IL-10 inhibits a broad spectrum of cellular immune responses. It suppresses the function of APCs and T cells by inhibiting proinflammatory cytokine production, co-stimulation, MHC class II expression, and chemokine secretion (1, 2). IL-10 has been associated with immunopathology in various immune-mediated and inflammatory diseases. For example, treatment with a combination of anti–IL-10R monoclonal antibody and Toll-like receptor 9 (TLR9) ligands had potent therapeutic antitumor effects (3, 4), indicating a role for IL-10 in the pathogenesis of cancer.

Viruses use a variety of strategies to avoid recognition by the host immune system (5–7). The active induction of immune suppression is one mechanism by which viruses escape clearance and thus establish a persistent infection (8). In humans, chronic viral infections afflict millions of people worldwide (9–11). Interestingly, elevated levels of IL-10 production have been associated with persistent infection by hepatitis C virus (HCV) (12, 13), HIV (14–18), and Epstein-Barr virus (1, 19). It was recently reported that upon HCV infection, intrahepatic CD8+ T cells from persistently infected subjects suppressed the in vitro proliferative responses of liver-derived lymphocytes in an HCV-specific and IL-10–dependent manner (20). Moreover, both CD4+ and CD8+ T cells have been shown to express high levels of IL-10 in HIV-infected individuals (14, 15). In addition, a higher frequency of IL-10–producing CD4+ T cells was observed in HIV-infected individuals with progressive disease or active HIV replication compared with individuals in the latent phase of disease (16–18).

To gain further insight into the role of IL-10 in the establishment and maintenance of persistent viral infections, we investigated whether this cytokine is involved in the persistence of lymphocytic choriomeningitis virus (LCMV) infection in its natural host, the mouse. LCMV is an arena virus that can cause either acute or persistent infection in vivo depending on the strain, route of infection, and dose of virus (21). Although adult mice infected with LCMV Armstrong rapidly clear the infection and establish a stable memory T cell pool (22–27), infection with a naturally selected isolate of LCMV Armstrong, the LCMV variant clone 13, results in a prolonged infection that persists (28–30). This chronic infection is associated with the functional impairment, exhaustion,
and deletion of virus-specific CD8+ T cells (31, 32), resulting in viral persistence, which was recently linked to expression of the programmed death 1 receptor (PD-1), an inhibitory receptor of the CD28 family (33–36).

DCs, which are key regulators of immune responses, play an important role in clearing viral infections. Upon engagement of DCs, naïve Th cells polarize into IFN-γ-producing Th1 or IL-4-producing Th2 effector cells (37, 38). It has previously been suggested that different DC subsets, for example CD11c+CD8α− and CD11c−CD8α+ DCs, have the potential to differentially induce Th2 and Th1 cells, respectively (39–41). Others have previously reported that, after LCMV clone 13 infection, LCMV-specific CD4+ T cells from TCR transgenic SMARTA mice produced higher levels of IL-10 than after LCMV Armstrong infection (42). In this study, we demonstrate that production of IL-10 during LCMV clone 13 infection is associated with viral persistence, because blockade of the receptor for IL-10 restored the antiviral immune response and resulted in viral clearance. This rapid resolution of viral infection after anti–IL-10R treatment was associated with diminished levels of endogenous IL-10 production and enhanced antiviral CD8+ T cell responses. Further analysis revealed that viral persistence during LCMV clone 13 infection was linked to a decline in the number of CD11c+CD8α− DCs.

CD11c+CD8α− DCs efficiently induced IL-10 secretion by antiviral CD4+ T cells, preventing viral clearance and therefore enabling viral persistence. Therapeutic IL-10R blockade broke the cycle of IL-10–mediated immune suppression, preventing IL-10 priming by CD8α− DCs and enhancing antiviral responses, thereby resolving infection in persistently infected mice.

These results highlight for the first time the role of IL-10 in the suppression of an antiviral immune response during a persistent viral infection. Upon blockade of signaling through the IL-10R, secretion of IL-10 was almost completely abrogated. We therefore propose that IL-10R blockade may be of therapeutic benefit in the treatment of chronic viral infections.

RESULTS Persistent infection with LCMV clone 13 is associated with increased IL-10 production

Previous studies reporting IL-10 production during chronic viral infections prompted us to examine the secretion of IL-10 during chronic and acute LCMV infections (1, 12–19). Splenocytes from mice infected with LCMV clone 13 or LCMV Armstrong were isolated at various time points after infection and cultured for 48 h in the absence of exogenous antigen (to allow for direct ex vivo presentation of viral antigens by splenic APCs). At the end of this culture period, we measured the concentration of IL-10 in the supernatants by ELISA. IL-10 production by splenocytes from mice infected with LCMV (both clone 13 and Armstrong strains) peaked at day 20 after infection (Fig. 1 A). However, splenocytes from mice infected with LCMV Armstrong secreted significantly less IL-10. This difference was apparent regardless of the initial dose of virus, as increasing the inoculum of LCMV Armstrong from 10^4 PFU to 2 × 10^6 PFU resulted in comparable kinetics of IL-10 secretion (unpublished data). By day 40, although IL-10 production was undetectable in LCMV Armstrong–infected mice, mice infected with LCMV clone 13 still produced significant amounts of this cytokine (Fig. 1 A). After in vitro LCMV clone 13 infection, we observed that IL-10 was produced by different immune cell types, including CD4+ and CD8+ T cells and CD11c+ DCs, whereas LCMV Armstrong infection induced less IL-10 production by CD4+ T cells and CD11c+ DCs and none by CD8+ T cells (Fig. 1 B). Interestingly, when splenic cell cultures were treated with a neutralizing anti–IL-10R monoclonal antibody (which specifically targets the IL-10R α chain) at the time of LCMV clone 13 infection in vitro, the levels of IL-10 produced by CD4+ T cells were reduced to those found in LCMV Armstrong–infected mice (Fig. 1 B).

We then analyzed in vivo IFN-γ secretion by CD8+ T cells at different time points after LCMV Armstrong and LCMV clone 13 infection (Fig. 1 C). Interestingly, CD8+ T cells from LCMV clone 13–infected mice secreted IFN-γ only during the early phase of infection (day 5), at a time when no IL-10 was yet detected (Fig. 1 A). However, IFN-γ secretion was almost completely lost at later time points, and high levels of IL-10 were produced (Fig. 1, A and C). In contrast, IFN-γ production by CD8+ T cells from LCMV Armstrong–infected mice peaked later (day 7) and was sustained in the form of a memory response (Fig. 1 C) (27). Secretion of TNF-α followed a similar profile (unpublished data). Increasing the dose of LCMV Armstrong did not induce earlier cytokine responses (unpublished data). These results indicate that LCMV clone 13 induces a faster antiviral response, possibly linked to a higher binding affinity of the virus to its α-dystroglycan receptor (43).

It has recently been shown that high expression of PD-1 is a characteristic of exhausted CD8+ T cells in LCMV clone 13–infected mice and that treatment with anti–PD-1 antibodies leads to the proliferation of antiviral T cells and the enhancement of viral clearance (33–36). We therefore analyzed PD-1 expression on T cells. We found that a small percentage of CD4+ and CD8+ T cells expressed PD-1 in naive mice (Fig. 1 D). Upon LCMV infection, significantly higher levels of PD-1 were expressed on CD8+ T cells isolated from LCMV clone 13–infected mice compared with LCMV Armstrong–infected mice (Fig. 1 D). In addition, the mean fluorescence intensity of PD-1 was increased in T cells from LCMV clone 13– versus LCMV Armstrong–infected mice (unpublished data).

Based on these findings, we hypothesized that the immune suppression mediated by LCMV clone 13 infection was caused by a shift from antiviral IFN-γ production to the secretion of the immunosuppressive cytokine IL-10. To investigate whether IL-10 was directly involved in the suppression of the antiviral immune response, we infected IL-10–deficient (IL-10−/−) mice with LCMV clone 13 and monitored viral clearance. 3 wk after LCMV clone 13 infection, we found...
plotted as IL-10 (pg/ml) produced per 10^6 cells and are means of three
present in the culture supernatants was measured by ELISA. Data are
PFU LCMV clone 13 and cultured for 48 h. The concentration of IL-10
were infected with 2\times10^6 PFU LCMV clone 13. Spleens were harvested 6 h after
and PD-1–expressing T cells are shown. Data show one representative mouse
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20, 40, and 94 after infection with 10^5 PFU LCMV Armstrong or 2 \times 10^6
BALB/c splenocytes were isolated and infected in vitro for 48 h with LCMV
experiments. (B) To reveal the origin of IL-10–producing cell subsets,
mice per time point. The experiment is representative of three similar
expression was detected on CD4^-/^- and CD8^-/^- cells. Data are representative of one experiment with five mice per group. (C) Sensitive direct in vivo intracellular cytokine assay was performed to detect in vivo cytokine expression in LCMV-specific T cells, as described by Liu et al. (reference 61). In brief, mice were injected i.v. with 0.5 mg BFA on days 5 and 7 after infection with 10^5 PFU LCMV Armstrong or 2 \times 10^6 PFU LCMV clone 13. Spleens were harvested 6 h after BFA injection, and direct in vivo intracellular cytokine detection was performed after staining with fluorescent antibodies to CD4, CD8, IFN-\gamma, and TNF-\alpha. Mean values per spleen of three to five individual mice are shown. The experiment is representative of two similar experiments. (D) PD-1 expression was detected on CD4^+ and CD8^+ T cells from naive or day 7 LCMV Armstrong– or LCMV clone 13–infected mice by labeling splenocytes with fluorescent antibodies to CD4, CD8, and PD-1. Percentages of PD-1–expressing T cells are shown. Data show one representative mouse per group (three mice total per group). (E) IL-10^-/- and wild-type mice were infected with 2 \times 10^6 PFU LCMV clone 13, and viral titers were detected in kidney and liver from three mice per group by RT-PCR 3 wk after infection. Data are shown as mean LCMV genome copies per mg organ ± SEM. (F) Cytokine expression was monitored in wild-type and IL-10^-/- mice 7 d after LCMV clone 13 infection by intracellular cytokine staining. Splenocytes were stimulated with BFA and GP33-41 or GP61-80 peptide for 6 h. Intracellular cytokine detection was performed by staining with fluorescent antibodies to CD4, CD8, IFN-\gamma, and TNF-\alpha. The IL-10/IFN-\gamma ratio of CD4^+ and CD8^+ T cells is shown as mean values of three individual mice per group. Statistical analysis was performed using the Student's t test. *, P < 0.01; **, P < 0.001; and ***, P < 0.0001.

Figure 1. Time course of T cell cytokine production after infection with LCMV clone 13: the role of IL-10 in promoting viral persistence. (A) Spleen cells were isolated from BALB/c mice on days 0, 5, 7, 14, 20, 40, and 94 after infection with 10^5 PFU LCMV Armstrong or 2 \times 10^6 PFU LCMV clone 13 and cultured for 48 h. The concentration of IL-10 present in the culture supernatants was measured by ELISA. Data are plotted as IL-10 (pg/ml) produced per 10^6 cells and are means of three mice per time point. The experiment is representative of three similar experiments. (B) To reveal the origin of IL-10–producing cell subsets, BALB/c splenocytes were isolated and infected in vitro for 48 h with LCMV Armstrong or LCMV clone 13 without or with anti–IL-10R antibody treatment (multiplicity of infection = 3). Next, CD4^+ , CD8^+ , and CD11c^-cells were enriched using MACS beads, and the supernatant of the purified populations was analyzed for IL-10 production by ELISA. The graph indicates the levels of IL-10 (pg/ml) in the supernatant of CD4^-/^- , CD8^-/^- , and CD11c^- cells. Data are representative of one experiment with five mice per group. (C) Sensitive direct in vivo intracellular cytokine assay was performed to detect in vivo cytokine expression in LCMV-specific T cells, as described by Liu et al. (reference 61). In brief, mice were injected i.v. with 0.5 mg BFA on days 5 and 7 after infection with 10^5 PFU LCMV Armstrong or 2 \times 10^6 PFU LCMV clone 13. Spleens were harvested 6 h after BFA injection, and direct in vivo intracellular cytokine detection was performed after staining with fluorescent antibodies to CD4, CD8, IFN-\gamma, and TNF-\alpha. Mean values per spleen of three to five individual mice are shown. The experiment is representative of two similar experiments. (D) PD-1 expression was detected on CD4^+ and CD8^+ T cells from naive or day 7 LCMV Armstrong– or LCMV clone 13–infected mice by labeling splenocytes with fluorescent antibodies to CD4, CD8, and PD-1. Percentages of PD-1–expressing T cells are shown. Data show one representative mouse per group (three mice total per group). (E) IL-10^-/- and wild-type mice were infected with 2 \times 10^6 PFU LCMV clone 13, and viral titers were detected in kidney and liver from three mice per group by RT-PCR 3 wk after infection. Data are shown as mean LCMV genome copies per mg organ ± SEM. (F) Cytokine expression was monitored in wild-type and IL-10^-/- mice 7 d after LCMV clone 13 infection by intracellular cytokine staining. Splenocytes were stimulated with BFA and GP33-41 or GP61-80 peptide for 6 h. Intracellular cytokine detection was performed by staining with fluorescent antibodies to CD4, CD8, IFN-\gamma, and TNF-\alpha. The IL-10/IFN-\gamma ratio of CD4^+ and CD8^+ T cells is shown as mean values of three individual mice per group. Statistical analysis was performed using the Student's t test. *, P < 0.01; **, P < 0.001; and ***, P < 0.0001.

that viral titers were significantly lower in liver and kidney from IL-10^-/- mice compared with wild-type control mice when measured by a highly sensitive RT-PCR method (Fig. 1 E). Resolution of LCMV clone 13 infection in IL-10^-/- mice was associated with a lower systemic IL-10 to IFN-\gamma ratio, as one would expect (Fig. 1 F). Sustenance of antiviral immune responses to clone 13 through genetic elimination of IL-10 has been described in detail in another study by Brooks et al. (44).

Anti–IL-10R antibody therapy decreases viral titers and leads to disease amelioration

IL-10 can inhibit immune responses by skewing the development of helper T cells and suppressing their function (45, 46). Based on our initial findings that cells from LCMV clone 13–infected mice secreted IL-10 and that mice deficient in IL-10 cleared this strain of LCMV more efficiently, we investigated whether blocking the IL-10–IL-10R signaling pathway could resolve LCMV clone 13–induced immune suppression and reestablish an antiviral Tc1/Th1 response. We therefore infected BALB/c mice with LCMV clone 13 and injected age-matched groups with either a neutralizing anti–IL-10R monoclonal antibody or an IgG1, isotype control antibody on days 0, 7, and 14, or with a therapeutic regimen on days 7 and 14 after infection. Disease severity was monitored by assessing bodyweight, spleen cell numbers, and viral titer over time. LCMV clone 13–infected mice treated with isotype control antibody lost body mass and weighed 30–40% less than age-matched naive mice (Fig. 2, A and C). These mice also exhibited a nonshiny, scruffy coat (unpublished data). In comparison, LCMV clone 13–infected mice treated with anti–IL-10R antibody lost weight less rapidly (Fig. 2 A) and exhibited a healthy shiny coat (not depicted). In addition, we monitored viral loads in lymphoid and nonlymphoid organs isolated from persistently infected mice treated with anti–IL-10R or IgG1. Viral titers were measured by conventional plaque assay (not depicted) and by a more recently developed, highly sensitive RT-PCR method (Fig. 2 B). At 6 and 26 wk after infection, numbers of LCMV genomic copies were low or nondetectable when measured by the sensitive RT-PCR method in organs from anti–IL-10R–treated mice; 36 wk after infection, numbers of LCMV genomic copies were low or nondetectable when measured by the sensitive RT-PCR method in organs from anti–IL-10R–treated mice; 36 wk after infection, viral titers remained undetectable in liver and kidney (Fig. 2 B). In contrast, viral genome copies were still detectable in these organs when mice were treated with IgG1 isotype antibody, indicating that resolution of chronic infection resulted from anti–IL-10R treatment rather than

infection. Data are shown as mean LCMV genome copies per mg organ ± SEM. (F) Cytokine expression was monitored in wild-type and IL-10^-/- mice 7 d after LCMV clone 13 infection by intracellular cytokine staining. Splenocytes were stimulated with BFA and GP33-41 or GP61-80 peptide for 6 h. Intracellular cytokine detection was performed by staining with fluorescent antibodies to CD4, CD8, IFN-\gamma, and TNF-\alpha. The IL-10/IFN-\gamma ratio of CD4^+ and CD8^+ T cells is shown as mean values of three individual mice per group. Statistical analysis was performed using the Student's t test. *, P < 0.01; **, P < 0.001; and ***, P < 0.0001.
progressive viral elimination over time. Viral titers in organs from control IgG1-treated mice remained high, and virus predominated in the kidney, characterizing the chronic infection (Fig. 2B). Furthermore, extremely low viral titers were detected by LCMV plaque assay in kidney, liver, and lung 6 wk after LCMV clone 13 infection in anti–IL-10R–treated mice.
mice, and no virus was detected 26 wk after infection (unpublished data).

Importantly, we also administered anti–IL-10R antibody in a therapeutic setting at days 7 and 14 after infection during the onset of lymphopenia and weight loss (by day 7 after infection, the total number of splenocytes was reduced by 41% in LCMV clone 13–infected mice compared with naive mice; unpublished data). Anti–IL-10R treatment was therapeutically effective during established infection as treated mice lost weight less rapidly (Fig. 2 C), did not exhibit any overt signs of disease, and showed a substantial reduction in viral titers in the liver, lung, and kidney 11 wk after treatment (not depicted). Additionally, a two to three log reduction in viral titer was observed when anti–IL-10R was given on days 5 and 12 after LCMV clone 13 infection, as well as on days 10 and 17 after infection (unpublished data).

Interestingly, there was no difference in anti-LCMV–specific antibody production after anti–IL-10R treatment (unpublished data), indicating that viral clearance was the result of an enhancement of the antiviral T cell responses. These results demonstrate that blocking IL-10 signaling results in clearance of an otherwise protracted LCMV clone 13 infection. Notably, anti–IL-10R treatment was effective even when given therapeutically during established infection.

**Anti–IL-10R treatment of persistently infected mice restores antiviral CD8+ T cell responses**

As reported previously (31, 47), we found that chronic infection was associated with deletion of virus-specific CD8+ T cells, leading to a drastic decrease in the number of spleen cells (unpublished data). We discovered that this state of lymphopenia was reversed after anti–IL-10R therapy both when treatment was initiated at day 0 or therapeutically at day 7 after infection (Fig. 3 A). In addition, endogenous IL-10 production by splenocytes isolated 3 wk after LCMV clone 13 infection was significantly reduced in anti–IL-10R–treated mice, indicating that early blockade of this cytokine prevented the long-term immunosuppressive effects mediated by LCMV clone 13 infection (Fig. 3 B).

We therefore assessed the effect of anti–IL-10R treatment on the LCMV-specific T cell response in detail. Splenocytes from anti–IL-10R– and control IgG1–treated BALB/c mice were isolated at various time points after LCMV clone 13 infection and stimulated with the MHC class I–restricted immunodominant LCMV peptide NP118-126. The number of IFN-γ–secreting CD8+ cells was measured by intracellular cytokine staining. Anti–IL-10R–treated mice had significantly higher numbers of IFN-γ–secreting CD8+ T cells compared with IgG1–treated mice, particularly at later time points (Fig. 3 C). Importantly, therapeutically treatment with anti–IL-10R on days 7 and 14 after LCMV clone 13 infection caused a similar enhancement of the LCMV-specific memory T cell response (Fig. 3 C). Thus, anti–IL-10R treatment resulted in reemergence of a potent antiviral IFN-γ–secreting CD8+ T cell response, indicating that events occurring early during T cell responses can profoundly affect the quality of T cell memory. It will be difficult to determine whether the enhancement of viral-specific CD8+ T cell responses was the cause or consequence of reduced viral loads; however, we believe that both alternatives are possible.

Finally, we investigated the effect of anti–IL-10R treatment on the PD-1 inhibitory pathway (33–36). To determine whether suppression of the PD-1 pathway could be involved in the enhancement of viral clearance after therapeutic anti–IL-10R treatment, we monitored expression of PD-1 by T cells in anti–IL-10R– and IgG1–treated mice 90 d after LCMV clone 13 infection. Low levels of PD-1 expression were detected on T cells from naive mice (Fig. 3 D, top). In comparison, T cells from LCMV clone 13–infected mice treated with IgG1 isotype antibody exhibited as much as a 25-fold induction of PD-1 expression compared with naive mice (Fig. 3 D, bottom). Importantly, a threefold reduction of PD-1 expression was observed on T cells from anti–IL-10R–treated mice compared with IgG1–treated mice (Fig. 3 D, middle). These results indicate that anti–IL-10R treatment decreased PD-1 expression on T cells, which could potentially contribute to abrogation of T cell exhaustion because treatment with anti–PD-1 antibodies in chronically infected mice has been shown to down-regulate PD-1 and circumvent virally induced CD8+ T cell exhaustion (33–36).

**Anti–IL-10R treatment reverses lymphopenia in LCMV clone 13–infected mice**

To further investigate the effect of anti–IL-10R treatment on the LCMV-specific immune response, we examined absolute numbers of different splenic cell subsets over time in mice infected with LCMV clone 13 and treated with anti–IL-10R or control IgG1 antibody. We discovered that even though LCMV clone 13 infection caused a reduction in the numbers of cells present in the spleen at day 21 after infection regardless of treatment, by day 150 after infection the numbers of CD4+, CD8+, B220+, and CD11c+ cells in anti–IL-10R–treated mice were similar to those in naive mice, whereas in IgG1–treated mice these cells remained scarce (Fig. 4 A). Notably, the numbers of CD11b+ cells in the spleen at day 150 after infection were comparable between groups, suggesting that the decrease in viral titers after anti–IL-10R treatment was not solely caused by enhanced clearance of virally infected macrophages.

**CD8α+ DCs are eliminated after LCMV clone 13 but not LCMV Armstrong infection**

We hypothesized that the loss of antiviral T cells after LCMV clone 13 infection would be mediated by APCs. DCs are the most potent APCs and have been shown to play a crucial role in the priming of LCMV-specific cytotoxic T cells (48). We therefore examined absolute numbers of different DC subsets (CD11c+CD8α− and CD11c+CD8α+) and their ratio after infection with LCMV Armstrong or LCMV clone 13. Over the course of infection, the number of CD8α− and CD8α+ DCs increased in LCMV Armstrong–infected mice but returned to that found in naive mice within 2 wk after infection.
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(Fig. 4 B). In contrast, although the number of CD8α− DCs from LCMV clone 13–infected mice 6 wk after infection was also comparable to that of naive mice, CD8α+ DCs gradually disappeared over time (Fig. 4 B). It has been previously shown that >50% of CD11c+ DCs carry viral particles at this time point in clone 13–infected mice (49), making these cells an excellent target for destruction and elimination by the immune system. In contrast, very few CD11c+ cells are infected at the same time point after LCMV Armstrong infection (49). Importantly, the numbers of CD8α+ DCs were significantly lower in LCMV clone 13–infected mice 6 wk after infection compared with mice infected with LCMV Armstrong (Fig. 4 B). Because the CD8α+ DC subset declined in LCMV clone 13–infected mice and the CD8α− DC subset remained at a stable level, the CD8α− to CD8α+ DC ratio was skewed toward CD8α− DCs 6 wk after infection (Fig. 4 C). To investigate whether this phenomenon was affected by therapeutic anti–IL-10R treatment, absolute numbers of CD8α− and CD8α+ DCs were monitored 6 wk after LCMV clone 13 infection in mice treated with anti–IL-10R on days 7 and 14 after infection. Anti–IL-10R treatment decreased CD8α− DC numbers more than twofold, as well as CD8α+ DC numbers to a lesser extent (Fig. 4 B). Hence, the resulting CD8α− to CD8α+ DC ratio after anti–IL-10R treatment resembled the ratio in LCMV Armstrong–infected mice 6 wk after infection (Fig. 4 C). In summary, these findings show that anti–IL-10R treatment in LCMV clone 13–infected mice resulted in a skewing of the CD8α− to CD8α+ DC ratio to levels seen in mice infected with the nonpersistent LCMV Armstrong strain.

Figure 3. Anti–IL-10R antibody treatment in LCMV clone 13–infected mice increases total cell numbers, decreases IL-10 production, induces a functional antiviral memory T cell pool, and reduces the percentage of PD-1–expressing T cells. (A) 6-wk-old BALB/c mice were infected with 2 × 10^6 PFU LCMV clone 13 and treated with an IgG1, isotype control antibody or i.p. with 250 μg of a neutralizing anti–IL-10R antibody on days 0, 7, and 14 (left) or with a therapeutic regimen on days 7 and 14 after LCMV clone 13 infection (right). On days 20 and 150 after LCMV infection, spleen cells were isolated and counted. Histogram bars represent mean values ± SD for three mice per group. The experiment is representative of three similar experiments. (B) Splenocytes from naive mice, LCMV clone 13–infected mice treated with IgG1, or anti–IL-10R antibody on days 0, 7, and 14 were isolated 3 wk after infection and cultured for 48 h. The amount of IL-10 present in the culture supernatant was measured by ELISA. Data are plotted as IL-10 (pg/ml) produced per 10^6 cells and are means of three mice per plate. Results are representative of three identical experiments. (C) Splenocytes from LCMV clone 13–infected BALB/c mice treated with IgG1 or anti–IL-10R antibody on days 0, 7, and 14 (left) or day 7 and 14 (right) were isolated at day 20 (early memory) and 150 (late memory) after infection. Cells were stimulated in the presence of NP118-126 peptide and BFA for 5 h. Intracellular cytokine staining was performed using fluorescent antibodies to CD8 and IFN-γ. The numbers of IFN-γ+CD8+ T cells per spleen are shown. Histogram bars represent mean values ± SD for three mice per group. The experiment is representative of three similar experiments. (D) BALB/c mice were treated with anti–IL-10R antibody or IgG1, isotype control antibody on days 7 and 14 after LCMV clone 13 infection. 90 d after infection, percentages of PD-1–expressing splenic CD4+ and CD8+ T cells were analyzed by gating on CD4+ (left) and CD8+ T cells (right) and staining for PD-1. Numbers in dot plots represent percentages of CD4+ and CD8+ T cells expressing PD-1. Data show one representative mouse per group (four mice total per group). Statistical analysis was performed using the Student’s t test. *, P < 0.01; **, P < 0.001; and ***, P < 0.0001.
Previous reports suggest that different DC subsets vary in their ability to prime effector T cells (39, 50, 51). In particular, evidence suggests that different DC subsets can induce T cells to produce different cytokines depending on the cytokine milieu in which they encounter antigen (40, 41, 46). We therefore investigated whether T cell priming by different DC subsets modulated the nature of the antiviral T cell response. CD11c⁺CD3⁻ splenic DCs were isolated from mice infected 7 d earlier with LCMV Armstrong or LCMV clone 13. In addition, one group of mice was treated with anti-IL-10R antibody at the time of infection. CD11c⁺CD3⁻ DCs from all groups were sorted into CD8α⁻ and CD8α⁺ DC subsets on day 7 after infection (Fig. 5). This early time point was chosen to allow for capture and processing of viral antigens by DCs directly in vivo during the early phase of LCMV Armstrong and LCMV clone 13 infection, which is
associated with viral dissemination and replication. At this time, viral antigen could be detected by RT-PCR (unpublished data). Because we had previously observed that CD4+ T cells from LCMV clone 13–infected mice produced elevated levels of IL-10, the isolated DCs were cultured for 5 d with naive GP61-80–specific CD4+ responder T cells isolated from TCR transgenic SMARTA mice (52). No exogenous antigen was added to the cultures to ensure that only viral antigen processed in vivo was presented by the DCs. The ability of the different DC subsets to stimulate LCMV-specific CD4+ T cells was determined by measuring the concentration of cytokines in the supernatants by ELISA at the end of the culture period. To rule out any contamination caused by cytokine release by APCs, DCs were irradiated, and levels of both IL-10 and IFN-γ were measured in the supernatants of DC cultures devoid of T cells. These background levels were <1 pg/ml (unpublished data).

IL-10 production by antiviral CD4+ T cells was induced exclusively by DCs isolated from LCMV clone 13–infected mice and was preferentially mediated by the CD8α− DC subset (Fig. 5 A), an observation that was confirmed when the different DC subsets were loaded with LCMV GP61-80 peptide (not depicted). Importantly, this corresponded to the time point at which CD8α+ DC numbers in LCMV clone 13–infected mice started to decline (Fig. 4 B). These results suggest that when CD8α+ DCs are infected and killed, the remaining CD8α− DCs that are left to prime T cells induce IL-10 production instead of a potent antiviral response. In contrast, CD8α− or CD8α+ DCs isolated from mice infected with LCMV Armstrong induced little to no IL-10 production by LCMV-specific CD4+ T cells (Fig. 5 A), indicating that differences between the two LCMV infections could merely reside in the type of cytokine responses induced. In vivo anti–IL-10R treatment completely abrogated the capacity of CD8α+ DCs to induce IL-10 secretion in CD4+ T cells (Fig. 5 A), indicating that signaling through the IL-10R was crucial for CD8α+ DCs to induce IL-10.

Additionally, both CD8α− and CD8α+ DCs from LCMV clone 13– and LCMV Armstrong–infected mice were able to stimulate IFN-γ production by CD4+ T cells, whereas CD8α+ DCs induced higher amounts of IFN-γ than CD8α− DCs. Anti–IL-10R treatment did not have an effect on IFN-γ production induced by the different DC subsets (Fig. 5 B).

Importantly, although the amount of IL-10 induced by CD8α− and CD8α+ DC subsets was dependent on the LCMV strain used (Fig. 5 A), the amount of IFN-γ was similar for both LCMV strains (Fig. 5 B). Therefore, the IL-10 to IFN-γ ratio was always significantly higher when DCs from LCMV clone 13–infected mice were used as APCs (Fig. 5 C).

Collectively, these data show that anti–IL-10R treatment abolished CD8α− DC-mediated IL-10 production in responder T cells early after infection and decreased absolutely CD8α− DC numbers over time, thereby inducing a shift toward IFN-γ+ and Tc1/Th1 responses, enabling viral clearance. Our observations that anti–IL-10R antibody led to resolution of protracted infection, enhanced antiviral T cell responses, and abrogated the ability of CD8α− DCs to induce IL-10 in CD4+ T cells, in conjunction with observed lower viral titers in mice deficient in IL-10, underscores the important role IL-10 plays in maintaining a chronic viral infection.
DISCUSSION

This study is the first to report that inhibiting IL-10 signaling in vivo leads to enhancement of viral clearance in a mouse model of persistent viral infection. Our data demonstrate that upon challenge with LCMV clone 13, the strong antiviral response elicited early after infection rapidly subsides and is followed by a state of chronic infection associated with generalized lymphopenia. In this paper, we show that chronic infection and deterioration in health is associated with systemic increase in the production of IL-10. Although the implication for the role of IL-10 in the maintenance of chronic viral infections has hitherto been unclear, our results show that lymphopenia is reversed, bodyweight recovers, and viral load is significantly reduced upon blockade of IL-10–IL-10R signaling ($P < 0.0001$). Furthermore, we show that anti–IL-10R treatment results in a reduction of CD8α− DC numbers and a subsequent abolishment of IL-10 production, thereby favoring a Tc1/Th1 cell–like environment inducing LCMV–specific IFN-γ+ T cell responses. Indeed, after LCMV clone 13 infection, CD8α+ DC numbers substantially were found to decrease because of viral infection and clearance, and we now show that CD8α− DCs will preferentially induce IL-10 production by T cells. Thus, it is possible that the generalized state of lymphopenia of LCMV clone 13–infected mice is mediated by APCs inducing IL-10–producing T cells, which results in a negative feedback loop. Our results suggest that distinct DC subsets may differentially regulate the cytokine balance of the immune response in vivo. The precise mechanism by which these DC subsets mediate their effects might be related to levels of IL-10 present in the microenvironment. We show that CD8α− DCs induce IL-10 production more efficiently than their CD8α+ counterparts. The ability to potentiate IL-10 secretion is drastically reduced upon blockade of signaling through the IL-10R. We believe that anti–IL-10R treatment modulates CD8α− T cell responses. Indeed, after LCMV clone 13 infection, CD8α+ DC numbers substantially were found to decrease because of viral infection and clearance, and we now show that CD8α− DCs will preferentially induce IL-10 production by T cells. Thus, it is possible that the generalized state of lymphopenia of LCMV clone 13–infected mice is mediated by APCs inducing IL-10–producing T cells, which results in a negative feedback loop. Our results suggest that distinct DC subsets may differentially regulate the cytokine balance of the immune response in vivo. The precise mechanism by which these DC subsets mediate their effects might be related to levels of IL-10 present in the microenvironment. We show that CD8α− DCs induce IL-10 production more efficiently than their CD8α+ counterparts. The ability to potentiate IL-10 secretion is drastically reduced upon blockade of signaling through the IL-10R. We believe that anti–IL-10R treatment modulates CD8α− DCs early upon infection and, as a result, dictates the nature of the cytokine response induced early during antiviral immunity.

The mechanisms by which IL-10 enables LCMV clone 13 to persist are unknown. Several variables associated with T cell priming have been shown to be important and may program qualitative differences into the effector and memory populations (53). IL-10 could either down-regulate proinflammatory responses in a general manner or, more specifically, inhibit the induction or expansion of antiviral CD8+ effector T cells. Importantly, we report for the first time that anti–IL-10R antibody treatment reduces PD-1 expression on CD8+ T cells, which has previously been shown to contribute to virally induced CD8+ T cell exhaustion (33–36). Interestingly, stimulation through TLR2, which has been shown to bind to LCMV, leads to the production of IL-10 and inhibits IFN-inducible protein 10 and IL-12 secretion by APCs. Thus, stronger binding of LCMV clone 13 to TLR2, as opposed to weaker TLR2 binding by LCMV Armstrong (54), could not only induce preferential production of IL-10 by DCs but also inhibit the secretion of cytokines that promote DC differentiation, maturation, and proliferation rather than survival (49). In addition, it is possible that the DCs that remain after LCMV clone 13 infection express higher levels of TLR2, thus perpetuating the cycle.

Subclasses of DCs have been shown to have the potential to differentially skew T cell cytokine production toward Th1 or Th2 cell profiles (40, 41, 55). Notably, it has been suggested that CD8α− DCs induce Th2 cell profiles, whereas CD8α+ DCs preferentially stimulate IFN-γ production and, therefore, induce Th1 cell profiles (39, 51). We now show that this may be true in the LCMV system, as we have analyzed the ability of CD8α− and CD8α+ DCs to polarize LCMV-reactive CD4+ T cells ex vivo. It appears that priming of cytotoxic T lymphocytes by DCs is achieved in a similar fashion in LCMV clone 13– and LCMV Armstrong–infected mice, but that stronger binding of LCMV clone 13 to its receptor on APCs, particularly CD8α+ DCs, leads to more efficient processing and presentation of viral antigen. Our finding that LCMV clone 13 infection leads to early induction of IFN-γ–secreting LCMV–specific CD8+ T cells strengthens this hypothesis. The strong cytotoxic function of such activated Tc1 or Th1 cells upon infection with LCMV clone 13 may result in killing of the APC subset responsible for their induction; i.e., DCs belonging to the CD8α+ subset. This hypothesis correlates with our finding that the number of CD8α+ DCs is reduced as early as day 7 after infection with LCMV clone 13 but not LCMV Armstrong.

Our results suggest that as the number of CD8α+ DCs decreases after chronic LCMV clone 13 infection, the CD8α− DC subset—which is more likely to prime T cells to produce IL-10 than Tc1/Th1 cytokines—will by default become the modulator of the T cell response. LCMV-specific CD4+ T cells activated in this context may provide inappropriate or insufficient antiviral help to other cell types, particularly CD8+ T cells, thus leading to persistent infection. Additionally, it is possible that the CD8α− DCs, which appear ill-equipped to propagate antiviral effector cells, will continue to support IL-10 production. The resulting high concentration of IL-10 in the milieu may thus lead to further modulation of DC function. In fact, IL-10 may directly decrease the viability of CD8α+ DCs, as has been previously suggested (40, 56). As a result, only disruption of IL-10 signaling will have the ability to break the vicious circle and enable the recovery of appropriate antiviral immunity by the infected host.

Mechanistically, our findings show that anti–IL-10R antibody treatment lowers the endogenous levels of IL-10, restores the ability to mount an antiviral CD8+ T cell response, and results in enhanced viral clearance, which all highlight the important role IL-10 plays in the maintenance of chronic infection. Upon treatment with anti–IL-10R, we observed a profound decrease in CD8α− and, to a lesser degree, in CD8α+ DC numbers, as well as a complete inability of CD8α− DCs to induce IL-10 production by CD4+ responder T cells, thereby favoring a systemic cytokine milieu that would better support IFN-γ+ and Tc1/Th1 cell responses.

Importantly, the inability of CD8α− DCs to induce IL-10 production by CD4+ T cells after anti–IL-10R treatment
underscores that IL-10 and signaling through the IL-10R is indeed involved in shaping the cytokine milieu with a potential impact on immune responses. Our results imply that IL-10R blockade is of great interest for the therapeutic treatment of chronic viral infections in humans.

MATERIALS AND METHODS

Mice, virus, and antibody treatment. 6-wk-old BALB/c, C57BL/6 (B6), and IL-10−/− (B6 background) mice were purchased from The Jackson Laboratory. LCMV-GP33-41-specific CD4+ TCR transgenic SMARTA mice (52) were obtained from S. Croo (La Jolla Institute for Allergy and Immunology, La Jolla, CA) and housed under specific pathogen-free conditions at the La Jolla Institute for Allergy and Immunology. LCMV plaques were purified three times on Vero cells, and viral stocks were prepared by a single passage on BHK-1 cells. Age-matched mice were infected i.v. with a single dose of either 2 × 106 PFU LCMV clone 13, 2 × 106 LCMV Armstrong, or an i.p. dose of 105 PFU LCMV Armstrong. For IL-10-blocking experiments, mice were infected i.p. with 250 μg anti-mouse IL-10R (IB1.3a; Becton Dickinson) monoclonal antibody or isotype control antibody (rat IgG1; BD Biosciences). All animal experiments were approved by the La Jolla Institute for Allergy and Immunology Animal Care Committee.

LCMV plaque assay. Organs (kidney, liver, spleen, and lung) were snap-frozen, weighed, and homogenized. In brief, three different dilutions of homogenized organ were prepared, and triplicates were used. Dilutions were incubated at 37°C, 5% CO2 for 1 h with Vero cell monolayers grown in 6-well plates (Costar). The plates were then overlaid with 1% agarose in minimal essential medium 199 (Invitrogen) containing 10% FCS (HyClone) and incubated at 37°C, 5% CO2 for 5 d. The wells were treated with 25% formaldehyde and stained with 0.1% crystal violet for 2 min. The agarose overlay was removed, infectious centers were counted, and the counts were averaged. Additionally, viral LCMV stock was used as a positive control.

RT-PCR. A detailed version of this assay will be published elsewhere (unpublished data). In brief, RNA was isolated from 50 μl of serum or 10 mg of tissue samples using RNAqueous (Ambion). All samples were frozen at −80°C until RNA extraction. RNA was eluted in a volume of 20 μl cDNA reaction with SuperScript III reverse transcriptase (Invitrogen), using primers GP-R and GP-F (CATTCCACCTTGACATTTGTCAGACTCT). A standard curve was generated using pSG5-GP plasmid (57), a gift from J.C. de la Torre (Scripps Research Institute, San Diego, CA). Data were analyzed using linear regression analysis software (Prism; GraphPad).

Quantification of cytokines. Cytokine quantification was performed by sandwich ELISA of supernatants. 2.5 × 105 splenocytes were incubated for 48 h in complete RPMI 1640 (Invitrogen) supplemented with 10% FCS (Sigma-Aldrich), 2 mM L-glutamine (Sigma-Aldrich), 50 μM 2-β-mercaptoethanol (Sigma-Aldrich), and 5 mM Hepes (Sigma-Aldrich) in 24-well plates (Invitrogen). For the detection of IL-10-secreting cell subsets from in vitro infected cultures, bulk splenocytes were infected in vitro with LCMV Armstrong or LCMV clone 13 (multiplicity of infection = 3) without or with 10 μg/mL anti-IL-10R antibody and incubated at 37°C in complete RPMI 1640. 48 h after infection, splenocytes were sorted into CD4+, CD8+, and CD11c+ populations by MACS, and equal numbers of each cell subset were incubated for 5 d at 37°C. Supernatant was removed at the time points indicated in the figures, and ELISA was performed in accordance with the manufacturer’s recommendations and standardized with mouse recombinant cytokine (BD Biosciences). Plates were read at 405 nm (Spectra Max 250; Molecular Devices). The sensitivity of the IL-10 ELISA was 10 pg/mL. Data shown correspond to the concentration of IL-10 (pg/ml) per 106 cells.

Intracellular cytokine staining, cell surface staining, and flow cytometry. To detect cytokine-producing cell subsets, splenocytes from LCMV-infected BALB/c or B6 mice were stimulated with 1 μg/ml NP32-39 or GP33-41 or GP53-60 peptide, respectively, in complete RPMI 1640 containing Brefeldin A (BFA; Sigma-Aldrich) at 37°C. 6 h later, cells were resuspended in staining buffer containing 1% FCS and 0.2% NaN3, labeled with anti-CD4 and anti-CD8, fixed in 1% paraformaldehyde, and permeabilized with 0.1% saponin buffer. Intracellular staining was performed with fluorescent antibodies to IFN-γ, TNF-α, or isotype controls. All antibodies were obtained from BD Biosciences. Additionally, PD-1 expression was detected on splenocytes by labeling with an anti-PE-conjugated PD-1 antibody (eBioscience). Events were acquired using a flow cytometer (FACSCalibur; Becton Dickinson) and analyzed using software (CellQuest; Becton Dickinson). 1–2 × 106 events were acquired, and live cells were gated based on forward/side scatter properties. When analyzing CD11c+ subsets, CD3-expressing cells were gated out to avoid contamination by T cells expressing myeloid markers (58, 59), and expression of the indicated cell surface molecules was detected on CD11c+ and CD8α+ cells. The number of cells was calculated by relating the frequency of cells from each subset to the overall number of cells per spleen.

Sorting of CD11c+CD8α− and CD11c+CD8α+ cells. Splenocytes from mice infected 7 d earlier with LCMV Armstrong or LCMV clone 13 with or without anti–IL-10R antibody treatment were incubated with HBSS medium containing 0.5 mg/ml collagenase D (Sigma-Aldrich) at 37°C, 5% CO2 for 30 min. 0.01 M EDTA was added to disrupt T cell–DC complexes. Next, cells were depleted of CD3-expressing cells (Dynal CD3-bend; Dynal), incubated with CD11c microbeads (Miltenyi Biotec), and positively selected using MACS columns. The enriched CD11c+ cells were labeled with APC-conjugated CD11c and PE-conjugated CD8a antibody (BD Biosciences) and sorted using a cell sorter (FACS Aria; Becton Dickinson) into CD8α+ and CD8α− CD11c+ subsets. 1.5 × 106 sorted CD11c+ cells were placed in 96-well plates in complete RPMI 1640 medium and irradiated with 2,900 rad. Assessment of the polarization of antigen-specific CD4+ T cells by DCs ex vivo was adapted from a previously described method (60). LCMV GP53-60-specific CD4+ T cells isolated from TCR transgenic SMARTA mice (52) were purified by negatively depleting CD8−, B220−, and CD11b−expressing cells with Dynal beads and, after enrichment of CD4+ T cells, with CD4-MACS beads (purity > 98%). 6 × 104 CD4+ cells were added to DCs in the presence or absence of 1 μg/ml GP53-60 peptide. Supernatants were isolated 5 d later and analyzed for the presence of IL-10 and IFN-γ by ELISA, as reported earlier in this paper.

Statistical analysis. Statistical analyses were performed using the Student’s t test: *P < 0.01; **P < 0.001; and ***P < 0.0001.

This work was supported by National Institutes of Health grant AI 58105, awarded to M.G. von Herrath.

The authors have no conflicting financial interests.

Submitted: 11 July 2006
Accepted: 8 September 2006

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