BRIEF DEFINITIVE REPORT

Cell migration plays a critical role in various processes of acquired immunity, such as egress of naive T cells from the thymus and their homing to secondary lymphoid organs (1). Such trafficking of immune cells is controlled by their directed migration toward chemotactic signals. Actin cytoskeleton undergoes continuous remodeling and serves as machinery for cell migration. The mDia family of formins and the Wiskott-Aldrich syndrome protein (WASP)–Arp2/3 system are two major actin nucleating–polymerizing systems in mammalian cells, with the former producing long straight actin filaments and the latter producing branched actin meshwork. Although much is known about the latter, the physiological functions of mDia proteins are unclear. We generated mice deficient in one mDia isoform, mDia1. Although mDia1−/− mice were born and developed without apparent abnormality, mDia1−/− T lymphocytes exhibited impaired trafficking to secondary lymphoid organs in vivo and showed reduced chemotaxis, little actin filament formation, and impaired polarity in response to chemotactic stimuli in vitro. Similarly, mDia1−/− thymocytes showed reduced chemotaxis and impaired egression from the thymus. These results suggest that mDia1 plays a distinct role in chemotaxis in T lymphocyte trafficking.

Impaired T lymphocyte trafficking in mice deficient in an actin-nucleating protein, mDia1

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Trafficking of immune cells is controlled by directed migration of relevant cells toward chemotactic signals. Actin cytoskeleton undergoes continuous remodeling and serves as machinery for cell migration. The mDia family of formins and the Wiskott-Aldrich syndrome protein (WASP)–Arp2/3 system are two major actin nucleating–polymerizing systems in mammalian cells, with the former producing long straight actin filaments and the latter producing branched actin meshwork. Although much is known about the latter, the physiological functions of mDia proteins are unclear. We generated mice deficient in one mDia isoform, mDia1. Although mDia1−/− mice were born and developed without apparent abnormality, mDia1−/− T lymphocytes exhibited impaired trafficking to secondary lymphoid organs in vivo and showed reduced chemotaxis, little actin filament formation, and impaired polarity in response to chemotactic stimuli in vitro. Similarly, mDia1−/− thymocytes showed reduced chemotaxis and impaired egression from the thymus. These results suggest that mDia1 plays a distinct role in chemotaxis in T lymphocyte trafficking.

The WASP–Arp2/3 system is under regulation by Cdc42 and Rac, two members of the Rho family of small GTPases, and actin meshworks induced by the WASP–Arp2/3 complex serve as an underlying structure for lamellipodia and filopodia, the structures that are induced by Rac and Cdc42, respectively (6), and promote directed cell migration (2, 5). The role of the WASP–Arp2/3 system in cell migration has been verified by the phenotypes of WAS patients with mutations in WASP and WASP-deficient mice, which showed defects in proliferation, differentiation, and migration of cells of the hematopoietic lineage (7, 8). In contrast, the mDia proteins were originally identified as Rho effectors (9). Straight actin filaments induced by mDia proteins are therefore used, for example, for stress fiber assembly (10, 11), a process regulated by Rac and Cdc42, respectively (6), and promote directed cell migration (2, 5). The role of the WASP–Arp2/3 system in cell migration has been verified by the phenotypes of WAS patients with mutations in WASP and WASP-deficient mice, which showed defects in proliferation, differentiation, and migration of cells of the hematopoietic lineage (7, 8). In contrast, the mDia proteins were originally identified as Rho effectors (9). Straight actin filaments induced by mDia proteins are therefore used, for example, for stress fiber assembly (10, 11), a process regulated by Rac and Cdc42, respectively (6), and promote directed cell migration (2, 5).
physiological roles in vivo. To address these issues, we generated mice deficient in one of the mDia proteins, mDia1 (9).

RESULTS AND DISCUSSION
We generated an mDia1-fl oxed strain of mice that allows deletion of the translation initiation exon, Exon 1, of mDia1 on Cre/loxP-mediated recombination (Fig. S1, A and B, available at http://www.jem.org/cgi/content/full/jem.20062647/DC1). mDia1+/fl ox heterozygotes were then crossed with EHa-Cre mice, in which Cre recombinase was expressed in the early embryo (12), to produce heterozygous mDia1+/− mice, which were intercrossed to produce homozygous mDia1−/− mice (Fig. S1 C). No expression of mDia1 protein was found in mDia1−/− mice, whereas expression of mDia2 and mDia3 proteins was not altered (Fig. 1 A). mDia1−/− mice were born with an expected Mendelian ratio (Fig. S1 D). Both male and female mDia1−/− mice developed without apparent abnormality and were fertile. Generated mDia1−/− mice were backcrossed for more than five generations to a C57BL/6 background and used for analysis with wild-type littermates.

Figure 1. Decreased number of T cells in secondary lymphoid organs of mDia1−/− mice. (A) Western blot analysis. The homogenates of the brain, lung, and thymus of wild-type (+/+), heterozygous (+/−), and homozygous (−/−) mice were subjected to immunoblot analysis for mDia1, mDia2, mDia3, and β-actin. (B) Weight of the body and lymphoid organs of wild-type (shaded bars) and mDia1−/− (open bars) mice (n = 3 for each group). The experiment was performed twice with reproducible results. (C) Immunohistochemistry of the spleen and lymph node. The spleen and lymph node of mDia1+/+ and mDia1−/− mice were stained for Thy1.2 (green) and B220 (red). Representative observation from samples of two mice. Bar, 100 μm. (D) Cell population analysis in the spleen, lymph node, and blood. The numbers of total cells, Thy1.2+ T cells, B220+ B cells, CD4+ T cells, CD8+ T cells, CD11c+ cells, and CD11b+ cells in the spleen, lymph node, and blood were determined in mDia1+/+ (shaded bars) and mDia1−/− (open bars) mice. (E) Cell population analysis in the thymus. The numbers of total cells and indicated subsets of thymocytes (DN, CD4−/CD8−; DP, CD4+/CD8−; CD4−SP, CD4+/CD8−; CD8−SP, CD4−/CD8+) in the thymus were determined in mDia1+/+ (shaded bars) and mDia1−/− (open bars) mice. The bottom panel shows the numbers of CD69+/CD62L+CD4−SP and CD69+/CD62L+CD8 SP cells in the thymus of mDia1+/+ and mDia1−/− mice (n = 4 for each group). Unless otherwise stated, experiments shown in D and E were performed using three mice for each group more than twice with similar results, and results from one experiment are shown. All data are shown as means ± SEM. *, P < 0.05 versus the number of the corresponding population in mDia1+/+ wild-type mice. The axillary and inguinal lymph nodes were used as peripheral lymph nodes for analysis in these experiments.
obtained from the same heterozygous mating as a control. 8–12-wk-old mice were used in subsequent analyses.

Systemic histological analysis of 8–9-wk-old mDia1−/− male and female mice (n = 3 each) using hematoxylin–eosin staining detected no apparent abnormality in various tissues (unpublished data). However, the wet weight of the spleen and peripheral lymph nodes (axillary and inguinal lymph nodes) tended to be lighter in mDia1−/− mice, whereas that of the thymus appeared no different between the genotypes (Fig. 1 B). Immunostaining for T cells and B cells in the spleen and lymph nodes revealed normal segregation of T cells and B cells but less density of T cells in both organs of mDia1−/− mice (Fig. 1 C). Consistently, the numbers of both CD4 and CD8 T cells were significantly reduced in the spleen and lymph nodes of mDia1−/− mice, whereas those of B cells, CD11c+ cells, and CD11b+ cells were not altered (Fig. 1 D). Notably, a significant reduction (P < 0.05) in the T cell numbers was also seen in the blood. In the thymus, although the numbers of the CD4−CD8−, CD4+CD8+, CD4−CD8+, and CD4−CD8+ thymocyte populations were not significantly different between wild-type and mDia1−/− mice, the numbers of the CD4+CD8− and CD4−CD8+ thymocytes tended to be higher in mDia1−/− mice, and those of the CD69hiCD62Lhi CD4 or CD8 single-positive thymocytes significantly increased in mDia1−/− mice (Fig. 1 E), suggesting impaired egression of mature thymocytes from the thymus in mDia1−/− mice.

Given that trafficking of thymocytes and T cells is achieved by their migration to chemokines (1), we examined the response of these cells toward chemokines in vitro. The transwell assay revealed impaired chemotaxis of mDia1−/− T cells toward CXC chemokine ligand (CXCL) 12 and CC chemokine ligand (CCL) 21, the two chemokines implicated in in vivo trafficking to secondary lymphoid organs (13). On the other hand, no significant impairment was found in chemotaxis of mDia1−/− B cells toward CXCL12 (Fig. 2 A). Similarly, the transwell assay revealed impaired chemotaxis of CD4+CD8+, CD4+CD8−, and CD4−CD8+ thymocytes toward CXCL12 (14) and CCL21 in the mDia1−/− genotype (Fig. 2 B). To assess egression more directly, we used organ culture of the thymus. Egression of CD4+CD8− and CD4−CD8+ thymocytes from thymic lobes in response to CCL21 was significantly decreased in mDia1−/− mice (Fig. 2 C). These results indicate that impaired chemotaxis of thymocytes and T cells underlies the phenotype observed in mDia1−/− mice. However, T cell trafficking to lymphoid organs in vivo may not be determined solely by the migratory activity of T cells but can be influenced by other factors, including the adhesive

Figure 2. Impaired migration of mDia1−/− T cells and thymocytes. (A and B) Impaired chemotaxis toward chemokines in vitro. Migration of T cells and B cells [A; n = 4 for each group] and thymocytes [B; n = 3 for each group] from mDia1+/− (shaded bars) and mDia1−/− (open bars) mice toward the indicated chemokines was examined using a transwell chamber. In B, populations of thymocytes that migrated to the bottom chamber were analyzed by flow cytometry after staining for CD4 and CD8. *, P < 0.05. (C) Impaired thymocyte egression from the thymus in organ culture. Thymocytes that egressed from the thymic lobe toward CCL21 were stained for CD4 and CD8 and analyzed by flow cytometry (n = 3 for each group). *, P < 0.05. (D) Adoptive transfer experiment. T cells were isolated from the spleen of mDia1+/+ and mDia1−/− mice and were labeled with different fluorescent dyes, administered to wild-type C57BL/6 mice or mDia1−/− recipient mice (n = 3 for each group). The ratios of the two populations in the blood and migrating to the spleen and the axillary and inguinal lymph nodes were analyzed after 2 h. All data are shown as means ± SEM.
property of high endothelium venules in the lymphoid organs (15). To examine this issue, we performed an adoptive transfer experiment, in which we labeled T cells from wild-type and mDia1<sup>−/−</sup> mice with different fluorescent dyes and infused them i.v. into wild-type or mDia1<sup>−/−</sup> recipients. The numbers of transferred cells that migrated into the spleen and lymph nodes were determined 2 h after the infusion. We observed that, compared with wild-type cells, only ~50% of mDia1<sup>−/−</sup> T cells migrated into the spleen and lymph nodes and that the extent of the reduction was not different in wild-type and mDia1<sup>−/−</sup> recipients (Fig. 2 D). Consistently, the higher numbers of mDia1<sup>−/−</sup> cells remained in the circulation.

Given the function of mDia1 in actin nucleation and polymerization (3, 4), we next examined the cellular response of mDia1<sup>−/−</sup> T cells to chemokine stimulation. The cells were initially stained with fluorescent phalloidin and subjected to flow cytometry, which revealed that the addition of CCL21 to wild-type T cells induced robust filamentous actin (F-actin) production in a time- and concentration-dependent manner (Fig. 3 A). In contrast, T cells from mDia1<sup>−/−</sup> mice exhibited a substantially attenuated response at all time points measured, and the extent of attenuation was more marked at the lower chemokine concentration. Then the cells were stained for F-actin and talin. Microscopic examination of these cells revealed that wild-type T cells exhibited not only robust F-actin formation but also cell polarization, and that both F-actin production and cell polarization were markedly suppressed in mDia1<sup>−/−</sup> T cells (Fig. 3 B). On the other hand, the addition of CXCL13 induced both actin filament production and polarization in mDia1<sup>−/−</sup> B cells (Fig. S2, available at http://www.jem.org/cgi/content/full/jem.20062647/DC1).

Because actin polymerization is also important in T cell proliferation to TCR stimulation, we next examined whether mDia1 is involved in this process. Compared with wild-type cells, mDia1<sup>−/−</sup> T cells exhibited impaired proliferation to stimulation with both anti-CD3 antibody (Fig. 4 A) and mixed lymphocyte reaction (Fig. 4 B). Thus, the loss of mDia1 affected not only T cell migration but also proliferation. Given that these processes are crucial in immune response, we suspected that an in vivo immune response was also impaired in mDia1<sup>−/−</sup> mice. To test this issue, we subjected mDia1<sup>−/−</sup> mice to the dinitrofluorobenzene (DNFB)-induced contact hypersensitivity, a model for a T cell–dependent immune response. Consistent with our hypothesis, the immune response examined as the hapten–induced ear thickness was significantly impaired in mDia1<sup>−/−</sup> mice (Fig. 4 C).

These results demonstrated that the deficiency of mDia1 significantly suppressed (P < 0.05) the T cell responses, not only cell migration and polarization but also cell proliferation. Given the intricate interaction of many signaling molecules in these biological processes (6), we examined the expression and abundance of molecules other than mDia1 involved in T cell migration and proliferation. Flow cytometry revealed no change in the expression of the chemokine receptors CXC chemokine receptor (CXCR) 4 and CC chemokine receptor (CCR) 7 (1) in mDia1<sup>−/−</sup> T cells (Fig. 5 A).

Immunoblot analysis revealed that mDia1 was apparently a sole mDia isoform expressed in T cells and no compensatory expression of other isoforms in mDia1<sup>−/−</sup> T cells. The mDia1 deficiency neither induced alteration in the amount of molecules such as RhoA, Rac1, and Cdc42 and another group of Rho effectors, ROCK 1 and 2, nor in the amount of the Arp2/3 subunit 34. However, substantial suppression was found in the

Figure 3. Impaired chemokine-induced actin polymerization and polarization of mDia1<sup>−/−</sup> T cells. (A) F-actin production to chemokine stimulation. T cells from mDia1<sup>+/+</sup> (blue line) and mDia1<sup>−/−</sup> (red line) mice were stained with different concentrations of CCL21 for the indicated times, stained with Oregon green–phalloidin, and analyzed by flow cytometry. Mean fluorescence intensity of the entire cell population was determined for each group and is shown with that of unstimulated mDia1<sup>−/−</sup> T cells. T cells from mDia1<sup>−/−</sup> mice were stained with Oregon green–phalloidin, and analyzed by flow cytometry. Mean fluorescence intensity of the entire cell population was determined for each group and is shown with that of unstimulated mDia1<sup>−/−</sup> T cells. T cells from mDia1<sup>−/−</sup> mice were stained with Oregon green–phalloidin, and analyzed by flow cytometry. Mean fluorescence intensity of the entire cell population was determined for each group and is shown with that of unstimulated mDia1<sup>−/−</sup> T cells.
amount of WASP in mDia1−/− T cells (Fig. 5 B). Comparison with dilutions of wild-type cell lysates suggests that the WASP amount in mDia1−/− cells was ~25% of that in wild-type cells. In contrast to these findings, RT-PCR analysis revealed that similar amounts of WASP transcript were present in wild-type and mDia1−/− cells (Fig. S3, available at http://www.jem.org/cgi/content/full/jem.20062647/DC1). We therefore treated wild-type and mDia1−/− T cells with cycloheximide and compared the rate of disappearance of WASP in the two genotypes of cells by immunoblot analysis. We found that the amount of WASP decreased more quickly in mDia1−/− T cells than in wild-type cells with half-lives of ~6 and 11 h, respectively (Fig. 5 C). To examine a mechanism for accelerated WASP degradation, we next examined the effects of a proteasome inhibitor, MG132. The addition of MG132 concentration dependent restored the amount of WASP in mDia1−/− T cells, which was doubled by incubation with 20 μM MG132 for 4 h. On the contrary, MG132 had no effect on WASP amounts in wild-type cells (Fig. 5 D). Thus, WASP appears to be degraded via a ubiquitin–proteasome pathway in the absence of mDia. Because WASP is one of the components of another major actin-nucleating–polymerizing system and is involved in the migration and proliferation of T cells, these findings raised a question as to whether the phenotype we observed is caused by the lack of mDia1 or partial depletion of WASP. To address this issue, we overexpressed WASP in wild-type and mDia1−/− T cells as an enhanced GFP (EGFP)–fusion protein to 17 and 19% of the level of endogenous WASP in wild-type and mDia1−/− T cells, respectively. We also expressed EGFP as a control. Transfection efficiency was ~5% for EGFP–WASP and ~13% for EGFP, indicating that EGFP–WASP was present in transfected mDia1−/− cells at the level of endogenous WASP in control cells. Cells expressing EGFP or EGFP–WASP were then subjected to the migration assay and F-actin staining. Overexpression of EGFP–WASP neither improved the migratory activity nor rescued impaired polarization and F-actin increase of mDia1−/− cells (Fig. 5 E). These results suggest that a defect in the migration of mDia1−/− T cells is primarily caused by the loss of mDia1 itself.

In this report, we revealed impaired trafficking of T lymphocytes in mDia1−/− mice. The in vitro chemotaxis analysis and the adoptive transfer experiment suggest that the impairment is intrinsic to mDia1−/− T cell lineage. These findings, together with impaired egression of thymocytes from the thymus in organ culture, indicate that the reduction of T cell density in the secondary lymphoid organs of mDia1−/− mice is caused by impairment in both thymocyte egress from the thymus and T cell homing to the lymphoid tissues. Our analysis has further demonstrated impaired chemokine-induced F-actin formation and cell polarization in mDia1−/− cells as a mechanism underlying the phenotype of the mDia1 deficiency. Previously, we used siRNA for mDia1 and showed that depletion of mDia1 in rat C6 glioma cells impaired their migration and polarization in vitro (16). Our present findings are consistent with those findings and suggest that the principles we found in cultured cells indeed operate in vivo and govern the behavior of lymphocytes in the body. Notably, WASP−/− mice or WAS patients or cells derived from them exhibit phenotypes partially overlapping those of mDia1−/− mice, such as decreased cellularity in the lymphoid organs and defects in T cell proliferation and migration (17, 18). Collectively, these results suggest that the two actin-nucleating systems work together to achieve lymphocyte functions, and the lack of either one cannot be complemented by the other.

In this respect, it is intriguing that WASP is depleted in the absence of mDia1, because this finding indicates not only functional but also physical interdependency of the two major actin-nucleating systems in the cell. Our results indicate that the WASP depletion in mDia1−/− cells is caused by accelerated degradation via the ubiquitin–proteasome system. Previously, phosphorylation of neural WASP by the Src family kinases was found to render it susceptible to ubiquination and degradation (19). Furthermore, WAVE2, a WASP family protein, is stabilized by making a multiprotein complex, and depletion of any other component of that complex triggers...
Given that the mDia family of proteins play elemental roles in actin filament dynamics, it is rather surprising that mDia1−/− mice exhibited relatively mild phenotypes. This is partly because of the redundancy of the mDia isoforms (3), but may also be caused by the presence of other cell-migration machinery. The latter possibility is consistent with our results on B cells (Fig. 2A). mDia1 is the major, almost sole, mDia isoform in B cells (unpublished data), and there is a negligible amount of WASP in B cells. Yet, there is no B cell phenotype in mDia1−/− mice, and mDia1−/− B cells responded normally to chemotactic stimuli. In summary, our present report has unveiled unforeseen relationships between the two actin-nucleating systems that not only function cooperatively but also regulate each other’s stability in the cell.

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Figure 5. Analysis on signaling molecules. (A) Flow cytometry for chemokine receptor expression. mDia1+/+ and mDia1−/− T cells were stained with anti-CXCR4 or anti-CCR7 antibody (continuous line) or control antibody (dotted line) and analyzed by flow cytometry. (B) Western blot analysis of lysates of mDia1−/− T cells. T cells were prepared from mDia1+/+ and mDia1−/− mice and subjected to immunoblot analysis for mDia1, mDia2, mDia3, RhoA, Rac1, Cdc42, ROCK-1, ROCK-2, WASP, or Arp2/3 subunit 34. The amount of WASP in mDia1−/− T cells was examined by serial dilutions of the mDia1+/+ lysates. (C) Degradation of WASP. T cells were treated with 10 μg/ml cycloheximide (CHX) for the indicated times, and cell lysates were subjected to immunoblot analysis with antibody to WASP (top). For comparison, two-fold amounts of lysates and longer color development were used for analysis of mDia1−/− T cells. The density of each band on the immunoblot was quantified by the densitometry and plotted (mDia1+/+, blue line; mDia1−/−, pink line) against incubation time with the density at 0 h as 100% (right). (D) Effects of proteasome inhibitor. T cells were treated with the indicated concentrations of MG132 for 4 h, and cell lysates were subjected to immunoblot for WASP. (E) Overexpression of WASP in mDia1−/− T cells. Wild-type or mDia1−/− T cells were transfected with pEGFP-WASP or pEGFP. Expression of EGFP-WASP was confirmed by Western blotting and flow cytometry (top left and top middle). Densitometry of the immunoblot of the cells transfected with 4 μg pEGFP-WASP showed that EGFP-WASP was expressed at 17.4 and 18.8% of the endogenous WASP in wild-type and mDia1−/− cells, respectively. Wild-type and mDia1−/− T cells transfected with 4 μg of pEGFP or pEGFP-WASP were subjected to the transwell migration assay (top right) or fluorescence microscopy after staining with Texas red–phalloidin (bottom left), followed by measurement of the fluorescence intensity of F-actin staining (bottom right; n = 30 cells each). Bar, 10 μm.
demonstrated an essential role of mDia1 in T lymphocyte trafficking. This work should lead to a better understanding of not only immune trafficking and its defects in humans but also of various pathological processes involving cell migration.

**MATERIALS AND METHODS**

**Mice.** Mice were bred at the Institute of Laboratory Animals of Kyoto University on a 12-h light/dark cycle under specific pathogen-free conditions. All experimental procedures were approved by the Committee on Animal Research of Kyoto University Faculty of Medicine.

Cell sorting, chemotaxis, and proliferation. Dissociated T cells and B cell suspensions were prepared from the spleen using an autoMACS system (Miltenyi Biotec) and were cultured as described previously (24). Chemotaxis was assayed using a transwell, as previously described (24). In brief, $3 \times 10^5$ T cells, B cells, or thymocytes were used, and cells that had migrated to the lower chamber after 3 h of incubation at $37^\circ C$ were collected and counted by flow cytometry. For expression of EGFP-WASP, 2–4 $\mu g$ of pEGFP-WASP were transfected into T cells from wild-type or mDia1 $^{-/-}$ mice using the Nucleofector system (Amaxa) with an efficiency of $\sim$5% and incubated for 18 h at $37^\circ C$. The cells were subjected to Western blotting and chemotaxis assay. For thymocyte egression, a thymic lobe from new-born mice was transferred to the top chamber of a transwell, and 600 $\mu l$ of the medium containing 100 nM CCL21 was added to the bottom chamber. After 24 h of incubation at $37^\circ C$, cells that had egressed from the lobe and migrated to the bottom chamber were collected and analyzed by flow cytometry. For T cell proliferation, T cells were isolated from the spleen of mDia1 $^{-/-}$ and mDia1 $^{+/+}$ mice, and $2 \times 10^5$ cells were cultured for 48 h with plate-bound anti-CD3 antibody, or $2.5 \times 10^5$ cells were cultured with CD11c $^+$ dendritic cells from BALB/c mice for 72 h. [3H]Thymidine uptake was measured as previously described (24).

Adaptive transfer. T cells were isolated from the spleen of mDia1 $^{-/-}$ or mDia1 $^{+/+}$ mice and were labeled differently with either 2 $\mu M$ PKH26 red fluorescent cell linker (Sigma-Aldrich) or 1 $\mu M$ 5,6-carboxyfluorescein diacetate (Invitrogen) (25). After washing, mDia1 $^{-/-}$ and mDia1 $^{+/+}$ T cells were mixed in equal numbers in PBS, and $5 \times 10^5$ cells were i.v. injected into recipient C57BL/6 or mDia1 $^{-/-}$ mice. After 2 h, the blood, spleen, and lymph nodes were removed from the recipient mice. A cell suspension was prepared, and the numbers of cells labeled with each dye were analyzed by flow cytometry.

Western blotting. The brain, lung, and thymus were isolated from adult mice, and T cells were prepared from the spleen. These organs and cells were lysed in RIPA buffer (20 mM Tris-HCl, pH 7.4, 150 mM NaCl, 2 mM EDTA, 1% Nonidet P-40, 1% Na deoxycholate, 0.1% SDS) containing a mixture of protease inhibitors (10 $\mu g$/ml aproten, 10 $\mu g$/ml leupeptin, 1 mM PMSF) and 5 mM 2-mercaptoethanol. The lysates were boiled and subjected to SDS-PAGE, and separated proteins were transferred to nitrocellulose membranes. The membranes were probed with antibodies to mDia1 (Bethyl Laboratories), mDia3, RhoA, ROCK-1, WASP (Santa Cruz Biotechnology, Inc.), ROCK-2, Rac1, Cdc42 (Transduction Laboratories), Arp2/3 subunit 34 (Imgenex), $\beta$-actin (Sigma-Aldrich), GFP (MBL International Corporation), or mDia2. Antibody to mDia2 (provided by Y. Ando, Kyoto University Faculty of Medicine, Kyoto, Japan) was raised in rabbits against a recombinant protein corresponding to Rho-binding domain of mDia2 expressed in Escherichia coli as a His-tagged protein. To examine the stability of WASP, T cells were incubated with 10 $\mu g$/ml cycloheximide for 0, 1, 2, 4, 8, 12, and 16 h. The cells were subjected to Western blotting for WASP. The effects of MG132 were examined by incubating T cells with 0, 1, 5, and 20 $\mu M$ MG-132 (Peptide Institute) for 4 h.

Fluorescence microscopy. Cryostat sections were prepared from the spleen and lymph node, fixed in cold acetone for 10 min, and stained with FITC- or PE-labeled anti-Thy1.2 or anti-B220 monoclonal antibodies. For chemokine responses, cells were cultured in the culture medium at $37^\circ C$ overnight on poly-L-lysine-coated coverslips, and the medium was changed with the fresh medium with or without 100 ng/ml CCL21 (R&D Systems) for T cells or 500 ng/ml CXCL13 (R&D Systems) for B cells. After 10 min, the cells were fixed with 4% paraformaldehyde in PBS, permeabilized with 0.1% Triton X-100, and stained with antibody to talin, Texas red-phalloidin, and TOPRO3. Talin was visualized by Alexa Fluor 488-labeled anti-mouse IgG monoclonal antibody. Samples were analyzed on a confocal laser scanning microscope (LSM510; Carl Zeiss MicroImaging, Inc.). Fluorescence intensity of F-actin staining with Texas red-phalloidin was determined in each cell by using MetaMorph (Universal Imaging Corp.), as previously described (16).

Flow cytometric analysis. Cell suspensions were prepared from the thymus, spleen, and lymph node and were incubated with FITC- or PE-labeled monoclonal antibody to CD4, CD8, Thy1.2, B220, CD11c, CD11b, CD62L, CD69, CXCR4, or CCR7 (29). For analysis of the CD69$^+$ CD62L$^+$ thymocyte subset, Cy5- or APC-labeled monoclonal antibody to CD4 or CD8 was used. Labeled cells were then subjected to flow cytometry. For analysis of F-actin production, 10$^6$ T cells were stimulated with CCL21 for various times and fixed with 4% paraformaldehyde in PBS, treated with 0.1% saponin, and stained with Oregon green–phalloidin (26).

Contact hypersensitivity. Mice were immunized by application of 25 $\mu l$ of 0.5% DNFB in 1:1 (vol/vol) acetone/olive oil to their shaved abdomens on day 0 and were challenged on day 5 by painting the ear with 20 $\mu l$ of 0.3% (wt/vol) DNFB. Ear thickness was measured $24$ h after challenge.

Statistical analysis. All data are shown as means ± SEM. Comparisons of two groups were analyzed using the unpaired two-tailed Student’s t test. P $<$ 0.05 was considered statistically significant. The analyses were performed by Prism software (version 4; GraphPad).

Online supplemental material. Materials and methods describe the generation and breeding of mDia1-deficient mice and the RT-PCR analysis of WASP mRNA. Fig. S1 shows a diagram of the loxP-frt/Neo cassette used in the targeting vector, schematic representation of the wild-type allele, the targeting vector, the targeted allele and the knockout allele for mDia1, and a PCR genotyping and ratio of wild-type, heterozygous, and homozygous littermates. Fig. S2 shows chemokine responses of mDia1 $^{-/-}$ T cells examined by F-actin staining. Fig. S3 shows RT-PCR and real-time PCR of WASP mRNA in mDia1 $^{-/-}$ and mDia1 $^{+/+}$ T cells. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20086264/DC1.

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REFEREENCES

Generation and breeding of mDia1-deficient mice. The mDia1-floxed mutant (available from the eXintegrator system [http://www.cdb.riken.go.jp/ scb/MartinG/html/exintegrator.html] under accession no. CD80437K) was established as follows. A genomic clone containing the mouse mDia1 gene was isolated from the RP23-407E2 mouse female (C57BL/6J) bacterial artificial chromosome library (BACPAC Resources Center). The loxP-frt/Neo cassette in pBSK (Fig. S1 A) was generated by modification of the frt/pgk-Neo-pA/frt in Bluescript (1) and the loxP/PGK-Neo-pA/loxP, pMCDT-A cassette (available at http://www.cdb.riken.go.jp/arg/cassette.html) (2), a gift from Shin-ichi Aizawa (RIKEN Kobe Institute, Kobe, Japan). A 4-kbp 5′ fragment (chromosome 18, nt 35066636–35062572) to exon 1 was inserted into multiple cloning site (MCS) 1 in the above cassette as the short arm. A 3.1-kbp fragment containing exon 1 (chromosome 18, nt 35062571–35059388) was ligated in a Hind3 site. A 5.8-kbp 3′ fragment to exon 1 (chromosome 18, nt 35059387–35053557) was inserted into MCS2 as the long arm. The targeting vector was linearized with Asc1 and introduced into TT2 embryonic stem cells by electroporation. Cell clones resistant to G418 were isolated, and homologous recombination was examined with PCR and Southern blotting (3). The positive embryonic stem cell clones were microinjected into eight cell-stage embryos from ICR strain mice, and chimeric offspring were generated (2). Chimeras were bred with C57BL/6N female mice twice, and resultant heterozygotes were crossed with EIIa-Cre mice (The Jackson Laboratory) to produce heterozygous mutants (mDia1+/−) lacking exon 1 and the neomycin gene. mDia1+/− heterozygous mice lacking the Cre gene were obtained by mating with C57BL/6N mice again and were intercrossed to produce mDia1+/− homozygotes. Genotyping analysis was performed by PCR using the genomic DNA from mice tails. PCR primer pairs were F1 (5′-CTCCAGTGCGGATGTGAACTGTG-3′) and R1 (5′-GTTCAGCTCCACCCTCTGTACAGC-3′) for examining the exon 1 deletion, or Cre1 (5′-GGGCTGGGATGTCACAGCTG-3′) and Cre2 (5′-GTGAAACAGCATTGCTGTCACTT-3′) for detection of the Cre gene. Loss of mDia1 protein was examined by Western blotting. Systemic histological analysis was performed on 8–9-wk-old wild-type and mDia1−/− male and female mice (n = 3 each) using hematoxylin-eosin staining. Organs examined included the brain, pituitary, liver, pancreas, kidney, spleen, thymus, heart, lung, adrenal gland, the gastrointestinal tract, mesenteric lymph node, testis, bladder, skeletal muscle, blood vessel, and bone (sternum and femur).

RT-PCR analysis of WASP mRNA. Total RNA was isolated by using an RNeasy mini kit (QIAGEN), according to the manufacturer’s instructions. RT-PCR was performed on 500 ng of total RNA using SuperScript one-step RT-PCR with Platinum Taq (Invitrogen) with a set of WASP primers (5′-AGAG-GAGGGCTCCCCACCTGTG-3′ and 5′-GTTGCTCCCGATATCAGCTTGG-3′, yielding a 211-bp amplified fragment) and, as a positive control, a set of hypoxanthine-guanine phosphoribosyltransferase primers (5′-GCTGGTGAAAAGGACCTCT-3′ and 5′-CACAGACTAGAACACCTGC-3′, yielding a 250-bp amplified fragment). cDNA synthesis and prenaturation conditions were as follows: 30 min at 50°C and 2 min at 94°C. PCR amplification conditions were as follows: 15 s at 94°C, 30 s at 57°C, and 1 min at 72°C for 35 cycles; and 10 min at 72°C as a final extension. For real-time PCR, complementary DNA was synthesized using SuperScript (Invitrogen). The amount of mRNA for WASP and GAPDH was quantified by a real-time PCR system (LightCycler 2.0; Roche Diagnostics). The primer sequences used were as follows: 5′-CACCACACAGCACCAAAATCA-3′ and 5′-CTTAGAGATAGTGGACACACTGGA-3′ as the forward and reverse primers, respectively, for WASP, and 5′-TGACGGGAAGCTCAGCTGGG-3′ and 5′-TCCACCCACGTGGTGAAC-3′ as the forward and reverse primers, respectively, for GAPDH. Data were analyzed by LightCycler Probe Design software (version 4.0; Roche Diagnostics).

REFERENCES