IL-10 signaling in CD4+ T cells is critical for the pathogenesis of collagen-induced arthritis

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Abstract

Introduction: IL-10 is a very important anti-inflammatory cytokine. However, the role of this cytokine in T cells in the pathogenesis of collagen-induced arthritis is unclear. The purpose of this study was to define the role of IL-10 signaling in T cells in the pathogenesis of collagen-induced arthritis.

Methods: IL-10 receptor dominant-negative transgenic (Tg) and control mice were immunized with bovine type II collagen to induce arthritis. The severity of arthritis was monitored and examined histologically. T-cell activation and cytokine production were analyzed using flow cytometry. T-cell proliferation was examined by [3H]thymidine incorporation. Antigen-specific antibodies in serum were measured by ELISA. Foxp3 expression in CD4+ regulatory T cells (Tregs) was determined by intracellular staining or Foxp3-RFP reporter mice. The suppressive function of Foxp3+CD4+ Tregs was determined in vitro by performing a T-cell proliferation assay. The level of IL-17 mRNA in joints was measured by real-time PCR. A two-tailed nonparametric paired test (Wilcoxon signed-rank test) was used to calculate the arthritis and histological scores. Student’s paired or unpaired t-test was used for all other statistical analyses (InStat version 2.03 software; GraphPad Software, San Diego, CA, USA).

Results: Blocking IL-10 signaling in T cells rendered mice, especially female mice, highly susceptible to collagen-induced arthritis. T-cell activation and proliferation were enhanced and produced more IFN-γ. The suppressive function of CD4+Foxp3+ regulatory T cells was significantly impaired in Tg mice because of the reduced ability of Tregs from Tg mice to maintain their levels of Foxp3. This was further confirmed by transferring Foxp3-RFP cells from Tg or wild-type (Wt) mice into a congenic Wt host. The higher level of IL-17 mRNA was detected in inflammatory joints of Tg mice, probably due to the recruitment of IL-17+γδ T cells into the arthritic joints.

Conclusion: IL-10 signaling in T cells is critical for dampening the pathogenesis of collagen-induced arthritis by maintaining the function of Tregs and the recruitment of IL-17+γδ T cells.

Introduction

Rheumatoid arthritis (RA) is an autoimmune disease characterized by chronic inflammation of the joint capsule and synovial membrane which results in cartilage injury, bone erosion and eventually joint destruction and deformity [1]. Collagen-induced arthritis (CIA) is a well-established animal model that has been studied extensively because of its similarities to human RA [2]. Although the etiology of RA remains unknown, it has been reported that a functional imbalance between proinflammatory cytokines and regulatory T cells (Tregs) is a key mechanism that underlies joint inflammation and disease progression in CIA as well as RA [3].

IL-10 is a pleiotropic cytokine with important immune-regulatory functions [4]. It has been implicated to have a potent anti-inflammatory role in several autoimmune disease models, including CIA [5]. IL-10 suppresses the expression of inflammatory cytokines such as TNF-α, IL-6 and IL-1 by activated macrophages [6]. IL-10 also affects T-cell proliferation and cytokine production [7,8]. Indeed, IL-10 affects many of the cell types in the immune system; however, the precise role of IL-10 signaling in CD4+ T cells in the pathogenesis of CIA has not been addressed.
It has been demonstrated in both animal model and human studies that naturally occurring CD4+CD25+Foxp3+ Tregs play a critical role in the prevention of autoimmunity and inflammatory arthritis [9,10]. Depletion of CD4+CD25+ T cells aggravated CIA [11,12], whereas transferring CD4+CD25+ cells to a disease-bearing animal ameliorated arthritis [12]. Although it has been well-documented that IL-10 can induce differentiation of naive CD4+ T cells into CD4+IL-10+ Tr1 cells [13,14], it is unclear whether IL-10 signaling in T cells affects the development or function of these Tregs.

Recently, a new subset of IL-17-producing CD4+ T cells, also called Th17 cells [15,16], has been implicated as an important mediator in tissue inflammation [17]. IL-17-deficient mice showed markedly suppressed CIA [18]. Resistance to CIA in p19−/− mice correlated with an absence of IL-17-producing CD4+ T cells, suggesting that the IL-23-IL-17 axis rather than the IL-12-IFN-γ axis is essential in promoting the development of CIA [19]. It has been reported that IL-10 suppresses Th17 cytokine secretion by macrophages and T cells in in vitro culture [20]. However, the role of IL-10 signaling in T cells in the differentiation of Th17 cells and how this regulation affects the pathogenesis of CIA is less clear.

In this study, in which we studied previously described IL-10 receptor dominant-negative transgenic (Tg) mice [21], we showed that when T cells fail to respond to IL-10, mice develop more severe arthritis and T cells are more activated and proliferate more against type II collagen (CII) antigen. Moreover, the suppressive function of Tregs in these Tg mice was significantly impaired as a result of attenuated expression of Foxp3. IL-10 signaling in CD4+ T cells might affect the cytokine profiles of CD4+ T cells as well as γδ T cells. These results indicate that IL-10 signaling in T cells is the key to the pathogenesis of CIA.

Materials and methods
Mice
DBA/1 and C57BL/6 mice (Thy1.2 or Thy1.1 background) were purchased from The Jackson Laboratory (Bar Harbor, ME, USA) and maintained under specific pathogen-free conditions at the Yale University School of Medicine. DBA/1 background IL-10 receptor dominant-negative Tg mice were obtained by backcrossing C57BL/6 Tg mice [21] with wild-type (Wt) DBA/1 mice for nine generations. The H-2kq and Tg genotypes were screened by PCR using tail DNA. The IL-10 GFP knockin mouse, designated IL-10-internal ribosomal entry site (IRES)-GFP-enhanced reporter (Tiger), and the Foxp3 red fluorescent protein (RFP) knockin mouse, designated Foxp3-IRES-mRFP (FIR) mice, were described previously [22,23]. All the animal procedures were performed with the approval of the Institutional Animal Care and Use Committee of the Yale University School of Medicine.

Reagents
Bovine CII for immunization, T-cell proliferation and ELISA, as well as Complete Freund’s Adjuvant (CFA) and Incomplete Freund’s Adjuvant (IFA), were all purchased from Chondrex (Redmond, WA, USA). BD GolgiPlug and antibodies for flow cytometry were purchased from BD (San Diego, CA, USA). A Foxp3 intracellular staining kit was purchased from eBioscience (San Diego, CA, USA). Cytokines and antibodies for T-cell differentiation in vitro were obtained from R&D Systems (Minneapolis, MN, USA). Phorbol 12-myristate 13-acetate (PMA) and ionomycin were purchased from Sigma-Aldrich (St Louis, MO, USA).

Collagen-induced arthritis model
The CIA model was employed as described previously [24]. Briefly, 100 μg of bovine CII dissolved in 0.1 M acetic acid was emulsified with 100 μl of CFA containing 2 mg/ml inactivated Mycobacterium tuberculosis. Mice were sensitized by intradermal injection at the base of the tail and boosted at day 21 with the same dose of CII and IFA intradermally. Mice were scored blindly for disease severity every other day after the boost using a grading scale from 0 to 3 that was described previously [24]. Each paw was graded with a maximum score of 12 per mouse.

Cell proliferation
Splenocytes from CIA mice were cultured with different concentrations of CII for 3 days. After pulsing with 1 μCi of [3H]thymidine per well for the last 18 hours, proliferation was measured as radioactivity incorporation (counts per minute (cpm)).

Activation markers
Splenocytes from CIA mice were stained with anti-CD3, anti-CD4, anti-CD8, anti-CD44 and anti-CD62L antibodies, and the surface markers for activation in CD4+ and CD8+ T cells were analyzed using an LSR II flow cytometer (BD).

ELISA for serum anti-CII antibodies and their isotypes
Serum samples were collected from different groups of immunized mice at different days postimmunization (days 0, 14, 28, 35 and 42), the levels of anti-CII antibody in different isotypes were measured by ELISA as described in our previous studies [25]. Briefly, ELISA plates were coated with bovine CII (0.5 μg/well). After the plates were blocked with 5% fetal bovine serum-PBS, 1:800 diluted (with blocking buffer) mouse sera
were added to duplicate wells and incubated for 2 hours at room temperature. The plates were then washed, and biotin-conjugated goat anti-mouse immunoglobulin G (IgG), IgG1 and IgG2a were added at a dilution of 1:2,000 for 1 hour, followed by the addition of streptavidin-horseradish peroxidase (Sigma-Aldrich). The plates were then developed, and antibody levels were measured as described previously [24].

Quantitative real-time PCR
Total RNA was isolated from joints by using the RNeasy Mini Kit (QIAGEN, Valencia, CA, USA). Genomic DNA was removed by digestion with DNase I. cDNA was synthesized using the iScript cDNA Synthesis Kit (Bio-Rad Laboratories, Hercules, CA, USA), and gene expression was examined using the Stratagene Mx3005P QPCR System with Brilliant SYBR Green QPCR Master Mix reagent (both from Agilent Technologies, Inc, Santa Clara, CA, USA). The data were normalized to a gapdh reference [26]. The forward and reverse primers for il17 and gapdh were as follows [27]: il17, TTTAATCCCTTGCCTGCAAAA, CTTTCTCCTCCTGCATTGACAC; and gapdh, TTCACCACCATGGA-GAAAGC, GGCATGGACTGTGGTCATGA.

Intracellular Foxp3 and cytokine staining
Single-cell suspensions were prepared from spleens, lymph nodes or thymi and stained with surface CD4 and CD25. After being fixed with freshly prepared fixation/permeabilization solution at 4°C for at least 30 minutes in the dark, cells were blocked with Mouse BD Fc Block (BD) in permeabilization buffer at 4°C for at least 30 minutes in the dark, cells were blocked with Mouse BD (IgG), IgG1 and IgG2a were added at a dilution of 1:2,000 for 1 hour, followed by the addition of streptavidin-horseradish peroxidase (Sigma-Aldrich). The plates were then developed, and antibody levels were measured as described previously [24].

Suppression assay
Enriched CD4+ T cells were sorted into a CD4+ CD25+ subset as the regulatory cells and a CD4+ CD25- subset as the responder cells. T-cell-free splenocytes treated with mitomycin C were used as antigen-present cells (APCs). The responder cells were cultured with the regulatory cells at ratios of 1:1, 1:0.5, 1:0.25 and 1:0.125 or without the regulatory cells as the positive control in the presence of brefeldin A for 4 hours. Cytokine production was measured as described previously.

Adoptive transfer
CD4+RFP+ Tregs were sorted from the spleens of B6 Thy1.1 IL-10 dominant-negative receptor Tg and Wt mice and were transferred into B6 Thy1.2 Wt mice at equal cell numbers. One day after transfer mice were immunized with CFA and bovine CII as described above. One week after immunization Foxp3 expression in transferred cells was analyzed by checking the levels of RFP.

Histopathologic assessment
On day 58 after CIA induction, mice were killed and their paws were dissected free and used for histologic examination. The joints were immediately fixed in 10% buffered formalin and decalcified in a decalcifying solution for about 3 to 5 days [24]. The tissues were then processed and embedded in paraffin. Five-micrometer tissue sections were prepared and stained with H & E using standard methods. All sections were stained at the Yale Pathology Laboratory. Sections were blindly observed and scored by histologists. Five arthritis severity factors were assessed using a grading scale from 0 (normal) to 5 (severe), based on a previously described scoring system [28].

Statistical analysis
Data presented are means ± SEM. Statistical testing was two-tailed with a 5% level of significance. A two-tailed nonparametric paired test (Wilcoxon signed-rank test) was used to analyze the arthritis and histologic scores. Student’s paired or unpaired t-test was used for all other statistical analyses (InStat version 2.03 software; GraphPad Software, San Diego, CA, USA).

Results
Blocking IL-10 signaling in T cells rendered mice highly susceptible to collagen-induced arthritis
To define the role of IL-10 signaling in T cells in the pathogenesis of CIA, we employed newly generated IL-10 receptor dominant-negative Tg mice in which T cells expressed high levels of Tg IL-10Ra that were truncated to remove its signaling domain [21]. Tg mice were backcrossed with DBA/1 mice 10 times to develop Tg mice on the CIA-susceptible background (DBA/1, H-2b). Four groups of DBA/1 mice (male Wt or Tg and female Wt or Tg) from the same cohort of littersmates at age 8 weeks were immunized with bovine CIA to induce arthritis using the standard CIA protocol described in our previous studies. It has been observed that even in the susceptible strain of mice, such as DBA/1 (H-2b), female mice are more resistant than male mice, with a lower incidence and less severity of CIA [28]. As expected, our experiment showed that male mice were highly susceptible to and females
relatively resistant to CIA (Figure 1A). All the male Wt mice developed progressive disease (with 100% incidence) and had a mean arthritis score of 8.25 ± 2.88 (n = 10) on day 35 postimmunization. In contrast, all the female Wt mice developed visible arthritis (also 100% incidence) with a significantly lower mean arthritis score of 5.00 ± 4.10 (n = 12) on day 35 postimmunization. Interestingly, female Tg mice developed severe progressive disease and had a mean arthritis score of 8.78 ± 2.33 (n = 10), slightly higher than those parameters in male Wt mice and male Tg mice (7.89 ± 2.49, n = 12). Further evidence in support of increased arthritis severity in female Tg mice was obtained by histopathologic analysis. On day 50 postimmunization, female Tg mice exhibited robust pannus formation and significant articular cartilage erosion, with pannus eroding and replacing the articular cartilage overlying the bone. In contrast, Wt female mice exhibited well-preserved joint spaces and articular cartilage surfaces, with minimal pannus formation. Among male mice, both Wt and Tg exhibited extensive pannus formation and destruction of bone and cartilage. One representative example is shown in Figure 1B. At 50 days postimmunization, the histologic score of Wt female mice was 3.8 ± 1.4 of a possible score of 25. All the other groups showed similar scores (Figure 1C) (female Tg, 13.7 ± 0.6; male Wt, 14.7 ± 0.8; male Tg, 13.3 ± 1.0, respectively; n = 4 mice in each group). Because the major difference in CIA disease severity was observed between female Wt and Tg mice rather than male Wt and Tg mice, only female mice were used for the rest of our study. Thus our data suggest that IL-10 signaling in T cells plays a critical role in suppressing the pathogenesis of CIA.

Figure 1 IL-10 receptor dominant-negative transgenic mice were highly susceptible to collagen-induced arthritis (CIA). (A) Sex- and age-matched DBA/1J transgenic (Tg) and wild-type (Wt) littermate control mice were immunized with bovine type II collagen as described in “Materials and methods.” The clinical arthritis score was observed, scored and recorded daily. Left (Wt males, n = 10, Tg males, n = 9, P > 0.05). Right (Wt females, n = 12, Tg females, n = 13; P < 0.01). The results of one of three repeated experiments are shown. (B) Representative H & E-stained sections of paw joints from Wt and Tg mice on day 50 postimmunization in CIA animals are shown (original magnification, ×10; scale bar, 100 μm). (C) Histological scores of paw joints on day 50 postimmunization. Each feature of arthritis (joint space exudate, synovitis, pannus formation and cartilage and bone degradation) was graded from 0 (normal) to 5 (severe). The values given are means and SEM (n = 4 per group; females, P < 0.01, and males, P > 0.05). The results are representative of two experiments.
IL-10 signaling in T cells controls T-cell activation and proliferation responses in collagen-induced arthritis mice

Activation and cytokine production of CD4+ T cells play an important role in the pathogenesis of CIA. To define the underlying mechanisms of increased arthritis severity in female Tg mice, T-cell activation and proliferation in CIA mice were first analyzed. Cells from draining lymph nodes on day 50 postimmunization were stained with anti-CD3, anti-CD4, anti-CD8, anti-CD44 and anti-CD62L antibodies. After gating on CD3+ T cells, the CD44hi T cells were recognized as activated cells. Activated CD4+ was found to be significantly higher in Tg mice than in Wt mice (Figure 2A, left), whereas activated CD8+ in Tg and Wt mice appeared to be comparable (Figure 2A, right), implying that IL-10 signaling in T cells controlled CD4+ T cell activation in inflammatory response. One representative dot plot is shown (Figure 2B). To test the impact of IL-10 signaling in antigen-specific T-cell proliferative response in immunized mice, cells recovered as described above were cultured in 96-well U-bottom plates in the presence of different concentrations of CII for 3 days, followed by pulsing with 1 μCi of [3H]thymidine per well for the last 18 hours, and proliferation was measured as radioactivity incorporated (in cpm). Cells from Tg mice showed significantly increased proliferative responses to specific antigen (Figure 2C), suggesting a critical role of IL-10 signaling in T cells in controlling lymphocyte cell proliferation in CIA. To further test whether IL-10 signaling in T cells would affect T-cell differentiation in CIA, we also analyzed IFN-γ-producing CD4+ T cells in draining lymph nodes upon immunization and found elevated percentages of IFN-γ+CD4+ cells in Tg mice (Figure 2D), suggesting that IL-10 signaling suppressed the

![Figure 2](http://arthritis-research.com/content/13/6/R212)

**Figure 2** Altered T-cell phenotype in IL-10 dominant-negative receptor transgenic mice upon induction of collagen-induced arthritis

(A) Sex- and age-matched DBA/1 transgenic (Tg) and wild-type (Wt) littermate control mice were immunized with type II collagen as described in “Materials and methods.” Cells from draining lymph nodes in female DBA/1 mice on day 28 postimmunization were stained with anti-CD3, anti-CD4, anti-CD8, anti-CD44 and anti-CD62L expression after gating on CD4+ (left) or CD8+ T cells (right). Wt, n = 3, Tg, n = 3; *P < 0.05. (B) One representative stain of CD44, CD62L in CD3+CD4+ (left) and CD3+CD8+ (right) T cells of three repeated experiments is shown. (C) The same sources of cells described above were cultured with different concentrations of collagen for 3 days. After pulsing with 1 μCi of [3H]thymidine per well for the last 18 hours, proliferation was measured as radioactivity incorporation in counts per minute (cpm). Stimulation Index = cpm experiment/cpm control (concentration of collagen = 0). The results of one representative experiment are shown. (D) The cells were also stimulated with phorbol 12-myristate 13-acetate (50 ng/ml) and ionomycin (500 ng/ml) in the presence of BD GolgiPlug for 4 hours then stained with CD4 and intracellular IFN-γ. The statistical analysis for the percentage of IFN-γ+ cells in the CD3+CD4+ gate is shown (Wt, n = 3, Tg, n = 3; P < 0.05). One representative stain is shown (E).
pathogenic Th1 response. One representative dot plot is shown (Figure 2E). Interestingly, no significant differences were observed in terms of the numbers and the activation states of γδ T cells in the spleens and draining lymph nodes of Wt and Tg mice (data not shown).

**Impaired suppressive function of regulatory T cells from transgenic mice**

It has been demonstrated that CD4⁺Foxp3⁺ Tregs play important roles in the suppression of CIA in animal models and human studies [9,10]. To define the role of IL-10 signaling in T cells in the development of naturally occurring Tregs (nTregs), the percentage and absolute number of CD4⁺CD25⁺Foxp3⁺ T cells in DBA/1 and B6 Tg and Wt mice were analyzed. Very similar percentages and numbers of CD25⁺Foxp3⁺ T cells were found in both Wt and Tg mice (Figures 3A and 3B), indicating that IL-10 signaling in T cells was not required for the development of nTregs. Similar results were observed in DBA/1 mice upon immunization (data not shown). To further define the function of Tregs, a standard in vitro T-cell suppression assay was performed. CD4⁺CD25⁺ T cells were sorted from B6 Wt or Tg mice as Tregs and CD4⁺CD25⁻ T cells from B6 Wt mice as responders. Tregs from both Wt and Tg mice were anergic to TCR activation (data not shown). Interestingly, the suppressor function of Tregs from Tg mice was significantly impaired in comparison to those from Wt mice (Figures 3C and 3D). To rule out the contamination of activated CD4⁺ T cells in the Treg population when using CD25 as the marker of Tregs, RFP-Foxp3 reporter mice [23] were used to sort Foxp3⁺ regulators and Foxp3⁻ responders and the suppress assay mentioned above was repeated. Similar results were obtained (Figure 3E). Our results demonstrate that IL-10 signaling in T cells is not required for Treg development, but rather for T cells’ suppressive function.

**IL-10 signaling in T cells is required for maintenance of Foxp3 expression**

To define the mechanisms underlying the reduced suppressive function of Tregs in Tg mice, we first analyzed IL-10 production in Tg mice. For this purpose, we used IL-10-Foxp3 double-reporter mice [22,23]. The percentage of IL-10-expressing cells was similar between Wt and Tg mice in both immunized and immunized states (Figures 4A and 4B), indicating that IL-10 signaling in T cells was not required for IL-10 production by Tregs and that IL-10 was not related to the impaired suppressive function of Tg Tregs. We next analyzed the transforming growth factor (TGF)-β production by Tregs in Wt and the Tg mice. Lymphocytes from Wt and the Tg mice were stimulated with PMA and ionomycin for 4 hours in the presence of brefeldin A, and intracellular Foxp3 and TGF-β were stained in the CD4⁺ T cell population. No significant differences in TGF-β production between Tregs from Wt and the Tg mice were detected (Figure 4C), suggesting that IL-10 signaling in T cells was not involved in TGF-β production. We then investigated whether IL-10 signaling in T cells would affect CTLA-4 expression level in Tregs. Cells from the lymph nodes of unimmunized and immunized Wt and the Tg mice were stained with Foxp3 and intracellular CTLA-4. CTLA-4 was highly expressed in CD4⁺Foxp3⁺ T cells, but not in CD4⁺Foxp3⁻ T cells; however, no significant differences were found in the levels of CTLA-4 in Tregs from Wt and Tg mice (Figure 4D). Because Foxp3 has been reported to be the master transcription factor controlling Treg development and determining the suppressive function of Tregs [29-31], we studied whether IL-10 signaling played any role in maintaining the stable expression of Foxp3. Similarly to the results shown in Figure 3A, the percentage of Foxp3⁺ T cells was very similar between Wt and Tg mice (Figure 4A, left). Interestingly, it decreased by 30% upon immunization (Figure 4A, right, and Figure 4E). A similar phenomenon was observed in DBA/1 mice using intracellular staining (data not shown). To define whether IL-10 signaling was critical for the maintenance of Foxp3 expression in Tregs, CD4⁺RFP⁺ T cells from Wt and Tg Thy1.1 mice were sorted and transferred to Thy1.2 Wt mice, followed by immunization with CII and CFA. One week later the transferred cells were analyzed for expression of RFP. Transferred Foxp3⁺ cells were observed to become RFP⁻ cells in both Tg and Wt Tregs (Figures 4F and 4G). However, Tregs from Tg mice showed significantly higher percentages of RFP⁻ cells than those from Wt mice, indicating that IL-10 signaling in Tregs was critical to maintaining the level of Foxp3 expression upon immunization.

**Increased IL-17⁺ γδ T cells and joint IL-17 expression in transgenic mice**

IL-17 is a critical cytokine in the pathogenesis of CIA [32]. To define the role of IL-10 signaling in T cells on IL-17 production by CD4⁺ T cells, we first compared the IL-17⁺CD4⁺ T cells from unimmunized mice. We found that Tg mice had increased Th17 cells in both peripheral and central lymph organs (Figure 5A), suggesting that IL-10 signaling negatively regulates naturally occurring Th17 cell development in steady state. To further define the role of IL-10 in Th17 cell differentiation, we sorted naïve CD4⁺ T cells (CD4⁺CD44⁺CD62L⁺ cells) from Tg as well as IL-10-knockout (IL-10Ko) mice versus Wt controls cultured under Th17 differentiation conditions in the presence or absence of IL-10, and the cells were then used to analyze cytokine production. No significant difference was found in terms of the percentage of IL-17-producing cells among Tg,
Ko and Wt mice with or without IL-10, indicating that IL-10 signaling had no effect on inducible Th17 cell differentiation (Figure 5B). To define the role of IL-17 in local inflammatory response, the level of IL-17 mRNA (il17) in the joints of CIA mice was analyzed using real-time PCR. We found that joint tissues from Tg mice expressed significantly higher levels of il17 than Wt mice did (Figure 5C), which was consistent with the severity of arthritis. To define the potential sources of IL-17 in the joints, cells from draining lymph nodes of CIA mice (Wt and Tg mice) were analyzed for IL-17A production. Interestingly, significantly higher percentages of γδ T cells from Tg mice produced IL-17 than those from Wt mice (Figure 5D), especially CD44hi γδ T cells. In contrast, we did not find significant differences in the number of IL-17-producing CD4+ T cells (Figure 5E), suggesting γδ T cells as the primary source of IL-17 in the inflammatory joints.

Figure 3 Impaired suppressive function of regulatory T cells from transgenic mice. (A) Splenocytes from C57BL/6 mice were stained with anti-CD4 and anti-CD25 antibodies, which was followed by Foxp3 intracellular staining, as described in "Materials and methods." The dot plots of CD25+ and Foxp3+ proportions after gating on CD3+CD4+ T cells are shown. The results shown are from one of three representative experiments. (B) The absolute number of regulatory T cells (Tregs) (CD4+CD25+Foxp3+) cells from the spleens of C57BL/6 mice were calculated and are shown here (n = 3 mice for each group, P > 0.05). (C) CD4+ T cells from the spleens of B6 wild-type (Wt) or transgenic (Tg) mice were enriched and sorted into a CD4+CD25+ subset as the regulatory cells and a CD4+CD25− subset from Wt C57BL/6 as the responder cells. The number of responder cells was fixed at 5 × 10^4. The inhibition function of Tregs was measured by in vitro assay as described in "Materials and methods." The percentage inhibition by different populations of Tregs was calculated by using the formula described in "Materials and methods." (D) The results shown are from one of three representative experiments. (E) B6 Wt and Tg mice were crossed with Foxp3-RFP mice. CD4+ T cells from the spleens of Wt or Tg red fluorescent protein (RFP) mice were enriched and sorted into a CD4+RFP+ subset as the regulatory cells and a CD4+RFP− subset from Wt C57BL/6 as the responder cells. The responder cells (5 × 10^4/well) were cocultured with an equal number of regulatory cells for the inhibition assay described above. The results from one of three representative repeated experiments are shown.
IL-10 signaling in T cells has no effect on antigen-specific antibody response

Antibody production against CII is an important factor in the development of CIA [33]. To define the impact of IL-10 signaling in T cells on collagen-specific antibody response, serum samples were collected from naïve mice as well as from different groups of immunized mice at different time points after immunization (days 0, 14, 28, 35 and 42), and the levels of anticollagen antibodies were analyzed by ELISA as described in our previous studies [24]. The level of collagen-specific antibody in serum increased upon immunization and was maintained at high levels between days 14 and 42. Surprisingly, there was no difference in collagen-specific
Figure 5 Increased IL-17 mRNA expression in the joints of transgenic mice (A) Cells from the lymph nodes and thymi of untreated B6 wild-type (Wt) or transgenic (Tg) mice were stimulated with phorbol 12-myristate 13-acetate (50 ng/ml) and ionomycin (500 ng/ml) in the presence of BD GolgiPlug for 4 hours, followed by staining with TCRβ, CD4, CD8, CD44 and intracellular IL-17. One representative stain is shown (right). The statistical analysis of the percentage of IL-17+ cells in TCRβ+CD4+CD8−CD44hi cells (three mice in each group) is shown (left). (B) Naive CD4+ T cells (CD4+CD44loCD62Llo) from B6 Wt, IL-10-knockout (IL-10Ko) and Tg mice were sorted and stimulated with coated anti-CD3 and anti-CD28 antibodies under Th17 differentiation conditions in the presence or absence of IL-10 (100 μg/ml). Seven days later IL-17 production was measured by intracellular staining. The results of one of three representative experiments are shown. (C) The relative levels of IL-17 mRNA from the joints of DBA/1 Wt and Tg mice with collagen-induced arthritis (on day 50 postimmunization) were measured by real-time PCR. (D) Cells from draining lymph nodes of immunized C57BL/6 Wt or Tg mice (on day 7 postimmunization) were stimulated with the same procedure described above for IL-17 intracellular staining. One representative stain showing lymphocytes (left) or CD3+TCRβ+ T cells (middle) upon gating is shown. The statistical analysis for the percentage of IL-17+ cells in CD3+TCRβ+ T cells (three mice in each group) is shown (right). (E) The same population of cells described in (D) was also stimulated for IL-17 intracellular staining upon gating on lymphocytes (left) or CD4+ T cells (middle). Statistics analysis for the percentage of IL-17+ cells in TCRβ+CD4+CD8−CD44hi cells (three mice in each group) is shown (right).
antibodies, including the total IgG, IgG1 and IgG2a iso-
types, among Wt and Tg mice (Figure 6). These results
suggest that blocking IL-10 signaling in T cells has no
direct effect on collagen-specific antibody production by
B cells in a CIA model.

Discussion
The CIA has been used as a model for studying the
pathogenesis and potential application of novel therapeu-
tic agents for human RA. Both T and B cells are
essential for the development of RA [34,35], and IL-10
plays different roles in regulating the function of T and
B cells [36]. It is therefore important to study the
unique role of IL-10 signaling in T cells (without affect-
ing B cells) in the pathogenesis of CIA. In this study, we
demonstrated that IL-10 signaling in T cells play a
critical role in the pathogenesis of CIA by affecting the
function of Tregs by stabilizing the expression of Foxp3.

To target IL-10 signaling selectively in T cells, a
unique IL-10 receptor dominant-negative Tg mouse was
recently developed with the Tg under the control of
CD4 promoter, which lacks the CD8 silencer element
and therefore is expressed in both CD4+ and CD8+ T
cells [21]. In these mice, CD4+ and CD8+ T cells
expressed high levels of the Tg IL-10Ra, which lacks the
cytoplasmic domain. As this receptor functions as a
dominant-negative receptor, its responsiveness to IL-10
is attenuated in these T cells. Using these mice, we
demonstrated that Tg mice developed more severe
arthritis upon induction in the absence of IL-10 signal-
ing (Figure 1). This was especially true for female mice.
Even in susceptible strains, female Wt mice are more

Figure 6 IL-10 signaling in T cells has no effect on type II collagen-specific antibody production. Serum samples from DBA/1J Wt or Tg
mice (six mice in each group) were collected on days 0, 14, 21, 35 and 42 after collagen-induced arthritis induction, and collagen-specific
antibodies (immunoglobulin G (IgG), IgG1 and IgG2a) were detected by ELISA. The results of one representative experiment are shown.
resistant than males, and the underlying mechanisms have not been studied systematically. Our observation that blockade of IL-10 signaling in T cells rendered female mice highly susceptible to CIA may shed some light on the molecular mechanisms of the gender differences. Of course, we did not rule out the possibility that IL-10 signaling might also affect the male mice if these mice were immunized with lower doses of CII and/or subdoses of adjuvant. Further studies are needed to explore this possibility.

**CD4+ T cells play an essential role in the pathogenesis of CIA as well as RA.** Autoreactive T cells are normally controlled by Tregs and other regulatory mechanisms under normal circumstances, whereas in autoimmune diseases Treg dysfunction causes autoreactive T-cell proliferative responses and tissue damage [37]. As one of the key suppressive cytokines, IL-10 plays an important role in controlling T-cell responses. Indeed, though T cells were unable to respond to IL-10, they were more activated and proliferated more vigorously upon CIA induction, with a more pronounced Th1 response (Figure 2). Our data suggest that a potential mechanism for this enhanced T-cell response could be the dysfunction of Tregs. Although IL-10 signaling in T cells was not required for the development of nTregs (Figures 3A and 4A), we clearly have demonstrated that the suppressive function of Tregs was significantly impaired in the absence of IL-10 signaling (Figures 3C and 3D).

Several elements are critical for the function of Tregs, including IL-10, TGF-β and CTLA-4 [38]. To define which factor was critical for the function of Tregs in connection of IL-10 signaling in T cells, we analyzed the levels of IL-10, TGF-β and CTLA-4 in Tregs of Wt and the Tg mice. To our surprise, no significant differences were detected between these two populations of Tregs (Figure 4), implying that IL-10 signaling in T cells was not required for the expression of these factors.

Foxp3 is a critical factor that determines not only the development and differentiation but also the suppression function of Tregs. Expression of Foxp3 can be regulated by multiple factors, including inflammatory cytokines [39]. One of the striking findings of our study is that we have demonstrated that in the absence of IL-10 signaling in T cells, the expression levels of Foxp3 were reduced upon immunization, on the basis of both ex vivo studies (Figure 4A) and Foxp3-RFP T-cell adoptive transfer experiments (Figures 4F and 4G). This phenotype was observed in C57BL/6 mice as well as in DBA/1 mice, suggesting that Foxp3 downregulation in Tg mice upon immunization may be associated with the disease mechanism in CIA (data not shown). By taking advantage of our Foxp3-RFP reporter mice, we were able to isolate a pure population of Tregs and track these cells in the congeneric hosts. We speculated that upon immunization, inflammatory cytokines might target Tregs and regulate expression of Foxp3. One of the critical functions of IL-10 signaling in these Tregs probably was to maintain the expression of this key molecule. Therefore, our results collectively indicate that IL-10 signaling in T cells is critical for the function of Tregs by maintaining the expression level of Foxp3. Although Foxp3 is an X-linked gene, no significant difference was observed in the expression levels of Foxp3 between male and female mice (data not shown). Further studies are needed to define the molecular mechanisms underlying how the IL-10 signaling pathway maintains the expression of Foxp3.

IL-17 has been defined as a critical cytokine that mediates autoimmune inflammation, especially in CIA and RA [40]. This has been evidenced by a series of studies using different gene-deficient mice, including p19−/−, inducible costimulatory molecule (ICOS−/−) and IL-17−/− mice [18,19,40]. Indeed, the expression levels of IL-17A were found to be significantly increased in the joints of Tg mice, which correlated with severe arthritis (Figure 5). Interestingly, the increased levels of IL-17 were not derived from CD4+ T cells; rather, they were most likely derived from γδ T cells (Figures 5D and 5E). Our results are consistent with those of previous studies in that Vγ4 Vδ T cells provided IL-17A in the pathogenesis of CIA [41]. How does IL-10 signaling in CD4+ T cells affect the production or recruitment of IL-17A-producing γδ T cells is unclear at present. Several possibilities might explain our findings. IL-10 signaling promoted both thymic and peripheral CD4+ T cells to produce IL-17A in naïve, but not in immunized, mice (Figure 5A), indicating a possibility that IL-10 signaling might change the threshold of IL-17 expression by CD4+ T cells. The increased percentage of IL-17+ γδ T cells in the draining lymph nodes of CIA mice could be a result of more local differentiation, more recruitment from other tissues, or both. Whether increased IL-17 production from CD4+ T cells in naïve mice has any impact on IL-17 production by γδ T cells upon immunization remains to be determined.

We have to emphasize that IL-10 signaling in T cells did not affect T-cell-B-cell collaboration or the production of CII-specific antibodies (Figure 6). IL-10 has been defined as a B-cell-stimulating factor and can promote antibody production [42,43]. Because the function of T cells but not B cells was affected in Tg mice [21], it was not surprising that we saw no differences in autoantibody production between Wt and Tg mice. More generally, this Tg mouse may provide a unique model in which to study the dysfunction of T cells without affecting antibody responses.

Our study clearly demonstrates that IL-10 signaling in T cells is critical for the pathogenesis of CIA by
maintaining the level of Foxp3 and consequently contributing to the dysfunctions of Tregs as well as to the increased expression level of the innate source of IL-17A by γδ T cells. Enhancement of IL-10 signaling in T cells may regulate the function of Tregs, which can dampen the harmful autoimmune responses.

Conclusions
Our results demonstrate a critical role of IL-10 signaling in T cells in the pathogenesis of CIA. Without IL-10 signaling, T cells, especially Tregs, lost their suppressive function, which in turn failed to control CD4+ T-cell activation. Enhancement of IL-10 signaling in T cells might serve as a therapeutically sound approach to the treatment of RA as well as other autoimmune inflammatory diseases.

Abbreviations
CFA: Complete Freund’s Adjuvant; CIA: collagen-induced arthritis; ELISA: enzyme-linked immunosorbent assay; GFP: green fluorescent protein; H & E: hematoxylin and eosin; IFIA: Incomplete Freund’s Adjuvant; IFN-γ: interferon-γ; IL: interleukin; PCR: polymerase chain reaction; TCR: T-cell receptor; Tg: transgenic; Wt: wild type.

Acknowledgements
This research was made possible by a grant from the American College of Rheumatology Research and Education Foundation Within Our Reach: Finding a Cure for Rheumatoid Arthritis campaign (to RAF and ZY). Dr Yin is supported by the National Natural Science Foundation of China (30725015 and 30890143), the National Basic Research Program of China (2010CB529104) and the International S&T Cooperation Program of China (2010DFA09000). Dr Yin was also supported by the Program of Introducing Talents of Discipline to Universities (B080111) and 07ZCKFSH0360 (Tianjin Science and Technology Commission). Zhong was supported by 09ZCKFSH0200 (Tianjin Science and Technology Commission).

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Authors’ contributions
JT performed most of the animal experiments as well as the in vitro experiments. MK prepared the IL-10 receptor dominant-negative transgenic mice. JH performed T-cell proliferation assays and intracellular cytokine staining. JC, one of the senior authors, engaged in discussion and made suggestions regarding the experiments. RF, one of the senior authors, was responsible for the design of the experiments and discussed the data. ZW helped to perform the animal experiments. Zhong helped to perform in vitro assays. LZ helped with the data analysis and interpretation of the results. ZY, the corresponding author, designed the experiments and wrote the paper. ZHao performed the histological scoring. XJ assessed the H & E staining and took the histologic pictures. All authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

Received: 21 August 2011 Revised: 11 November 2011 Accepted: 22 December 2011 Published: 22 December 2011

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