In vivo equilibrium of proinflammatory IL-17+ and regulatory IL-10+ Foxp3+ RORγt+ T cells

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The nuclear hormone receptor retinoic acid receptor–related orphan receptor γt (RORγt) is required for the generation of T helper 17 cells expressing the proinflammatory cytokine interleukin (IL)-17. In vivo, however, less than half of RORγt+ T cells express IL-17. We report here that RORγt+ T cells include Foxp3+ cells that coexist with IL-17–producing RORγt+ T cells in all tissues examined. The Foxp3+ RORγt+ T cells express IL-10 and CCL20, and function as regulatory T cells. Furthermore, the ratio of Foxp3+ to IL-17–producing RORγt+ T cells remains remarkably constant in mice enduring infection and inflammation. This equilibrium is tuned in favor of IL-10 production by Foxp3 and CCL20, and in favor of IL-17 production by IL-6 and IL-23. In the lung and skin, the largest population of RORγt+ T cells express the γδ T cell receptor and produce the highest levels of IL-17 independently of IL-6. Thus, potentially antagonistic proinflammatory IL-17–producing and regulatory Foxp3+ RORγt+ T cells coexist and are tightly controlled, suggesting that a perturbed equilibrium in RORγt+ T cells might lead to decreased immunoreactivity or, in contrast, to pathological inflammation.

Abbreviations used: BAC, bacterial artificial chromosome; CIA, collagen-induced arthritis; DSS, dextran sodium sulfate; EGFP, enhanced GFP; LTi, lymphoid tissue inducer; RA, retinoic acid; RORγt, RAg receptor–related orphan receptor γt; T reg, T regulatory.

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T cells adapt immune responses not only to the antigens, but also to the type of pathogen they face. Cell-based pathogens, such as viruses, intracellular bacteria, and tumors evoke a CD4+ Th1 type of response characterized by the production of IFN-γ. In contrast, extracellular-pathogen–based pathways, such as worms and allergens, evoke a Th2 type of response characterized by the production of IL-4, IL-5, and IL-13 (1). Recently, some bacteria and fungi have been found to evoke a Th17 type of response characterized by the production of IL-17 (or IL-17A), IL-21, IL-22, IL-6, and TNF-α, as well as IL-23 by APCs (2–6). The proinflammatory Th17 response induces the recruitment of neutrophils and is involved in autoimmune inflammatory diseases. However, the reactivity of T cells is modulated by T regulatory (T reg) cells. These include so-called natural Foxp3+ T reg cells that are induced in the thymus (7) as well as adaptive Foxp3+ T reg cells that are induced in the periphery and produce TGF-β and/or IL-10 (8, 9).

The generation of these distinct T cell effector types is under control of specific cytokines and transcription factors (2). In mice, TGF-β is required for the generation of both Th17 and T reg cells. The switch to the Th17 pathway is induced by IL-6 and/or IL-21 (4–6, 10–12), and requires IL-23 for full effector maturation (2, 3). However, TGF-β together with IL-6 induces CD4+ T cells to produce both IL-17 and IL-10, a population that is not able to induce inflammation in mice (13).
In contrast, IL-23 generates proinflammatory cells that produce only IL-17. In accordance with these findings, spontaneous colitis is induced in mice deficient for IL-10 (14), and the inflammatory disease is prevented by concomitant deficiency in IL-23 (15). The immunosuppressive IL-10 can be produced by all types of T cells, including a subset of T reg cells (16), and is induced by IL-27 in Th1 and Th2 cells (17–19).

The equilibrium within RORγt+ T cells requires the forkhead/winged-helix transcription factor Foxp3 (23–25). Mice deficient in Foxp3 lack natural T reg cells (24) and are highly susceptible to inflammatory disease (26).

The development of inflammatory pathologies, such as experimental autoimmune encephalomyelitis (27, 28), an animal model of multiple sclerosis, collagen-induced arthritis (CIA) (29), and inflammatory bowel disease (15, 30), depends on Th17 cells and IL-23. In humans, a coding variant of IL23 confers protection to Crohn’s disease (31). The Th17–IL-23 pathway in also involved in resistance to a growing list of bacterial and fungal pathogens in mice (11, 32–35), and Th17 cells are readily isolated from patients with Candida infection (36). However, a comprehensive picture of the class of pathogens targeted by this pathway remains to be drawn. T reg cells, on the other hand, and IL-10 in particular, have been shown to limit inflammation and adaptive immunity to a variety of pathogens (37), as well as to promote long-term T cell memory (38). Collectively, these studies demonstrate the essential role of the Th17 and the T reg cells in immunity, immunoregulation, and immunopathology. They also suggest that a balanced expansion of these Th subsets during immune responses might ensure efficient yet restrained immunity that limits damage to self.

The nuclear hormone receptor RA receptor–related orphan receptor γt (RORγt) is a marker for Th17 cells and is required for their generation (39). Using an enhanced GFP (EGFP) reporter mouse, we found that RORγt+ Tαβ cells included IL-10–producing Foxp3+ T reg cells. These cells coexisted with IL-17–producing RORγt+ Tαβ cells in all tissues examined in an equilibrium regulated by Foxp3 and CCL20 in favor of Foxp3+ cells, and by IL-6 and IL-23 in favor of IL-17–producing cells. Foxp3 was shown to bind to RORγt, suggesting that it directly regulates RORγt activity. The equilibrium within RORγt+ Tαβ cells was maintained during massive inflammation of the intestine or the lungs, but it was perturbed in cancer. We suggest that the balanced expansion of RORγt+ Tαβ cell subsets during inflammation might ensure efficient yet regulated immunity, and that perturbation of this equilibrium might lead to inadequate immune reactivity to tumors, pathogens, or self.

RESULTS

Diversity and distribution of RORγt+ T cells in vivo

To optimize the visualization of RORγt+ cells, we have generated transgenic Rorc(γt)-GfpTG mice expressing EGFP under control of the Rorc(γt) locus on a bacterial artificial chromosome (BAC) (40). An estimated 5–10 copies of the BAC transgene were integrated, as assessed by quantitative PCR. The expression pattern of EGFP in Rorc(γt)-GfpTG mice was identical to the one described previously in Rorc(γt)/Gfp mice carrying one allele of Rorc(γt) inserted with EGFP (39, 41).

The expression pattern of EGFP was further confirmed by immunofluorescence histology of GFP and endogenous RORγt in various organs from Rorc(γt)-GfpTG mice. In all tissues analyzed, we found a strict correlation between GFP and RORγt protein expression (not depicted). High expression of GFP was found in fetal lymphoid tissue inducer (Ltii) cells, Ltii-like cells in the adult intestine, in immature CD4+CD8+ thymocytes (not depicted), and in a subpopulation of peripheral T cells (Fig. 1 A). The highest frequency of RORγt+ T cells was observed in the intestine, as previously reported (39), which were localized in the lamina propria of the villi, whereas CD3– RORγt+ cells, i.e., Ltii-like cells, were clustered in cryptopatches that are precursor structures to isolated lymphoid follicles (Fig. 1 C) (42). RORγt+ T cells were also found in all organs or tissues tested, including the LNs, spleen, lung, skin, and bone marrow (Fig. 1 A). Assessing the homogeneity of these populations within different organs, we observed that up to 50% of RORγt+ T cells in the lung and skin expressed the γδ TCR instead of the αβ TCR (Fig. 1 B). RORγt+ Tγδ cells expressed the highest levels of EGFP (Fig. 1 A) and none of the CD4 or CD8 coreceptors (Fig. 1 D).

RORγt+ T cells include both Th17 and IL-10–producing Foxp3+ T reg cells

We assessed whether RORγt was a marker for IL-17–producing cells in both Tαβ and Tγδ populations, and stained RORγt+ T cells for IL-17, IFN-γ, and Foxp3. Surprisingly, even though a majority of IL-17–producing cells expressed RORγt (Fig. 2 A), only 15–50% of RORγt+ Tαβ cells expressed IL-17, depending on the organ examined, and few but detectable numbers of RORγt+ Tαβ cells produced both IL-17 and IFN-γ (Fig. 2 B). In contrast, a majority (50–90%) of RORγt+ Tγδ cells expressed high levels of IL-17. Even more surprising, 15–50% of IL-17+ RORγt+ Tαβ cells expressed the T reg cell markers Foxp3 and CD25 (Fig. 2, B and C), with the highest incidence in LNs, even though Foxp3+ RORγt+ Tαβ cells represented only a minority of the total population of Foxp3+ T cells (Fig. 2 A). No RORγt+ Tαβ cells were found to express both IL-17 and Foxp3, and no RORγt+ Tγδ cells expressed Foxp3 (not depicted). The coexpression of RORγt and Foxp3 protein at the single cell level was confirmed by immunofluorescence histology of the LNs (Fig. 2 D). Furthermore, Foxp3+ RORγt+ Tαβ cells were functionally T reg cells, as they suppressed in vitro proliferation of activated CD4+ T cells (Fig. 2 E).

We next measured the expression, by the different subsets of RORγt+ T cells isolated by FACS (Fig. S1, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1), of a restricted panel of genes involved in the Th17
T reg cell pathways using quantitative qRT-PCR arrays. As compared with RORγT+ T cells, RORγT+ T cells expressed higher levels of Rorc, Il17a, Il17f, il22, and Il23r, as expected, as well as Il10, Ios, Cx3cl1, and Mmp9 (Fig. S2). Expression of these genes was further examined in the individual RORγT+ T cell subsets. As cells could not be fixed and stained for Foxp3 expression before isolation for RNA recovery, Foxp3+ RORγT+ T cells were isolated on the basis of CD25 expression. As expected, CD25− RORγT+ Tαβ cells enriched in IL-17-producing cells (Fig. 2 C), as well as RORγT+ Tγδ cells, expressed high levels of Il17a, Il17f, and Il22 transcripts (Fig. 3 A and not depicted). CD25+ RORγT+ Tαβ cells expressed detectable levels of Il17a and Il17f transcripts, probably due to the presence of a small population of IL-17–producing CD25+ cells (Fig. 2 C). In addition, RORγT+ Tγδ cells, but not RORγT− Tγδ cells, expressed Csf2, which codes for GM-CSF involved in the mobilization of granulocytes and monocytes.

CD25+ (Foxp3+) RORγT+ Tαβ cells specifically expressed high levels of the expected Foxp3 transcript, as well as Il10, Ios, Cx3cl1, and Mmp9. IL-10 expression was confirmed at the protein level, and Foxp3+, but not Foxp3− RORγT+ Tαβ, cells represented a major fraction of total IL-10–producing cells in LNs (Fig. 3 B), and to a lesser extent in the gut (Fig. S3, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1). The transcript coding for CCL20 was specifically expressed by Foxp3+ RORγT+ Tαβ cells and barely detected in other T cell populations. Furthermore, CCR6, the receptor for CCL20, was expressed by both CD25+ and CD25− RORγT+ Tαβ cells (Fig. S4), suggesting that CD25+ RORγT+ Tαβ cells can regulate the recruitment of RORγT+ Tαβ cells. Interestingly, Icos (43), Ccl20 (13, 44), and Mmp-9 (45, 46) have been associated with Th17 cells previously, but not with T reg cells, whereas Cx3cl1 is normally expressed by activated endothelial cells (47).

Generation and regulation of IL-17+ or Foxp3+ RORγT+ cells

We further assessed how the generation of IL-17–producing or Foxp3+ RORγT+ T cells in vivo was affected by the absence of factors shown to be required for the generation of Th17 or T reg cells, such as IL-6 (4–6, 10–12) and Foxp3 (24, 26). In IL-6–deficient mice, the number and proportion of RORγT+ Tαβ cells, in particular of IL-17–producing RORγT+ Tαβ cells, were significantly decreased, whereas the proportion of IL-10–producing Foxp3+ RORγT+ T cells was unaffected (Fig. 4, A, C, and D). This confirms the requirement of IL-6 for the generation of Th17 cells, but also shows that IL-6 is not required for the generation of IL-10–producing RORγT+ T cells, even though in vitro it acts in synergy with TGF-β to induce IL-10–producing CD4+ T cells (13, 17). In contrast, IL-17–producing RORγT+ Tγδ cells were unaffected by the absence of IL-6 (Fig. 4 B). Furthermore, in IL-12Rβ1–deficient mice, unresponsive to both IL-12 and IL-23 (2, 3), the proportion of IL-17–producing RORγT+ Tαβ cells was slightly decreased (Fig. 4, C and D).
Figure 2.  RORγt+ T cells include Th17 and Foxp3+ T reg cells. (A and B) Flow cytometry analysis of cells isolated from Rorc(γt)-GfpTG mice. Cells were restimulated in vitro with PMA/ionomycin for 5 h and subjected to intracellular staining for GFP, IL-17, Foxp3, and/or IFN-γ. Plots are gated on TCR-β+ cells (A) and GFP-TCR-β+ or GFP-TCR-β- cells (B). Numbers indicate mean percent cells in quadrants ± SD obtained with at least three Rorc(γt)-GfpTG mice. (C) CD4+ T cells isolated from the spleen and mesenteric LNs of Rorc(γt)-GfpTG mice were sorted into two populations based on their expression of GFP and CD25. Sorted cells were subsequently restimulated in vitro and stained for IL-17 and Foxp3. Results are representative of three independent experiments. (D) Immunofluorescence histology staining for GFP, RORγt, Foxp3, and nuclei (DAPI) of a mesenteric LN isolated from a Rorc(γt)-GfpTG mouse. Arrowheads indicate cells expressing RORγt and/or Foxp3. Bar, 20 μm. (E) Proliferation assay of CD4+ T cells isolated from the spleen and mesenteric LNs of Rorc(γt)-GfpTG mice and sorted into two populations based on their expression of GFP and CD25. CFSE-labeled CD25 responder T cells (2.5 × 10^4) were cultured alone (left) or together with either GFP-CD25+ T cells or GFP+CD25+ T cells (2.5 × 10^4 or 0.625 × 10^4 cells). Cells were cultured in duplicates for 3 d in the presence of 10^5 irradiated spleen cells and anti-CD3ε antibody. Numbers indicate frequency of proliferating cells (± SD) obtained from three independent experiments.
Marrow chimeras were generated by adoptive transfer of bone marrow from Foxp3sf × RorcH9253t-GfpTG and WT mice into irradiated lymphopenic hosts (Fig. S5, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1). In such chimeras, RORH9253t + T cells do not express Foxp3, but cells from WT mice protected the host from a scurfy phenotype (not depicted). Foxp3sf RORH9253t + T cells failed to generate IL-10–producing cells and generated a reduced proportion of IL-17–producing cells as compared with Foxp3wt RORH9253t + T cells in control chimeras (Fig. 5, A and B). This shows that Foxp3 protein or Foxp3 + cells are required for the generation of IL-10–producing RORH9253t + T cells and, surprisingly, also favor the differentiation of IL-17–producing RORH9253t + T cells.

To test whether Foxp3 regulates RORH9253t + T cells through direct interaction with RORγt, N-terminally FLAG-tagged RORγt was expressed in HEK293T cells together with Foxp3. Foxp3 was specifically coimmunoprecipitated with FLAG-RORγt, indicating that the two factors interact (Fig. 5 C). Collectively, these data suggest that Foxp3 in accordance with the important but not essential role of IL-23 in the generation of Th17 cells (2, 3).

It has recently been shown that Th17 cells isolated from SKG mice, which spontaneously develop arthritis, as well as human Th17 cells, express CCR6 (44, 48, 49). We therefore assessed the impact of CCR6 on the generation of IL-17–producing or Foxp3+ RORγt + T cells. In CCR6–deficient mice (50), contrary to IL6– or IL12Rβ1–deficient mice, the proportion of IL-17–producing RORγt + Tαβ cells was significantly increased (Fig. 4, C and D). As CD25+ (Foxp3+) RORγt + Tαβ cells specifically express Ccl20 transcripts (Fig. 3 A), these cells might regulate the generation of IL-17–producing RORγt + Tαβ cells through CCR6.

We next assessed whether Foxp3 regulates the generation of IL-17– or IL-10–producing RORγt + Tαβ cells. Mice bearing the Scurfy mutation in the Foxp3 gene (Foxp3sf mice) lack T reg cells (24, 26). In these mice, however, the analysis of RORγt + T cells was complicated by generalized autoimmunity, and the development of immature RORγt + CD4+CD8+ thymocytes was abnormal (not depicted). Therefore, mixed bone

![Figure 3](http://www.jem.org/cgi/content/full/jem.20080034/DC1)

**Figure 3.** RORγt+ T cells express genes involved in the Th17 or T reg cell pathway. (A) Cells isolated from the spleen and mesenteric LNs of Rorcγt-GfpTG mice were sorted into eight distinct populations based on their expression of GFP, CD3, TCR-β, TCR-γ, CD4, and CD25 (Fig. S1), and gene expression was assessed using real-time PCR. Ct values were normalized to the mean Ct of five housekeeping genes. Data are the mean of two or three independent experiments. (B) Foxp3+ RORγt+ T cells express IL-10. Cells isolated from LNs of Rorcγt-GfpTG mice were restimulated in vitro with PMA/ionomycin for 5 h and subjected to intracellular staining for GFP, IL-17, Foxp3, and IL-10 or an isotype control. Numbers indicate percent cells in quadrants. Results are representative of at least three individual experiments.
Equilibrium between IL-17+ or Foxp3+ RORγt+ cells during inflammation

Given the prominent role of IL-17+-producing cells in inflammation and of IL-10-producing cells in immune regulation, we assessed if and how the relative frequencies of the different RORγt+ T cell subsets were modified in mice enduring severe intestinal inflammation, virus infection, or cancer. Dextran sodium sulfate (DSS) was administered orally to Rorcγt-GfpTG mice until the development of chronic intestinal inflammation (Fig. 6 A). A high number of T cells was recovered in DSS-treated mice and both the frequency and the number of RORγt+ Tαβ cells increased, but the ratio of IL-17–producing to Foxp3+ RORγt+ T cells remained constant. Similar results were obtained in mice exposed to intranasal infection with influenza A virus, even though their lungs sustained heavy leukocyte infiltration (Fig. 6 B). In contrast, in Rorcγt-GfpTG × Apcmin/+ mice bearing colonic and ileal polyps (52), no significant infiltration of leukocyte and RORγt+ T cells was observed in the colon and the terminal ileum (Fig. S7, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1), whereas the mesenteric LNs were largely hyperplastic. The ratio of IL-17–producing to Foxp3+ cells in these LNs was significantly decreased, even though the frequency and total number of RORγt+ T cells were increased (Fig. 6 C). Similar results have recently been obtained comparing Th17 and Foxp3 cells in mice injected with B16 melanoma (53). These data indicate that the relative frequency of IL-17+-producing or Foxp3+ RORγt+ Tαβ cells can remain constant during severe inflammation or infection. However, chronic tumors can alter this balance in favor of the generation or recruitment of Foxp3+ RORγt+ Tαβ cells producing the immunosuppressive IL-10.

induces the generation of IL-10–producing RORγt+ T cells and regulates RORγt activity directly. On the other hand, RORγt is required for the generation of Th17 cells (39), but not for the generation of "mainstream" Foxp3+ and/or IL-10–producing T cells. In Rorcγt-CreTG × Rosa-StopDta mice, which lack RORγt+ T cells due to their specific expression of the lethal diphtheria toxin subunit A (51), no or few IL-17–producing cells were detected (not depicted), and the proportion of total Foxp3+ or IL-10–producing cells was unaffected (Fig. S6, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1).

Equilibrium between IL-17+ or Foxp3+ RORγt+ T cells during inflammation

Given the prominent role of IL-17+-producing cells in inflammation and of IL-10–producing cells in immune regulation, we assessed if and how the relative frequencies of the different RORγt+ T cell subsets were modified in mice enduring severe intestinal inflammation, virus infection, or cancer. Dextran sodium sulfate (DSS) was administered orally to Rorcγt-GfpTG mice until the development of chronic intestinal inflammation (Fig. 6 A). A high number of T cells was recovered in DSS-treated mice and both the frequency and the number of RORγt+ Tαβ cells increased, but the ratio of IL-17–producing to Foxp3+ RORγt+ T cells remained constant. Similar results were obtained in mice exposed to intranasal infection with influenza A virus, even though their lungs sustained heavy leukocyte infiltration (Fig. 6 B). In contrast, in Rorcγt-GfpTG × Apcmin/+ mice bearing colonic and ileal polyps and tumors (52), no significant infiltration of leukocyte and RORγt+ Tαβ cells was observed in the colon and the terminal ileum (Fig. S7, available at http://www.jem.org/cgi/content/full/jem.20080034/DC1), whereas the mesenteric LNs were largely hyperplastic. The ratio of IL-17–producing to Foxp3+ RORγt+ T cell in these LNs was significantly decreased, even though the frequency and total number of RORγt+ Tαβ cells were increased (Fig. 6 C). Similar results have recently been obtained comparing Th17 and Foxp3 cells in mice injected with B16 melanoma (53). These data indicate that the relative frequency of IL-17–producing or Foxp3+ RORγt+ Tαβ cells can remain constant during severe inflammation or infection. However, chronic tumors can alter this balance in favor of the generation or recruitment of Foxp3+ RORγt+ Tαβ cells producing the immunosuppressive IL-10.
population level (39), but not in individual cells expressing Foxp3. In contrast to their in vivo counterparts, however, few in vitro–derived Foxp3+ RORγt+ T cells expressed IL-10, which was rather expressed by RORγt+ cells. IL-10 was not induced by TGF-β alone but required the addition of IL-6, as found previously (13, 17), or the addition of IL-21. IL-10 production was inhibited in the presence of RA, even though RA favored the generation of Foxp3+ RORγt+ T cells and decreased the generation of IL-17–producing cells in the presence of TGF-β and IL-6. A combination of TGF-β with IL-6 (with or without RA) or IL-21 generated IL-17–producing as well as IL-10–producing or Foxp3+ RORγt+ cells, showing that the engagement of these pathways is not mutually exclusive at the cell population level, in agreement with recent reports (13, 17). RORγt+ T cells producing both IL-17 and IL-10 could be generated in the presence of TGF-β and IL-6, but Foxp3+ RORγt+ T cells produced neither IL-10 nor IL-17 (not depicted). Collectively, these data show that even though naive CD4+
T cells differentiate in vitro into IL-17–producing, IL-10–producing, or Foxp3+ RORγt+ Tαβ cells, substantial differences exist with the in vivo RORγt+ Tαβ cell subsets in the phenotype of the cells producing IL-10, and the proportion of Foxp3+ cells expressing RORγt.

**DISCUSSION**

In this study, we have shown that RORγt+ T cells include both IL-17– and IL-10–producing Foxp3+ T reg cells, which are endowed with proinflammatory and regulatory functions, respectively. In addition, RORγt+ T cells include Tγδ cells producing high levels of IL-17 in an IL-6–independent pathway. The relative frequency of IL-17– or IL-10–producing RORγt+ Tαβ cells is maintained constantly during infection or inflammation, suggesting that a robust mechanism maintains an equilibrium between these two effector arms. However, this equilibrium is perturbed in mice harboring intestinal tumors. IL-6, Foxp3, and CCR6, and to a lesser extent IL-23, are essential regulators of this equilibrium.

Ivanov et al. (39) reported that RORγt+ T cells or IL-17–producing cells were only present in the intestinal lamina propria. These authors therefore suggested that the intestinal flora induces the differentiation of RORγt+ IL-17–producing T cells. We show now that although these cells are more abundant in the intestine, sizeable populations of RORγt+ T cells were found in all organs examined, including the LNs, spleen, lung, skin, and bone marrow. This extended distribution of RORγt+ T cells was not due to infection, as specific pathogen–free mice were used in all experiments. This discrepancy might be explained by the sensitivity of the reporter mice that have been used. Ivanov et al. (39) used a knock-in mouse encoding one copy of the Egfp reporter gene inserted into the Rorc locus (41), whereas we used a transgenic mouse encoding 5–10 copies of a BAC carrying a similar Egfp insertion and therefore induces a brighter reporter expression. The more extended distribution of RORγt+ IL-17–producing T cells, in particular in the LNs and spleen, might better explain the occurrence of Th17-dependent inflammatory disease in the brain or joints of mice enduring experimental autoimmune encephalomyelitis or CIA (27–29, 39).

Foxp3+ RORγt+ Tαβ cells were an important fraction of RORγt+ T cells. They produced the immunosuppressive IL-10 and functioned as genuine T reg cells in vitro. Furthermore, these cells, but not IL-17–producing RORγt+ T cells, also expressed high levels of the transcript coding for CCL20. This transcript was undetected in RORγt– T cells, suggesting that Foxp3+ RORγt+ T cells were the main source of CCL20 in T cells, even though it might be expressed by a small subset of RORγt+ T cells. In accordance with our results, McGeachy et al. (13) have reported that CD4+ T cells

![Figure 6](image-url)
**Figure 7.** Generation and differentiation of RORγt+ Teβ cells in vitro. MACS-sorted naïve (CD62L+) CD4+ T cells from the spleens of Rorcγt−/Gfp− mice were stimulated in duplicates with anti-CD3ε and anti-CD28 in the presence of blocking anti–IFN-γ and anti–IL-4 antibodies and the indicated cytokines or RA. After different periods of time, cells were restimulated with PMA/ionomycin for 5 h and analyzed by flow cytometry for the expression of GFP, Foxp3, IL-17, and IL-10. All plots are gated on TCR-β+ cells, except plots for IL-10 that are gated on GFP+TCR-β+ cells. Numbers indicate percent cells in quadrants. Data are representative of three independent experiments.

cultured in the presence of TGF-β and IL-6, which produce both IL-17 and IL-10, express CCL20, in contrast to cells cultured with IL-23 alone, which produce only IL-17. CCL20 is the ligand for CCR6, suggesting that Foxp3+ RORγt+ T cells regulate the recruitment of CCR6+ T cells. Th17 cells isolated from SKG mice, which spontaneously develop arthritis, as well as human Th17 cells, express CCR6 (44, 48, 49). CCR6 is also expressed by a subset of "memory-like" Th17 cells in the lung (45) and by IL-17R in inflamed joints (46), as well as elevated levels of the fractalkine gene Cx3cl1. The functional consequences of elevated expression of Ios, Mmp9, and Cx3cl1 by Foxp3+ RORγt+ T cells remain to be explore, but this expression pattern appears paradoxical as the product of these genes might promote rather than modulate inflammation through the recruitment and activation of effectors, as well as through tissue remodeling through MMP-9. It might be suggested that the role of Foxp3+ RORγt+ T cells is not merely to regulate inflammation through IL-10, but also to partici- rate in the recruitment of inflammatory cells such as Th17 cells to an immunosuppressive environment. On the other hand, we cannot formally exclude that Foxp3+ RORγt+ T cells contain distinct functional subsets of IL-10–producing and/or CCL20–, ICOs–, MMP–, and CX3CL1–expressing cells.

Of note, 30–50% of RORγt+ Teβ cells did not express IL-17, IL-10, or Foxp3. We could not purify these cells from IL-17 producers for transcript analysis, as no markers were found to distinguish these cells before intracellular cytokine staining. It is possible that non–IL-17 and non–IL-10 producers contained uncommitted RORγt+ T cells, i.e., not yet committed to either the Th17 or the T reg cell pathway. They could also represent a third committed subpopulation that remains to be characterized. In favor of the first hypothesis, only few IFN-γ– and no IL-4–producing cells (not depicted) could be detected in RORγt+ T cells.

RORγt+ Tγδ cells have been reported previously to represent 50% of total Tγδ cells in the lamina propria of the small intestine, a majority of them expressing IL-17 (39). In the CIA model, 60–80% of Vγ4+ Tγδ cells and an equivalent number of CD4+ Teβ cells produce IL-17 in the draining LNIs (58). Furthermore, Tγδ cells are a primary source of IL-17 in the lung before and during infection with *Mycobacterium tuberculosis* (59) or *Mycobacterium bovis* (60), and Vβ1+ Tγδ cells produce IL-17 during intraperitoneal infection with *Escherichia coli* (61). In accordance with these data, we found...
that RORγT+ Tγδ cells were a major source of IL-17 in the lung and skin, as >80% of them produced IL-17, representing half of the total RORγT+ T cell population in these compartments. RORγT+ Tγδ cells expressed no Foxp3, but produced the highest levels of RORγT, as well of Il12r and cIf2 coding for GM-CSF. The latter induces the differentiation of granulocytes and monocytes, as well as the production of proinflammatory cytokines such as TNF-α and IL-6 (62). In contrast to RORγT+ Tαβ cells, IL-17–producing RORγT+ Tγδ cells were not affected by the absence of IL-6 in deficient mice, indicating that RORγT+ Tγδ cells and IL-17–producing RORγT+ Tαβ cells are generated by distinct pathways. Collectively, these data suggest that RORγT+ Tγδ cells play an important role in inflammation to pathogens through the production of IL-17 and GM-CSF; however, their precise role in vivo remains to be assessed.

RORγT is required for the generation of Th17 cells and has been shown to induce Th17 cells when transduced into mouse naive CD4+ T cells (39). Furthermore, naive CD4+ T cells cultured in the presence of TGF-β and IL-6 or IL-21 generate Th17 cells (4–6, 10–12) and IL-17–producing RORγT+ T cells (Fig. 4). We find that a large fraction of RORγT+ Tαβ cells expressed Foxp3, in particular in the LNs, as well as the immunosuppressive IL-10, and functioned as T reg cells. It has been shown previously that culture conditions that favor the generation of Th17 cells from naive CD4+ T cells, i.e., in the presence of TGF-β and IL-6, also induce the generation of IL-10–producing cells (13, 17). But in contrast to the in vivo situation, IL-10 was produced by Foxp3+ rather than Foxp3+ RORγT+ T cells, indicating that these culture systems might not faithfully replicate in vivo pathways. Interestingly, RA blocked the generation of IL-10–producing cells, whereas it favored the generation of Foxp3+ RORγT+ T cells. Furthermore, cultures of CD4+ T cells in the presence of TGF-β and IL-6 or IL-21 generated RORγT+ T cells expressing both IL-17 and IL-10, as observed previously (13, 17). However, RORγT+ T cells producing both cytokines could not be detected in vivo, exposing potential conflicts between in vivo and in vitro differentiation pathways.

Most cells cultured in the presence of TGF-β alone expressed RORγT after 1–2 d, but then differentiated into Foxp3+ RORγT+ T cells that did not express IL-17 after 2–3 d. It has been previously reported that TGF-β alone induces significant levels of Rori(γT) transcripts but no detectable Il17 or Il23r transcripts (4–6). These data indicate that RORγT is necessary but not sufficient for the generation of Th17 cells from naive CD4+ T cells, in accordance with the observation that cells cultured in the presence of TGF-β and IL-6 express RORγT after 1–2 d, but express IL-17 only after 2–3 d of culture (Fig. 4) (39). Collectively, these data indicate that TGF-β is opening a RORγT-dependent differentiation pathway in CD4+ T cells that leads either to Th17 or T reg cells, depending on the presence of matura-

The ratio of Th17 to T reg cells within RORγT+ T cells varied between organs, with the highest proportions of Th17 cells in the intestine and lung, and of Foxp3+ RORγT+ T cells in the LNs. This ratio was tuned in favor of Th17 cells by IL-6 and IL-23, as the ratio of IL-17–producing to Foxp3+ RORγT+ T cells decreased in IL-6– or IL-12Rβ1-deficient mice. However, in IL-6–deficient mice, the total number of RORγT+ Tαβ cells was also significantly diminished, indicating that IL-6 is involved more generally in the generation of RORγT+ Tαβ cells, as observed previously (6, 39). Conversely, the ratio of Th17 to T reg cells within RORγT+ T cells was tuned in favor of T reg cells by Foxp3 and CCL20, as inferred from scurfy mice or CCR6–deficient mice. Foxp3 might regulate the fate of “uncommitted” RORγT+ T cells into IL-17– or IL-10–producing cells by directly modifying the molecular function of RORγT, as Foxp3 was coimmunoprecipitated with RORγT. Foxp3 might also act through Foxp3+ RORγT+ T cells or “mainstream” Foxp3+ T cells regulating RORγT+ T cells. Interestingly, cells derived from scurfy mice generated a decreased proportion of RORγT+ Tαβ cells and IL-17–producing RORγT+ Tαβ cells, indicating a more complex role for Foxp3 in the generation and regulation of RORγT+ Tαβ cells that remains to be elucidated.

A robust mechanism maintains an equilibrium between Th17 to T reg cells within RORγT+ T cells. During massive inflammation induced by DSS in the intestine, the total number of RORγT+ T cells increased 10-fold in the intestine and >10-fold in influenza A virus–infected lungs. However, the numbers of IL-17– and IL-10–producing (Foxp3+) RORγT+ T cells were increased in comparable proportions. Regulating IL-17 versus IL-10 production might promote inflammation while limiting “collateral” damage, a necessary compromise between effective immunity and tissue integrity. IL-10–deficient mice, which lack one of these effector arms, generate increased proportions of IL-17–producing CD4+ T cells and develop spontaneous colitis in an IL-23–dependent pathway (15). In contrast, Apemin/+ mice that developed large colon and ileal polyps and tumors by 3–4 mo of age generated an increased proportion of IL-10–producing RORγT+ T cells. Similar observations have been made in mice injected with the B16 melanoma, which develop an increased ratio of Foxp3+ cells versus IL-17–producing cells as early as 8 d after tumor injection (53). The mechanisms used by tumors to alter the equilibrium between IL-17– and IL-10–producing RORγT+ T cells remain to be investigated, but this phenomenon is reminiscent of the IL-10–mediated immunosuppression induced by diverse viruses and bacteria (63). In contrast, the equilibrium between IL-17– and IL-10–producing RORγT+ T cells might be disrupted in favor of Th17 cells in chronic inflammatory diseases such as rheumatoid arthritis, multiple sclerosis, or inflammatory bowel disease. Tools to control the fate of RORγT+ T cells might therefore prove useful either to boost or to limit IL-17–producing cells. Potential targets include RORγT as well as any factors that regulate the differentiation of RORγT+ T cells into IL-17– or IL-10–producing effectors.
MATERIALS AND METHODS

Mice. BAC-transgenic Rorγt-Gfp and Rorγt-Cre mice were generated as described previously (40). The coding sequence for EGFP or Cre, including the stop codon, was inserted into exon 1 of Rorc in place of the endogenous ATG translation start codon, on a 200-kb BAC (Invitrogen) carrying at least 70 kb of sequence upstream of the Rorc translation start site. Transgenic mice used in transfer experiments were backcrossed eight times to C57BL/6 mice. CCR6-deficient (50), mice used in transfer experiments were backcrossed eight times to C57BL/6.

In vitro T cell differentiation. Spleen and mesenteric LN cells from Rorc( sf )×Gfp mice were pooled and T cells were purified using a CD4 + T cell isolation kit (Miltenyi). Purified cells were then mixed in a 1:1 ratio with lin− bone marrow cells from CD45.1 + C57BL/6 mice, and a total of 4 × 105 was injected into each recipient. 250–500 pg of high quality total RNA was subjected to one linear mRNA amplification cycle using the MessageBooster kit for quantitative RT-PCR (Epigen Bioscience Corporation). Real-time PCR was performed on a PTC-200 thermocycler equipped with a Chromo4 detector (Bio-Rad Laboratories). Data were analyzed using the Bio-Rad Laboratories.

Immunofluorescence histology. Tissues were washed and fixed overnight at 4°C in a fresh solution of 4% parafomaldehyde (Sigma-Aldrich) in PBS. The samples were then washed for 1 d in PBS, incubated in a solution of 30% sucrose (Sigma-Aldrich) in PBS until the samples sank, embedded in OCT compound 4583 (Sakura Finetek), frozen in a bath of isopentane cooled with liquid nitrogen, and stored at −80°C. Frozen blocks were cut at 8-μm thickness, and sections were collected onto SuperFrost/Plus slides (VWR). Slides were dried for 1 h and processed for staining or stored at −80°C. For staining, slides were first hydrated in PBS-XG (PBS containing 0.1% Triton X-100 and 1% normal goat serum; Sigma-Aldrich) for 5 min and blocked with 10% bovine serum in PBS-XG for 1 h at room temperature. Endogenous biotin was blocked with a biotin blocking kit (Vector Laboratories) and slides were washed and incubated with the indicated mAb conjugates for 40 min in a total volume of 100 μl PBS-F. For intracellular cytokine staining, cells were stimulated for 10 h in the presence of 10 ng/ml PMA, 1 μg/ml ionomycin, 0.5 μg/ml brefeldin A (Sigma-Aldrich), and 2 μg/ml monensin (eBioscience) to block intracellular cytokine staining. Cells were incubated with a GolgiPlug kit (BD Biosciences) for 1 h and then stained with mAbs specific for intracellular cytokines. Stained cells were then washed and stained with a Fix & Perm kit (eBioscience) and cells were analyzed on a FACSCanto II (BD Biosciences). FlowJo software (Treestar).
into Rag2−/−Il2rb−/− mice (64) irradiated at 600 rad. After 4 wk, reconstitution of the mice was assessed in peripheral blood. Reconstituted mice were killed and analyzed 6 wk after transfer.

Online supplemental material. In Fig S1, the purification of the different subsets of RORγt− T cells is shown. Fig. S2 shows that RORγt− T cells express genes involved in the Th17 or Th2 cell pathway. In Fig. S3, expression of IL-10 by intestinal RORγt− Th2 cells is depicted. Fig. S4 shows the expression of CCR6 by RORγt− Th2 cells. Fig. S5 shows the generation of mixed bone marrow chimeras, and Fig. S6 shows the generation of Foxp3+ T cells in mice a/b of RORγt− cells. In Fig. S7, immunofluorescence histology of terminal ileum from a WT or a 4-mo-old Rorc(ΔF)-Cgr (G) mouse developing polyps and carcinoma is shown. Figs. S1–S7 are available at http://www.jem.org/cgi/content/full/jem.20080034/DC1.

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