panels), indicating that additional domains are necessary for full FGD2 activity and interaction with Cdc42. Full-length FGD2 was found to mediate Cdc42 activation in several additional experiments (not shown), using either theACK1-Crib GST or WASP-Crib GST fusion proteins in the pull-down assay. We did not see evidence of endogenous Cdc42 activation in COS-7 cells by recombinant FGD2 expression, indicating that in these cells, the levels of active Cdc42 were too low to be reproducibly measured. In addition, we did not see consistent evidence of activation of either Rac1 or RhoA using PAK1-Crib GST or Rhotekin RBD-
FGD2 Activates JNK1 via Cdc42—Guanine exchange factors can direct downstream signaling of cognate GTPases (41); therefore, we wished to test if FGD2 could activate mitogen-activated protein kinase activity via Cdc42. In co-transfection studies, overexpression of EGFP-tagged FGD2, but not EGFP alone, led to increased JNK1 activity in the presence of Cdc42 but not Rac1 (Fig. 9A). The elevated JNK1 activity, as measured by the phosphorylation of the substrate, GST-ATF-2, was dependent on wild type FGD2 sequence in the DH domain, since a DH mutant of FGD2 (FGD2<sup>GEFΔA</sup>) had a diminished effect on JNK1 activation (Fig. 9B, lane 3). In contrast, mutation of residues in the FYVE domain (FGD2<sup>FYVEKT</sup>) had no effect on the enhancement of JNK1 activity (Fig. 9B, lane 4). A constitutively active mutant of Cdc42, EGFP-Cdc42<sup>Q61L</sup> demonstrated the highest activation of JNK. This supports the conclusion that FGD2 utilizes the DH domain to activate JNK1 through Cdc42 but not Rac1, despite the ability of active Rac1 to also mediate JNK1 activation (Fig. 9B, lane 10).

DISCUSSION

The present study provides the first characterization of FGD2 expression, cellular localization, and protein function. Our data support the predictions that mouse FGD2 has RhoGEF activity and can regulate Cdc42 and downstream JNK1 activation. Furthermore, we have discovered that Fgd2 has striking developmental and tissue-restricted expression in antigen-presenting cells of the immune system. Particularly, B lymphocytes express high levels of FGD2, but significant expression was also detected in bone marrow macrophages and to lower levels in mature bone marrow dendritic cells. In both immature and mature B cells, FGD2 expression is down-regulated by prior B cell receptor signaling. Another B cell mitogen, LPS, fails to do so, suggesting that FGD2 expression in B cells is regulated specifically by BCR signals. The down-regulation of Fgd2 message after BCR cross-linking does not appear to be permanent, however, since memory B cells also appear to express high levels of Fgd2 mRNA (data from gene expression omnibus accession number GDS1695). These data indicate that Fgd2 mRNA expression remains high in resting B and memory cells but declines upon differentiation into plasma cells. Finally, we show that FGD2 is

GST beads (supplemental Fig. 1). Collectively, these results support the conclusion that when expressed exogenously, FGD2 has RhoGEF activity specific for Cdc42.

FIGURE 6. The C-terminal PH domain drives FGD2 co-localization with cortactin in membrane ruffles. HeLa cells expressing either EGFP alone (top panel), EGFP-FGD2 fusion protein (second panel), EGFP-FGD2<sup>FYVEKT</sup> (third panel), or EGFP-FGD2<sup>S PH2</sup> (fourth panel) were grown on coverslips prior to fixation and permeabilization with Triton X-100. The samples were stained for cortactin, a protein localizing to newly polymerized actin filaments, using a monoclonal antibody, followed by Alexa 586-goat anti-mouse IgG (red). Cells expressing EGFP-FGD2<sup>S PH2</sup> were also stained for the endosomal marker EEA1 (bottom).
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associated with early endosomes in a phosphatidylinositol 3-kinase-dependent manner and with plasma membrane ruffles, suggesting multiple possible avenues for regulation of activity and cellular function. Since the amino acid sequences of human and mouse FGD2 are 83% identical and are expressed in a similar tissue distribution (based on the Symatlas data base (42)), we predict human FGD2 probably carries out the same biochemical and biological functions as the mouse FGD2 protein studied here.

FIGURE 7. Recombinant FGD2 directly binds to phospholipids. His-thioredoxin protein alone (TRX; left) or His-thioredoxin-FGD2 fusion protein (TRX-FGD2; right) was incubated with membranes spotted with listed concentrations of phospholipids. After washing, bound protein was detected with an anti-His antibody and chemiluminescence. PI, phosphatidylinositol; PI(3)P, phosphatidylinositol-3-phosphate; PI(4)P, phosphatidylinositol-4-phosphate; PI(5)P, phosphatidylinositol-5-phosphate; PI(3,4)P2, phosphatidylinositol 3,4-bisphosphate; PI(3,5)P2, phosphatidylinositol 3,5-bisphosphate; PI(4,5)P2, phosphatidylinositol 4,5-bisphosphate; PI(3,4,5)P3, phosphatidylinositol 3,4,5-trisphosphate.

FIGURE 8. FGD2 activates Cdc42. A, the “specificity patch” within the DH domains of RhoGEFs specific for Cdc42 (ASEF, PEM2, intersectin, and FGD1,3,4) are aligned with the corresponding FGD2 sequence. The conserved leucine mediating effective binding to Cdc42 is shaded in gray. B, lysates from COS-7 cells transfected with Myc-Cdc42 and EGFP, EGFP-tagged FGD2, FGD2-DHPH domain (DHPH), or FGD2FYVEKT1 mutant (FYVEKT) were immunoprecipitated with ACK1-crib-GST beads. Immunoprecipitated beads (IP; Ack, upper panels) or whole cell lysates (WCL, lower panels) were electrophoresed and immunoblotted with anti-Myc (right panels) or anti-EGFP (left panels). Lysates from transfected COS-7 cells expressing dominant active (Cdc42DA) and negative (Cdc42DN) EGFP-tagged Cdc42 mutants were used as controls (left panels). This experiment was performed three times with reproducible results.

FIGURE 9. FGD2 enhances JNK1 activity via Cdc42 but not Rac1. A, COS-7 cells were transiently transfected with FLAG-JNK1 cDNA together with EGFP or EGFP-FGD2, in the presence of either empty vector (V), Myc-tagged Rac1 (R), or Myc-tagged Cdc42 (C). JNK1 was immunoprecipitated with anti-FLAG beads and analyzed for kinase activity by incubation with [32P]ATP and the JNK1 substrate ATF-2-GST. Eluates from the FLAG beads were electrophoresed and transferred to membranes; JNK1 activity was determined by the autoradiography of phosphorylated ATF-2. Western blotting from electrophoresed cell lysates confirmed equal expression of transfected JNK1 (lower panel). B, FGD2 requires Cdc42 and an active GEF domain but not an active FYVE domain to stimulate JNK1 activity. JNK1 kinase assays were performed as in A, but COS-7 cells were transfected with Myc-Cdc42 (left panels) or Myc-Rac1 (right panels) and either EGFP alone, EGFP-tagged FGD2, EGFP-tagged FGD2DA mutant, or EGFP-tagged FGD2FYVEKT mutant. EGFP-tagged constitutively active mutants of Cdc42 (lane 5) or Rac1 (lane 10) were used as positive controls. The expression of the various components was confirmed by Western blotting with antibodies against either Myc or EGFP.
The FGD2-restricted expression pattern in antigen-presenting cells implies that it has a unique role in controlling the biology of these cells. This is underscored by the finding that FGD2 expression is tightly regulated at least during B lymphocyte development. Potential cellular functions that might be modulated by FGD2 include antigen uptake via endocytosis, antigen presentation via regulation of vesicle trafficking, and cell migration via cytoskeletal rearrangements. In accordance with these potential FGD2 functions, we demonstrate that FGD2 is partly localized to early endosomes, through FYVE domain-mediated binding to phospholipids. To our knowledge, FGD2 is the only known RhoGEF family member shown to have a functional FYVE domain and endosomal binding activity. Although both FGD1 and FGD3 have a FYVE domain, sequence analysis shows that FGD1 and FGD3 have species-conserved alterations of the critical tryptophan residue in the WXXX motif of the FYVE domain (see supplemental Fig. 2). Mutation of this residue in EEA1 abrogates endosomal localization, indicating that both FGD1 and FGD3 may have reduced affinity for phosphatidylinositol 3-phosphate and endosomes (43). FGD4, or Frabin, appears to have a conventional FYVE domain motif, but endosomal localization of this protein has not been described, perhaps due to the presence of an F-actin binding domain that may predominantly direct the protein to the actin cytoskeleton (44).

Although the Rab and Arf families of GTPases are best known for their roles in vesicle budding, docking, fusion, and movement within the mammalian endocytic and secretory pathways (45–47), functions for the Rho family of GTPases in vesicle pathways are being elucidated (48). A role for Cdc42 in intracellular trafficking in yeast has been identified, where Cdc42 is involved in vacuolar membrane docking and fusion (49, 50). Recent studies have also demonstrated that an effector of Cdc42, ACK1, is intimately involved in regulating the endocytosis and degradation of the EGFR (51). In addition, in maturing bone marrow-derived dendritic antigen-presenting cells, which express FGD2, the endocytic capacity correlates with the level of active Cdc42 (52). In this system, as dendritic cells mature, the ability to uptake further antigens by endocytosis is blocked, and mature dendritic cells have reduced levels of active Cdc42 when compared with immature dendritic cells. The intracellular localization of FGD2 at endocytic vesicle membranes would suggest that FGD2 is in a unique position to regulate Cdc42 and possible vesicle formation and trafficking processes in dendritic or other professional antigen-presenting cells.

We also found a portion of FGD2 localizing to membrane ruffles. In contrast to FGD1, which associates with cortical actin via cortactin binding by its Src homology 3 binding domain, we show that FGD2 is recruited to membrane ruffles through binding of phosphoinositides by its C-terminal PH domain. The presence of FGD2 in cortactin-rich membrane ruffles may relate to a role of this protein in the regulation of cell polarization and migration, since membrane ruffles often form at the leading edge of migrating cells. Another possibility is that FGD2 controls macropinocytic endocytosis in antigen-presenting cells. Membrane ruffles are essential parts of the macropinocytic machinery, which controls high volume nonspecific endocytosis. Blockade of Rho family GT Pases by toxin B completely inhibits macropinocytosis of immature spleen or bone marrow dendritic cells, as does injection of dominant negative Cdc42 or Rac1 (53). Presumably, FGD2 would exert control over this endocytic pathway via Cdc42, rather than Rac1, since we have not been able to detect direct activation of Rac1 by FGD2. Membrane ruffling is classically viewed as controlled by Rac-specific RhoGEFs. Constitutive ruffling by dendritic cells, which reflects macropinocytosis, appears to be different, however, since previous studies have shown that it is independent of Rac (54).

The recruitment of FGD2 to both endosomal membranes and the actin-rich lamellipodia is driven by two competing signals. In the absence of the C-terminal PH domain, early endosomal recruitment of FGD2APH2 appears to be enhanced, with a corresponding loss of membrane ruffle localization. In contrast, the FYVE domain mutant FGD2FYVEKT lacked endosomal localization but appeared to have augmented recruitment to the cortactin-rich areas of the cell. We conclude that wild type FGD2 is receptive to both signals and can be recruited to either compartment, depending on the levels of specific phospholipids and phosphatidylinositol 3-kinase activity within cellular compartments.

Our biochemical analysis of FGD2 predicts a specific role of FGD2 in the activation of Cdc42 and not of Rac1. Based on studies of the RhoGEF’s DBs and intersectin, the structure of FGD2 (and indeed all Fgd family members) predicts a role as a Cdc42 activator and seems inconsistent with Rac activation (4). All Fgd members have a conserved leucine residue in the DH domain, which, when changed to an isoleucine in intersectin, leads to a specificity change from Cdc42 to Rac1 (4). Overexpression of FGD2 led to elevated levels of active GTP-bound Cdc42, and FGD2 activated JNK1 activity when expressed with Cdc42, but not Rac1, consistent with Cdc42 specificity. FGD1, FGD3, and FGD4 (Frabin) are known to activate Cdc42 directly (17–19), we presume the same to be true for FGD2, although so far, we have been unable to generate a soluble recombinant FGD2 that contains any exchange activity in vitro.

A recent publication reports that in certain wild strains of mice, overexpression of Fgd2 RNA is associated with an altered allelic map to the T complex on chromosome 17, leading to altered inheritance of the T complex (22), a phenomenon called t-complex ratio distortion (55). Although normally expressed at very low levels in testes, this study shows that an altered form of Fgd2 RNA is overexpressed in T complex-containing sperm, which in conjunction with other distortion genes can influence the fertilization and thus transmission of the T complex genetic region. The authors propose that FGD2 is acting upstream in the activation of Smok (sperm motility kinase) (22). Cdc42 is expressed in sperm cells, along with other Rho GTPases (56); however it is not clear if the FGD2-mediated distortion effects on sperm fertility in this model are dependent upon Cdc42 function or through a so far uncharacterized regulation of the Smok kinase pathway. Since the Smok kinase described in mice is not conserved across species, and Fgd2 is highly conserved in mammals, we predict that FGD2 will play roles in other Cdc42-dependent signaling pathways. Future studies of the immune
response of FGD2-deficient animals may help elucidate this novel protein’s function in vivo.

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