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Orai1 Function Is Essential for T Cell Homing to Lymph Nodes

Milton L. Greenberg,⁎†,1 Ying Yu,⁎†,1,2 Sabrina Leverrier,⁎†,3 Shenyuan L. Zhang,⁎†,3 Ian Parker,⁎† and Michael D. Cahalan⁎†

In T lymphocytes, Ca2+ release–activated Ca2+ (CRAC) channels composed of Orai1 subunits trigger Ag-induced gene expression and cell proliferation through the NFAT pathway. We evaluated the requirement of CRAC channel function for lymphocyte homing using expression of a dominant-negative Orai1-E106A mutant to suppress Ca2+ signaling. To investigate homing and motility of human lymphocytes in immunocompromised mouse hosts, we transferred human lymphocytes either acutely or after stable engraftment after a second transfer from the same blood donor. Human and mouse lymphocyte homing was assessed, and cells were tracked within lymph nodes (LNs) by two-photon microscopy. Our results demonstrate that human T and B lymphocytes home into and migrate within the LNs of immunocompromised NOD-SCID mice similar to murine lymphocytes. Human T and B cells colocalized in atrophied or reconstituted mouse LNs, where T cells migrated in a random walk at velocities of 9–13 μm/min. Expression of Orai1-E106A inhibited CRAC channel function in human and mouse T cells, and prevented homing from high endothelial venules into murine LNs. Ca2+ signals induced by CCL21 were also inhibited in T cells expressing Orai1-E106A. With CRAC channels inhibited, the high-affinity form of LFA-1 failed to become active, and T cells failed to migrate across endothelial cells in a transwell model. These results establish a requirement for CRAC channel–mediated Ca2+ influx for T cell homing to LNs mediated by high-affinity integrin activation and chemokine-induced transendothelial migration. The Journal of Immunology, 2013, 190: 000–000.

Sustained Ca2+ influx through Ca2+ release–activated Ca2+ (CRAC) channels is essential for Ag-dependent T cell activation and proliferation via NFAT-driven gene expression (1, 2). In addition, after TCR engagement, CRAC channel activation elevates cytosolic calcium ([Ca2+]i) and acts to halt T cell migration and facilitate prolonged interactions with APCs (3). RNA interference screening and mutagenesis have identified two proteins that activate and form the CRAC channel: STIM1 proteins (STIM1 and STIM2 in mammals) as the endoplasmic reticulum (ER)–resident Ca2+ sensor; and Orai proteins (Orai1, Orai2, and Orai3 in humans) as the Ca2+-selective, pore-forming subunit (4–9). In T cells and other cells of the immune system, Ca2+ store depletion induces STIM1 to migrate as oligomers from the bulk ER to junctions immediately adjacent to the plasma membrane and mediates Ca2+ influx by opening Orai channels directly (10). Although the properties of CRAC channels and their role in T cell Ca2+ signaling have been described in detail (11, 12), the functional roles of CRAC channels in human T cell activation and immune defense remain incompletely characterized. In particular, CRAC channels may also be required for lymphocyte trafficking and migration. To address this, we developed methods to engraft and visualize human lymphocytes in the context of immunocompromised mouse models, and expressed a dominant-inhibitory Orai1 construct to evaluate the functional role of CRAC channel activity in lymphocyte homing to lymph nodes (LNs) in vivo.

Several mutations in Orai1 or STIM1 have been identified in human patients with SCID, leading to inhibition of CRAC channel function and lethal immune suppression (7, 13–16). In patients with the homozygous Orai1-R91W mutation, CRAC channels fail to open, and T cells cannot take up Ca2+ after stimulation (7). In contrast, Ca2+ influx in T cells isolated from Orai1-deficient mice was only partially inhibited (17, 18), perhaps because of compensation by other Orai isoforms. We therefore sought an alternative method to evaluate CRAC channel function. A charge-neutralizing mutation of the critical glutamate residue to alanine in Drosophila Orai or human Orai1 has a dominant-negative effect on the CRAC channel, resulting in a nonconducting channel (4, 5). In this study, we use expression of the dominant-negative Orai1-E106A to silence the CRAC channel and inhibit Ca2+ influx to evaluate the functional role for Orai1 in homing.

T cell homing from the blood into secondary lymphoid organs (SLO) occurs by migration across the high endothelial venules.

Abbreviations used in this article: B2, β2-microglobulin; [Ca2+]i, intracellular calcium concentration; CF, cyan fluorescent protein; CMF, EC, endothelial cell; eGFP, enhanced GFP; ER, endoplasmic reticulum; HEV, high endothelial venule; LN, lymph node; SLO, secondary lymphoid organ; Tg, thapsigargin.

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(HEVs) and is required for adaptive immunity. This process is sequentially mediated by selectins, including CD62L, that facilitate tethering, cellular adhesion, and rolling within the HEV lumen (19), and chemokines and chemokine receptors that activate integrins and direct migration from the venules (20, 21). The chemokine receptor CCRI7 and its ligands CCL19 and CCL21 play a significant role in T cell homing and facilitate activation of the β2-integrin complex LFA-1, which allows interactions with ICAM-1 on the surface of the endothelium (22, 23). Elevations in [Ca2+]i have been shown to activate LFA-1 and induce lateral clustering on the T cell surface (24). CCL21 treatment induces Ca2+ influx in human T cells (25, 26), but it is unknown whether CCL21-induced Ca2+ signaling through Orai1 channels facilitates integrin activation and homing into LN.

Two-photon microscopy enables real-time visualization of immune cell migration and dynamics within the SLO of mice (27). However, it is not feasible to directly manipulate and image human immune cells in a native in vivo environment. Therefore, we sought an in vivo system using an immunocompromised murine host to visualize human T cell motility. NOD.SCID mice are characterized by defects in innate and adaptive immunity, including atrophied LNs devoid of T and B lymphocytes and reduced NK cell activity (28). Several NOD.SCID strains have been validated as recipients for reconstitution with human hematopoietic cells and are useful for establishing durable human xenografts (29–31). In this study, we first established human lymphocyte xenografted models for two-photon imaging of human cells in an in vivo environment, and characterized the motility of human T and B lymphocytes in comparison with murine cells in the immunocompromised murine host LNs. We used both the original NOD.SCID strain and NOD.SCID.B2 mice with B2-microglobulin (B2) knocked out. The latter strain lacks lymphoid and NK cells, and has been shown to permit enhanced engraftment of human T cells (32). We then used this model system in conjunction with expression of the dominant-negative Orai1-E106A mutant to investigate the role of Ca2+ influx through CRAC channels for integrin activation, chemokine responses, and homing of human and murine T cells.

**Materials and Methods**

**Mice**

NOD.B6.129-Pkdcdexnull/J (NOD.SCID) mice obtained from Jackson Laboratory (stock no. 001303; Bar Harbor, ME) were housed and monitored in a specific pathogen-free environment with sterile food and water in our animal facility. To inhibit NK cell activity, we injected NOD.SCID mice i.v. with 20 μM anti-NK cell Ab (rabbit anti-Asialo GM1; Wako Bioproducts, Richmond, VA) according to manufacturer’s instructions 3–4 d before adoptive transfer of human T cells. NOD.Cg-Pkdcdexnull B6.129-CreIscSeJ mice (NOD.SCID.B2) and C57.BL/6J mice were obtained from Jackson Laboratory (stock no. 002570, 000664). Mice used were between 8 and 18 wk of age. NOD.SCID.B2 mice were reconstituted with human peripheral blood leukocytes (PBLs) as described previously (30). A total of 3 × 107 human PBLs were injected i.p., and experiments were performed 3 wk later. All of the experimental procedures were approved by the Animal Care and Use Committee of University of California, Irvine.

**Purification of human monocytes and isolation of CD4+, CD8+, CD19+, and CD3+ cells**

Human PBLs were isolated from blood of voluntary healthy donors by Ficoll-Hypaque (gradient = 1.077 g/dl) density gradient centrifugation. Isolation of human CD4+ and CD8+ T cells was performed using human CD4+ and CD8+ T cell isolation kits (Miltenyi Biotec, Bergish Gladbach, Germany) according to manufacturer’s instructions. PBLs were first suspended in labeling buffer and incubated with anti-CD9 mAb-coated microbeads to prepare human CD8+ T cells. CD8+ T cells were isolated by positive selection using a magnetic cell separation system (MACS; Miltenyi Biotec). To prepare human CD4+ T cells, we isolated the remaining cells by negative selection with a CD4+ T cell isolation mixture. CD19+ B cells were isolated by positive selection according to manufacturer’s instructions (Miltenyi Biotec). In some experiments, human CD3+ T cells were isolated using EasySep Human CD3+ Isolation kit (Stem Cell Technologies, Vancouver, BC, Canada), according to manufacturer’s instructions. The purity of CD4+, CD8+, CD19+, or CD3+ cell populations was confirmed to be >95% by flow cytometry.

**Cell adoptive transfer in mice**

For human T cells, purified CD4+ and CD8+ T cells and CD19+ B cells were separately labeled with CellTracker dyes (Invitrogen, Grand Island, NY) CFSE (4 μM), 5-(and-6)-[(4-chloromethyl)benzoyl]amino-4,4’-dihydroxypropanoic acid (CMTMR; 8 μM), or 4-chloromethyl-6,8-difluoro-7-hydroxycoumarin (CMF-HC; 20 μM) for 10–60 min at 37˚C and adoptively transferred into NOD.SCID mice. For comparison, mouse CD4+ T cells were isolated from LNs of wild-type C57.BL/6J mice by magnetic negative selection (Miltenyi Biotec), labeled with CFSE, and transferred into NOD.SCID mice. CD3+ human T cells were labeled with CMTMR and transferred into reconstituted NOD.SCID.B2 mice. For all adoptive transfer experiments, 5 × 106 labeled cells of each cell type were transferred by tail-vein or retro-orbital injection.

**Transfection and retroviral transduction**

Primary human T cells were transfected by nucleofection (Amaxa, Allendale, NJ), using the high-viability protocol with enhanced GFP (eGFP), eGFP-tagged Ori1-E106A mutant, or mock transfected with no plasmid as indicated. Human cells were used for experiments 24 h after transfection. Constructs were designed previously (33). Retroviral transduction of purified mouse T cells was performed as described previously (34). Thy1.1-cyan fluorescent protein (CFP) or Thy1.1-eGFP-Ori1-E106A retroviral vectors were transduced into 293T cells and retroviral supernatant was collected. Activated primary mouse T cells were spin-fected using retroviral supernatant and 4 μg/ml Polybrene (Sigma-Aldrich, St. Louis, MO) for 1 h at 1800 rpm and incubated for 24–48 h at 37˚C. Transduced mouse T cells expressing the retrovirus (Thy1.1+) were magnetically isolated from nonexpressing cells (Thy1.1−) using a biotinylated anti-Thy1.1 Ab according to manufacturer’s instructions (Miltenyi Biotec). After purification, Thy1.1−CFP+ cells were labeled with CMTMR and Thy1.1−eGFP-Ori1-E106A− cells were labeled with CFSE. Also, Thy1.1−CFP+ were labeled with CMTMR, and Thy1.1−eGFP-Ori1-E106A− were labeled with CFSE. For determination of homing efficiency, fluorescently labeled Thy1.1− or Thy1.1+ populations were coadaptively transferred into wild-type C57.B16 recipient mice.

**Two-photon imaging and analysis**

Multidimensional (x, y, z, time, emission wavelength) two-photon microscopy was used to image fluorescently labeled lymphocytes in explanted mouse LNs, using a 780-nm femtosecond pulsed laser as described previously (35). Fluorescence emission was split by 510- and 560-nm dichroic mirrors into three detector channels, used to visualize CFP-labeled (green), CMTMR-labeled (red), and CMF-HC-labeled (blue) adoptively transferred cells. Tissue was oriented with the hilum away from the dipping objective (Olympus 20×, NA 0.9) of the upright microscope to image the LNs. The node was maintained at 36–37˚C by perfusion with medium (RPMI 1640) bubbled with carbogen (95% O2/5% CO2). Three-dimensional image stacks were sequentially acquired, and the x, y, z, coordinates of individual lymphocytes in the intact lymphoid organ were used to create individual cell tracks. Imaging volumes of x = 200 μm, y = 162 μm, and z = 50 μm were acquired at 18- to 20-s intervals using MetaMorph software (Universal Imaging, Sunnyvale, CA). This volume collection was repeated for up to 40 min to create a four-dimensional data set. Data were processed and analyzed using Imaris software (Bitplane, Oxford, UK). All clearly visible cells were tracked. From each video segment to reduce selection bias in our analysis of motility and trajectory.

**Immunohistochemistry**

LN specimens were fixed in 4% neutrally buffered formaldehyde. Five- to 6-μm-thick paraffin-embedded serial tissue sections were mounted on positively charged glass slides. Sections were taken for HE staining or immunofluorescence staining. Sections were bathed in xylene, then treated with a graded series of alcohol (100, 95, and 75% ethanol [v/v] H2O) and rehydrated in PBS (pH 7.5). Sections were treated with Retriviagen A (pH 0.0, 0.1 M), and microwave treated for 10 min for Ag retrieval. Sections were then treated with 3% H2O2 in PBS to block endogenous peroxidase activity for 10 min followed by incubation with blocking solution containing 1% normal goat serum for 20 min. Samples were then stained with anti-pan-
node addressin mAb MECA-79 (1:20 dilution, rat IgM; BD) overnight at 4°C. Secondary detection Abs were Cy5-conjugated goat anti-rat IgM+IgG (Jackson Laboratories). Slides were analyzed by fluorescence microscopy using a Zeiss Axioskop microscope (Carl Zeiss, Thornwood, NY).

**Flow cytometry**

Cells were analyzed using FACSCalibur flow cytometer with Cell Quest software or a LSRII with FACSDivina software (Becton Dickinson, Mountain View, CA). Anti-human CD45, CD4, CD8, CCR7, and CD62L, and the corresponding isotype control Abs were obtained from eBioscience (San Diego, CA). Conformation-specific Abs against human LFA-1 (MEM-83) and VLA-4 (12G10) were purchased from Abcam (Cambridge, MA). TRITC-conjugated secondary Abs were purchased from KPL (Gaithersburg, MD). Annexin V-PE was purchased from BD. Cells were suspended in buffer containing 0.5% BSA and 2 mM EDTA in PBS (staining buffer), and stained for 20–30 min with specific PE-Cy5-conjugated mAbs. Unbound Abs were removed by two washes with staining buffer, and cells were resuspended in staining buffer and analyzed. For homing experiments, mouse LNs and spleen were collected 18 h after adoptive transfer and made into single-cell suspensions with 70-μm nylon cell strainers (BD Falcon, Mountain View, CA). Cells were gated for appropriate forward and side scatter. Data were saved as FCS 2.0 list mode files and analyzed with FCS Express 3.1 or FlowJo.

**Single-cell Ca2+ imaging and intracellular Ca2+ measurement by flow cytometry**

Ratiometric single-cell Ca2+ imaging was performed as described previously (36). Cells were loaded by incubation for 30 min with 3 μM fura-2 AM (Invitrogen) and analyzed after 2 μM thapsigargin (TG; Sigma-Aldrich) treatment. Data were analyzed with MetaFlour software (Universal Imaging) and Origin software (OriginLab, Northampton, MA). For flow cytometry, cells were analyzed on a BD LSRII flow cytometer with FACSDivina software (BD). eGFP- or eGFP-Orai1-E106A–transfected CD3+ human T cells were loaded by incubation for 30 min with 8 μM fura red, AM (Invitrogen) at 37°C for 30 min, washed once, and resuspended in 2 mM Ca2+ RPMI 1640 supplemented with 1% FCS at 5 × 105 cells/ml. An 800-μl aliquot was warmed to 37°C before stimulation. Cells were maintained at 37°C and analyzed at 500 events/s. After establishment of the baseline, the tube containing the cells was removed, stimulant was added, and the tube was replaced. Recording was continued for up to 200 s. Data were saved as FCS 2.0 list mode files and analyzed with FlowJo Kinetics tool. [Ca2+]i, was determined as described previously (37).

**Chemokines and transwell assays**

Chemokines were obtained from PeproTech (Rocky Hill, NJ) and prepared according to manufacturer’s instructions. Transwell assay plates with 5-μm pore size were obtained from Corning (Tewksbury, MA). HUVECs were isolated as described previously (38). HUVECs were plated in M199 (Invitrogen) supplemented with 10% FCS and endothelial cell (EC) supplement (BD) on 1% gelatin-coated inverted transwells and allowed to proliferate to confluency. HUVECs and transfected human T lymphocytes were plated in M199 (Invitrogen) supplemented with 10% FCS. The percentage migrated into the lower chamber was measured as a fold increase over control.

**Soluble ICAM-1 binding assay**

Transfected human CD3+ T cells were suspended in 2 mM Ca2+ RPMI 1640 supplemented with 1% FCS. Cells were treated with 100 ng/ml rmCCL21 in the presence of ICAM-1/Fc (20 μg/ml; R&D Systems) andallophycocyanin-conjugated anti-human IgG1 (Fc specific; Southern Bio-technology) for 10 min at 37°C. Change in ICAM-1 binding determined by flow cytometry was measured as a fold increase over control.

**Statistical analysis**

Statistical significance was determined using the Student t test. Two-way analysis of covariance was used where indicated and was calculated using R Statistical Computing Software (Vienna, Austria). A p value <0.05 was considered significant. Data are presented as mean ± SEM.

**Results**

**Live-cell imaging of human lymphocytes in immunocompromised mice**

To establish a system for in vivo imaging of human lymphocytes, we initially transferred human lymphocytes into NOD.Scid mice by i.v. injection (Supplemental Fig. 1A). Purified CD4+ and CD8+ T cells and CD19+ B cells distributed randomly throughout the atrophied LN, not localizing into any compartmentalized structure (Supplemental Fig. 2A, 2B, 2C), and actively migrated in a stop-and-go pattern of motility (Fig. 1A, Supplemental Video 1). Human CD4+ T cells migrated with a mean velocity of 9.3 ± 0.4 μm min−1 (Fig. 1B). For comparison, adaptively transferred wild-type mouse CD4+ T cells also distributed randomly throughout the LN of NOD.Scid mice (Supplemental Fig. 2D) and migrated with slightly higher mean velocities averaging 10.9 ± 0.2 μm min−1 (Supplemental Fig. 2E). Migrating human CD4+ T cells traversed a wide area, and their displacement from the origin increased in proportion to the square root of time (Fig. 1C, 1D). These properties are consistent with a default random-walk pattern of motility of human T cells in the immunocompromised mouse host. Analysis of turning angles between consecutive imaging frames revealed a mean turning angle of 69 ± 4.5 degrees (Fig. 1E). Human CD8+ T cells performed similarly to human CD4+ cells, migrating with a mean velocity of 9.7 ± 0.4 μm min−1 (Fig. 1F), with displacement increasing as a square root function of time (Fig. 1G, 1H) and with a similar mean turning angle (Fig. 1I). Human CD19+ B cells colocalized with and migrated more slowly than CD4+ or CD8+ T cells, with a mean velocity of 5.8 ± 0.3 μm min−1 (Fig. 1J), displacing about half the distance of human T cells (Fig. 1K, 1L), and along more tortuous paths with a mean turning angle of 94 ± 4.8 degrees (Fig. 1M). This analysis of human lymphocyte motility is in agreement with previous two-photon imaging, which showed that murine T cells migrate faster and along straighter paths than B cells (39). We conclude that human T and B cells migrate similarly to mouse T and B cells in the host LN environment, with velocities that are only slightly slower than mouse cells. Despite the lack of organization into separate T zones and follicles, both human and mouse T cells migrated faster than colocalized B cells, implying an intrinsically faster mechanism for crawling.

**Functional CRAC channels are required for human T cell homing in NOD.Scid mice**

The human xenograft model provides an opportunity to observe human lymphocytes that have been manipulated by transfection before adoptive transfer. Purified human CD4+ T cells were transfected with control eGFP or with the dominant-negative eGFP-Orai1-E106A CRAC channel mutant and rested overnight in culture. Expression of eGFP-Orai1-E106A had no effect on cell viability before transfer as determined by Annexin V staining, an indicator of apoptosis (Supplemental Fig. 3A). Aliquots of cells were evaluated by flow cytometry, revealing a transfection efficiency of >50% (Fig. 2A). Ca2+ imaging confirmed that, as expected, human T cells expressing eGFP-Orai1-E106A showed impaired Ca2+ influx readdition of external Ca2+ after Tg treatment to deplete ER Ca2+ stores (Fig. 2B). T cells were transferred into an NOD.Scid recipient and analyzed by flow cytometry (Supplemental Fig. 1B) to quantify human T cell homing to the SLO in the xenograft model. Control eGFP human T cells efficiently homed to both the LNs and spleen. In contrast, eGFP-Orai1-E106A human T cells were almost entirely deficient in homing to the LN and, to a lesser extent, the spleen (Fig. 2C). To confirm that the homing defect is not due to differential CD62L or CCR7 expression, we quantified the surface phenotype of control eGFP and eGFP-Orai1-E106A–transfected human T cells, eGFP-Orai1-E106A human T cells maintained CD62L expression at marginally lower levels than the control population (Fig. 2D). CCR7 expression was unchanged in both populations (Fig. 2E).

We evaluated whether suppression of Ca2+ signaling through Orai1 channels could lead to an accumulation of Orai1-E106A–transfected human T cells.
T cells in the HEV after adoptive transfer. To determine the localization of T cells relative to the HEV, we isolated LNs from NOD.SCID mice after adoptive transfer of equal numbers of control CMTMR-labeled and eGFP-Orai1-E106A–transfected human T cells (Supplemental Fig. 3B). Fixed LNs were sliced from the base of the hilum and stained for peripheral node addressin, a carbohydrate epitope specific for the HEV (40). Both control and eGFP-Orai1-E106A+ human T cells were observed in regions that contained HEVs (Fig. 3A). Control human T cells (red) were positioned both within and outside of HEVs (blue) that perfuse the LN cortex, whereas T cells expressing eGFP-Orai1-E106A+ were observed predominantly within the HEVs (Fig. 3B). Fewer total eGFP-Orai1-E106A+ T cells were observed in selected areas containing HEVs, compared with control T cells (Supplemental Fig. 3C). Collectively, these findings indicate that human T cells require CRAC channel function for LN homing across the HEVs in our xenograft model.

Migration and homing in reconstituted immunocompromised mice

To extend these results on the requirement of CRAC channel function for homing, we performed additional validation studies using NOD.SCID.B2 mice that had been previously reconstituted with human PBLs. Reconstitution of immunocompromised mice was achieved by i.p. injection of purified human PBLs. Three weeks after PBL injection, LNs and spleens from NOD.SCID.B2 mice increased in size. Mean LN perimeter grew from 0.39 to 0.82 cm; mean spleen perimeter increased from 2.0 to 3.5 cm (Supplemental Fig. 4A). H&E staining of nonreconstituted NOD.SCID.B2 LN slices showed a lack of structure and nuclear...
density. Cellular density increased after 3 wk of reconstitution by human PBL, but cellular density was lower and follicular structure undefined compared with wild-type LNs (Supplemental Fig. 4B). To confirm that human T cells could migrate with characteristic motility behavior in the reconstituted NOD.SCID.B2 LN, CD3+ T cells from the original donor were transferred into the host animal after reconstitution (Supplemental Fig. 1C). Time-lapse two-photon imaging revealed robust T cell migration in the reconstituted LN (Fig. 4A, Supplemental Video 2), where T cells migrated with a mean velocity of 12.8 ± 0.4 μm min⁻¹ (Fig. 4B). Human CD3+ T cells migrated away from their starting coordinates, with mean displacement increasing as a square root function of time, typical of a random walk (Fig. 4C, 4D). The average CD3+ step-wise turning angle was 110.9 ± 2.5 degrees (Fig. 4E). Normal representative T cell migration and increased LN cellularity indicated that reconstituted NOD.SCID.B2 mice can also be used for analysis of transfected human immune cells.

To validate our finding that human T cells expressing the dominant-negative Orai1 mutant are unable to home to the LN and spleen in vivo, we performed a homing study in reconstituted NOD.SCID.B2 mice. Equal numbers of mock-transfected, CMTMR-labeled control and Orai1-E106A–transfected human T cells were adoptively transferred into NOD.SCID.B2 mice that had previously been reconstituted with blood from the same donor. Whole-organ flow cytometry revealed that control CMTMR-labeled human T cells homed to the SLO, representing 0.8 and

**FIGURE 2.** Human T cells expressing Orai1-E106A cannot home into the SLO of NOD.SCID mice. (A) Expression of eGFP and eGFP-Orai1-E106A in human T cells 24 h after transfection. (B) Averaged Tg-dependent influx of Ca²⁺ in human T cells expressing eGFP or eGFP-Orai1-E106A. Data are representative of three different experiments. (C) Human lymphocyte homing capacity in NOD.SCID mouse LN and spleen 18 h after adoptive transfer. Percentage of human cells expressing eGFP or eGFP-Orai1-E106A in SLO determined by flow cytometry. (D) Surface expression of CD62L on eGFP⁺ (left) and eGFP-Orai1-E106A⁺ human T cells (right), 18 h posttransfection. Isotype control staining represented in dotted gray lines. (E) Surface expression of CCR7 on eGFP⁺ (left) and eGFP-Orai1-E106A⁺ human T cells (right), 18 h posttransfection. FACS plots are representative of three separate experiments.

**FIGURE 3.** Human T cells expressing Orai1-E106A remain localized to the HEV. (A) Localization of human CD3⁺ T cells in the LN of a NOD.SCID mouse. This image shows CMTMR-labeled control (red) or eGFP-Orai1-E106A–transfected (green) human CD3⁺ T cells in a LN slice from a NOD.SCID mouse. HEVs were labeled with anti–peripheral node addressin Ab MECA-79. Inset shows labeled human cells localized in the HEV and cortex. (B) The percentage of human T cells found within the HEV.
0.2% of the total cells in the LN and spleen, respectively (Fig. 4F). Human T cells expressing the Orai1-E106A mutant lost their homing capacity to the SLO, representing 0.1% of the total cellularity in both organs (Fig. 4F). The majority of the cells recovered by flow cytometry were CMTMR+ controls (Fig. 4G), demonstrating that eGFP-Orai1-E106A+ human cells cannot home to the SLO in the human-reconstituted mouse model.

Homing block in mouse T cells with disrupted CRAC channel function

The results so far point to a requirement for CRAC channel function in the homing of human T cells in acutely and stably engrafted immunocompromised mice. To determine whether functional CRAC channels are also required for the homing of mouse T cells into the LN, we engineered MSCV-IRES retroviral vectors containing the variant Thy1.1 gene as a marker for expression, together with either eGFP-Orai1-E106A or a control CFP construct. We then transduced wild-type mouse T cells and examined their Ca2+ responses after store depletion by single-cell Ca2+ imaging. As expected, Ca2+ influx was suppressed in mouse T cells expressing eGFP-Orai1-E106A, relative to control CFP+ T cells (Fig. 5A). Using Thy1.1 coexpression and magnetic bead separation to purify cells, we labeled control and eGFP-Orai1-E106A–expressing T cells with CMTMR or CFSE, respectively, and evaluated homing after cotransfer into a wild-type recipient (Supplemental Fig. 1D). By flow cytometric analysis of recovered cells, Thy1.1− cells that did not express retroviral constructs were effectively recovered from LN and spleen (Fig. 5B). In addition, Thy1.1+CFP+ T cells, representing control cells that were successfully transduced, also were able to home to the SLO, representing >8% of the total cellularity (Fig. 5C, top gate). In contrast, adoptively transferred Thy1.1+eGFP-Orai1-E106A+ mouse T cells represented <0.2% of the total cellularity of the LN and spleen (Fig. 5C, bottom gate). We conclude that mouse T cells deficient in CRAC channel function are unable to home to the SLO of syngeneic mice, in agreement with our results using human cells in the xenograft models.

Orai1-E106A expression disrupts CCL21-dependent Ca2+ signaling and LFA-1 activation

Because treatment with CCL21 (the chemokine implicated in T cell homing) evokes Ca2+ signals in human T cells (25, 26), we hypothesized that CRAC channels mediate this response. To test for CRAC channel activity during the chemokine response, we used time-lapse flow cytometry to measure Ca2+ influx after Tg or CCL21 treatment in untransfected control human T cells and in cells transfected with either eGFP or eGFP-Orai1-E106A (Supplemental Fig. 1E). To achieve this, we used the Ca2+ indicator dye fura red, which reports increases in [Ca2+]i by a decrease in

FIGURE 4. Human T cell motility and homing in the SLO of reconstituted NOD.SCID.B2 mice. (A) Time-lapse two-photon image in an explanted reconstituted NOD.SCID.B2 LN, showing transferred human CD3+ T cells labeled with CMTMR (pseudocolored green). Panels depict cell positions at the times indicated, together with superimposed tracks depicting progressive cellular migration since the beginning of the record. (B) Mean cellular velocities of human CD3+ T cells shown with the mean ± SEM, 74 individual cell tracks. (C) Three-dimensional tracks of 10 human CD3+ T cells monitored throughout over 10-min intervals, normalized to starting coordinates. (D) Mean displacement of the 10 T cells as a square root function of time. (E) Distribution of turning angles between individual CD3+ T cell steps over 10-min intervals shown with mean turning angle ± SEM. (F) Homing of control CMTMR+ or eGFP-Orai1-E106A+ human CD3+ T cells in the reconstituted NOD.SCID.B2 mouse. T cell gates are drawn as indicated. (G) Percent ± SEM of recovered cells after adoptive transfer, detected by flow cytometry (three individual donors). p < 0.001.
fluorescence signal. As expected, Tg treatment resulted in a sustained [Ca\(^{2+}\)]i increase in both untransfected and eGFP+ control T cell populations (Fig. 6A). In contrast, cytosolic Ca\(^{2+}\) did not change in eGFP-Orai1-E106A+ T cells. Recombinant mouse CCL21 stimulation resulted in a rapid and transient increase in [Ca\(^{2+}\)]i, in both control untransfected and eGFP+ human T cells. However, human T cells expressing the dominant-negative eGFP-Orai1-E106A construct did not respond to CCL21 (Fig. 6B). Measurements of peak [Ca\(^{2+}\)]i levels demonstrate that human T cells deficient in CRAC channel function are unable to mobilize Ca\(^{2+}\) after Tg or CCL21 treatment (Fig. 6C).

Because CCL21 treatment has been previously shown to induce T cell binding to ICAM-1 (22), we tested whether suppression of CCL21-dependent Ca\(^{2+}\) signaling could lead to a defect in migration across a confluent EC monolayer in vitro. Using a standard chemotaxis assay, we measured migration of control eGFP+ or eGFP-Orai1-E106A+ human T cells in response to CCL21 across an uncoated polycarbonate transwell. Both T cell populations migrated robustly in response to the chemokine gradient (Fig. 7A), revealing that Orai1 function is not required for chemotaxis in the uncoated transwell. However, eGFP-Orai1-E106A+ human T cells were deficient in migrating across transwells coated with ECs independent of the CCL21 concentration (Fig. 7B), suggesting a role for Orai1 in facilitating T cell–EC interactions independent of CCR7. Because chemokine-induced activation of integrins is required for transendothelial migration and LN entry (22, 23), we investigated the role of Orai1 in integrin activation after CCL21 treatment (Supplemental Fig. 1F). Flow cytometry revealed that CCL21-induced activation of LFA-1 integrin was abolished in eGFP-Orai1-E106A+ human T cells (Fig. 7C), although VLA-4 \(\beta_1\) integrin was unaffected (Fig. 7D). Chemokine treatment has been shown to enhance binding of soluble ICAM-1 to leukocytes, demonstrating LFA-1 activation (41). CCL21 treatment enhanced ICAM-1 binding to control T cells, and binding was reduced in eGFP-Orai1-

**FIGURE 5.** Mouse T cells expressing Orai1-E106A cannot take up Ca\(^{2+}\) or home to the SLO of wild-type syngeneic mice. (A) Fura-2 fluorescence ratios in transduced mouse CD3+ T cells loaded with fura-2. (B) Homing of control mouse T cells after transduction. Successfully transduced cells (Thy1.1+) were purified via positive selection. Unbound mouse T cells (Thy1.1-) transduced with the CFP vector were labeled with CMTMR, and Thy1.1- cells transduced with the eGFP-Orai1-E106A vector were labeled with CFSE and adoptively transferred. Whole organs were analyzed with T cell gates and are drawn as indicated. (C) Homing of positively isolated (Thy1.1+) mouse T cells. Cells were labeled: CFP+ with CMTMR and eGFP-Orai1-E106A+ with CFSE.

**FIGURE 6.** Inhibition of Ca\(^{2+}\) signals evoked by Tg or CCL21 treatment in human T cells expressing Orai1-E106A. (A) Flow cytometry analysis, using the FlowJo Kinetics tool, of human T cells loaded with fura red. After 90 s to establish a baseline, cells were treated with 2 \(\mu\)M Tg. Mean fluorescence changes (lower fluorescence = higher [Ca\(^{2+}\)]) are shown in untransfected human T cells (top), eGFP+ human T cells (middle), and eGFP-Orai1-E106A+ human T cells (bottom). (B) Fluorescence changes in untransfected human T cells (top), eGFP+ human T cells (middle), and eGFP-Orai1-E106A+ human T cells (bottom) after addition of 100 ng/ml recombinant mouse CCL21. Data are from one donor and are representative of three donors. (C) Mean peak [Ca\(^{2+}\)], signals evoked by Tg and CCL21 in control and eGFP-Orai1-E106A+ human T cells.
We conclude that Orai1 is required for CCL21-induced Ca\(^{2+}\) signaling in human T cells and subsequent LFA-1 activation, transendothelial migration, and LN homing.

**Discussion**

We show that CRAC channel function is required for T cell homing into SLO. Our results are demonstrated in a combined in vivo and in vitro assessment of human T cells in mouse xenograft models, as well as in retrovirally transduced mouse T cells in syngeneic recipient mice. By inhibiting native CRAC channels with a dominant-negative Orai1 mutant (E106A), our approach avoids potential compensation in Orai1 knockout mouse models that retain residual CRAC channel function. Moreover, these results establish a system for live two-photon imaging of labeled or transfected human lymphocytes in an in vivo human xenografted mouse environment.

**Homing and motility of human lymphocytes in xenografted mice**

Using human lymphocytes transferred acutely into immunocompromised mice or into reconstituted human xenografted mice, we describe two-photon imaging of human T and B lymphocytes within intact LNs in a preparation that does not require human fetal tissue. Our methods differ from a recent study that used two-photon microscopy to image human T cell migration in mice reconstituted using fetal human blood, liver, and thymus (42). Both NOD.SCID and NOD.SCID.B2 mice retain peripheral LNs that support engraftment of human lymphocytes. These two strains were chosen because LNs from other immunocompromised mouse strains, including Rag2\(^{-/-}\) and NSG, are even more atrophied (43, 44), rendering them unsuitable for imaging. NOD.SCID mice require depletion of resident NK cells before transfer of human lymphocytes, and LNs remain small after engraftment, although imaging was possible. Conversely, NOD.SCID.B2 mice readily support human xenografts, allowing for expansion of lymphocyte populations and stable engraftment of human PBLs. After reconstitution, LNs are enlarged and easily used for tracking individual labeled cells from the same blood donor by two-photon microscopy.

Our results demonstrate that adoptive transfer, both acutely and into reconstituted SCID mouse models, enables quantitative imaging and analysis of human cells. Human T cells homed into and actively migrated through the atrophied mouse host LNs of NOD.SCID mice with velocities that were somewhat slower than mouse T cells in the same environment (9.3 ± 0.4 versus 10.9 ± 0.2 µm min\(^{-1}\), re-
respectively), similar to velocities reported for the fetal human blood, liver, and thymus mouse model (42). Compared with the acutely transferred model, human T cells transferred into reconstituted mice migrated with faster mean velocity (12.8 ± 0.4 μm min⁻¹), closely similar to wild-type mouse T cells in the LN (27). Human T and B cell relative velocities mirrored the migration of mouse lymphocytes; both human and mouse T cells migrate more rapidly than CD19⁺ B cells in the xenograft environment (Fig. 1), as originally observed for mouse T and B cells in mouse LN (39). Displacement analysis confirmed that, like mouse B cells, human B cells migrate with lower velocities and motility coefficients than T cells, reflecting slower migration. Our results suggest that more rapid migration is an intrinsic characteristic of T cells.

CRAC channel activity required for homing

Our results used the dominant-negative Orai1-E106A mutant to suppress CRAC channel function in mouse and human T cells, and thereby elucidate mechanisms by which these ion channels regulate T cell function. Orai1-E106A has been used previously to determine the requirements for Orai1 clustering at the immunological synapse (33), to identify STIM1–Orai1 interacting domains (45), to determine regulation of voltage-gated Ca²⁺ channels by STIM1 (46), to reveal mechanisms for protection against apoptotic signaling through CD95 (47), and to demonstrate Ca²⁺ release from secretory granules (48). In Orai1-deficient mice, Ca²⁺ entry is not fully abolished in T cells (17, 18). Moreover, whereas human patients with the homozygous Orai1-R91W mutation do not have functional CRAC channels (7), knock-in mice with the homozygous mutation exhibit slower homing and lower cellularity in secondary LN compared with wild-type mice (49). In comparison with these knock-in and knockout mouse models, our use of Orai1-E106A expression specifically and completely suppresses CRAC channel function in human and mouse T cells, which we demonstrate to result in a severe homozygous defect in both the xenograft or wild-type mouse environment (Figs. 2, 4, 5).

We further show that transendothelial migration and homing of lymphocytes to LN requires Ca²⁺ influx through Orai1 channels. During homing, T cells must adhere to ECs and then extravasate from the HEV to the LN paracortex. Consistent with a role of Ca²⁺ influx, our results demonstrate that Orai1 assists in LFA-1 (α₃β₂ integrin) activation by providing the initial Ca²⁺ signal in response to CCL21 treatment. We previously showed that inhibition of Kv1.3 channels, which are required for sustained Ca²⁺ signaling in lymphocytes (12), disrupts VLA-4 (α₅β₇ integrin) activation in effector memory T cells and inhibits their migration in inflamed tissues (50). Our results do not fully account for where eGFP-Orai1-E106A T cells reside after transfer. Other studies also support a requirement for Ca²⁺ signaling in T cell migration. T cell-specific deletion of the STIM proteins that activate Orai1 Ca²⁺ channels abolishes chemokine-induced Ca²⁺ influx and diminishes chemokinesis/toward chemokine gradients (51). Orai1 has also been proposed to mediate migrational guidance in response to high-affinity LFA-1–ICAM interactions after shear stress in neutrophils (52) and in chemotaxis of platelets (53). Moreover, CCR7 signaling enhances overall lymphocyte motility (chemokinesis) in the LN after homing (54–57). Thus, CCR7 signaling and the resulting CRAC channel activation mediated by STIM1–Orai1 interaction may modulate both homing and chemokinesis by enhancing intercellular adhesion. Our findings extend the scope of immune system functions controlled by Orai1 to include homing, integrin activation, and transendothelial migration.

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Disclosures

The authors have no financial conflicts of interest.

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Orai1 Function is Essential to Facilitate T Cell Homing to Lymph Nodes

Data Supplement

• Supplemental Figures 1-4

• Supplemental Videos 1 and 2

Supplemental Figure 1: Diagrams of experimental protocols. A, Timeline of adoptive transfer for FACS, confocal, and two-photon microscopy of NOD.SCID mice for imaging experiments in Figure 1. B, Timeline of adoptive transfer for FACS of NOD.SCID mice for homing experiments in Figure 2. C, Timeline representing reconstitution protocol, FACS, and two-photon microscopy with reconstituted NOD.SCID.B2 mice in Figure 4. D, Timeline of experiments for homing of transduced wildtype mouse T cells in wildtype hosts in Figure 5. E, Timeline of timelapse flow cytometry to measure Ca^{2+} responses in human T cells in Figure 6. F, Timeline of treatment and staining for measuring integrin activation on human T cells in Figure 7C.

Supplemental Figure 2: Human and mouse lymphocyte subsets home to the lymph node of NOD.SCID mice. A, Localization of CD4^{+} (green), CD8^{+} (red), and CD19^{+} (blue) human lymphocytes in the NOD.SCID mouse LN analyzed by confocal microscopy. B, Localization of human lymphocytes analyzed by 2-photon microscopy. Light blue color at the top of the imaging volume is second-harmonic generation emitted from fibrous collagen in the LN capsule following two-photon excitation. C, Purity of labeled human lymphocytes. Following dye labeling, lymphocytes were stained with PE-Cy5 conjugated antibody against human CD4, CD8, and CD19. D, Localization of mouse (green) and human (red) CD4^{+} T cells in the NOD.SCID mouse LN. E, Mean cellular velocities of mouse CD4^{+} T cells in the NOD.SCID mouse LN shown with the mean and standard error of the mean, 575 individual cell tracks.
Supplemental Figure 3: *eGFP-Orai1-E106A expression does not influence Annexin V staining.*

A, Staining of Annexin V-PE on untransfected (left), eGFP\(^+\) (middle), and eGFP-Orai1-E106A\(^+\) (right) 18 hr post transfection. B, Flow cytometry profile of an aliquot of cells prior to adoptive transfer in Fig. 3, showing roughly equivalent transfer of both control CMTMR\(^+\) and eGFP-Orai1-E106A\(^+\) human T cell populations. C, Quantification of the total number of human T cells within selected images of lymph node diffuse cortex that contain HEV. Note the number of control and eGFP-Orai1-E106A\(^+\) cells includes cells outside and within HEV.

Supplemental Figure 4: *Reconstitution increases the size and cellularity of SLO from NOD.SCID.B2 mice.* A, SLO perimeter of reconstituted and control NOD.SCID.B2 mice. Perimeter of the spleen and lymph nodes ± SEM. P < 0.001 B, Hematoxylin and eosin (H&E) staining of LN slices from control wildtype C57.Bl6, NOD.SCID.B2, and reconstituted NOD.SCID.B2 mice.

Supplemental Video 1. Imaging human lymphocyte motility in the NOD.SCID mouse LN.
Images of human CD4\(^+\) (green), human CD8\(^+\) (red), and human CD19\(^+\) (blue) lymphocytes within NOD.SCID LN explants were acquired by two-photon microscopy. Time-lapse video shows robust lymphocyte motility in the LN. Tick marks, 40 \(\mu\)m. (1.9 MB)

Supplemental Video 2. Imaging human T cell motility in the reconstituted NOD.SCID.B2 mouse LN.
Images of human CD3\(^+\) (green) T cells within reconstituted NOD.SCID.B2 LN explants were acquired by two-photon microscopy. Time-lapse video shows lymphocyte motility in the LN. Tick marks, 20 \(\mu\)m. (0.8 MB)