T cell depletion is commonly used in organ transplantation for immunosuppression; however, a restoration of T cell homeostasis following depletion leads to increased memory T cells, which may promote transplant rejection. The cytokine IL-7 is important for controlling lymphopoiesis under both normal and lymphopenic conditions. Here, we investigated whether blocking IL-7 signaling with a mAb that targets IL-7 receptor α (IL-7Rα) alone or following T cell depletion confers an advantage for allograft survival in murine transplant models. We found that IL-7R blockade alone induced indefinite pancreatic islet allograft survival if anti–IL-7R treatment was started 3 weeks before graft. IL-7R blockade following anti-CD4– and anti-CD8–mediated T cell depletion markedly prolonged skin allograft survival. Furthermore, IL-7 inhibition in combination with T cell depletion synergized with either CTLA-4Ig administration or suboptimal doses of tacrolimus to induce long-term skin graft acceptance in this stringent transplant model. Together, these therapies inhibited T cell reconstitution, decreased memory T cell numbers, increased the relative frequency of Tregs, and abrogated both cellular and humoral alloimmune responses. Our data suggest that IL-7R blockade following T cell depletion has potential as a robust, immunosuppressive therapy in transplantation.
IL-7 receptor blockade following T cell depletion promotes long-term allograft survival

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Introduction

T cell depletion by antibodies is one of the most potent immunosuppressive therapies and is increasingly used as an induction therapy in organ transplantation (1). However, T cell homeostasis after depletion therapy leads to a predominance of memory T cells (1–3), which are more potent than naive T cells in mediating graft rejection and present as a major obstacle to achieving tolerance. Mice undergoing T cell homeostatic proliferation following depletion therapy rejected cardiac allograft despite costimulatory blockade by CTLA-4Ig, a treatment capable of inducing tolerance in nondepleted mice (4, 5). In human, kidney transplant patients who had received T cell depletion therapy by high-dose alemtuzumab, but no maintenance immunosuppression, uniformly developed acute rejection within the first month after transplantation (6), a period during which there was still a severe T cell lymphopenia but most of the remaining T cells were effector memory T cells (7).

T cell reconstitution after depletion therapy comprises de novo thymopoiesis and homeostatic proliferation of remaining peripheral T cells, and both processes are IL-7 dependent (8, 9). IL-7 signals through the IL-7 receptor (IL-7R) which is composed of 2 chains, the common γ chain and the α chain (IL-7Rα or CD127) (10). IL-7 plays an essential, nonredundant role in lymphopoiesis, since IL-7– or IL-7Rα knockout mice have severe T and B cell lymphopenia (11, 12) and infants with IL-7Rα mutations have severe T cell lymphopenia necessitating bone marrow transplantation (13). IL-7 has also been shown to be necessary for the homeostatic proliferation of both naive and memory CD4+ and CD8+ T cells in lymphopenic conditions (14–18). Therefore, in the setting of organ transplantation, the blockade of IL-7/IL-7R signaling is expected to prolong the effects of T cell depletion therapy, reduce the number of memory T cells, and increase immunoregulation, leading to better graft acceptance (19).

In this study, we investigated the role of IL-7R blockade by an anti–IL-7Rα mAb, first given alone in an islet allograft model and then given after T cell depletion by a combination of anti-CD4 and anti-CD8 mAbs in a more stringent skin allograft model. We also elucidated the mechanisms underlying the therapeutic efficacy of IL-7R blockade in transplantation.

Results

IL-7R blockade reduces almost all lymphocyte subset numbers and increases Treg frequency. The anti–IL-7Rα mAb (A7R34) used in our study was previously shown to block IL-7R and reduce lymphocyte numbers when given at 2 mg every other day (qod) for 2 weeks (20). In this study, we tested a lower dose of A7R34 and were able to produce similar effects. Naïve BALB/c mice were injected with either PBS or A7R34 400 μg qod for 3 weeks and sacrificed. Anti–IL-7Rα–treated mice had significantly lower numbers of total lymphocytes, T cells, CD4+ T cells, CD8+ T cells, and B cells in the LNs, spleen, and peripheral blood and drastically reduced numbers of thymocytes in the thymus compared with control mice (Supplemental Figure 1, A–D; supplemental material available online with this article; doi:10.1172/JCI66287DS1). Interestingly, we found a significant increase in the percentage of CD4+ T cells expressing programmed death 1 (PD-1) and an increase in the percentage of CD4+CD25+FOXP3+ Tregs in the LNs and spleens of treated mice compared with those of control mice (Supplemental Figure 1, A and B), in concordance with recent publications (21, 22). On
the other hand, we found that mice sacrificed 4 days after having received A7R34 at a dose as high as 2 mg/d had not yet exhibited significant reduction in lymphocyte numbers (data not shown), indicating that A7R34 does not have important lytic activity, also in agreement with studies from other groups (21, 23).

IL-7R blockade alone started 3 weeks before graft induces islet allograft tolerance and abrogates both cellular and humoral alloimmune responses. We next investigated the effects of IL-7R blockade in a fully mismatched islet allograft model in which C57BL/6 islets were transplanted into BALB/c mice previously rendered diabetic by streptozotocin (STZ). Anti–IL-7Rα mAb given at 400 μg qod from the day of graft (D0) until rejection did not significantly prolong graft median survival time (MST) compared with no treatment (29 vs. 21 days; \( P = 0.16 \)). Being guided by the aforementioned experiment in naive mice, we gave anti–IL-7Rα mAb from 3 weeks before grafts (D–21) to posttransplant day 90 (PTD90) and obtained indefinite graft survival of more than 180 days in 5 of 6 treated mice (\( P = 0.0002 \) compared with no treatment) (Figure 1A). Subsequent left nephrectomy to remove the islet grafts in tolerant mice led to a prompt increase in blood glucose (data not shown), confirming that the maintenance of normoglycemia was due to graft function.

Untreated and some treated mice that received anti–IL-7R from D–21 to PTD30 were sacrificed a few days after graft rejection and on PTD70, respectively, for immunological analyses. Using mixed lymphocyte reaction (MLR) with 3H-thymidine incorporation, we showed that the treatment of islet graft recipients with anti–IL-7R from D–21 significantly reduced T cell proliferation in response to stimulation by donor splenocytes (B), but did not alter T cell responses to third party splenocytes (C). (D) IFN-γ ELISPOT showed that mice treated with anti–IL-7R from D–21 had significantly fewer IFN-γ–secreting T cells in response to stimulation by donor splenocytes. (E) Mice treated with anti–IL-7R from D–21 had lower levels of DSAs. (F) BALB/c nude mice underwent islet grafts from C57BL/6 donors and then received 5 × 10^6 of either T cells from BALB/c mice that had been treated with anti–IL-7R from D–21 and become tolerant to C57BL/6 islet grafts or T cells from naive BALB/c mice (\( n = 4 \) for each groups). T cells from tolerant mice were less effective in inducing islet graft rejection than those from naive mice. *\( P < 0.05 \); **\( P < 0.01 \); ***\( P < 0.001 \).

Figure 1

IL-7R blockade started 3 weeks before graft induces islet allograft tolerance and abrogates both cellular and humoral alloimmune responses. (A–E) BALB/c mice previously rendered diabetic by STZ received C57BL/6 islet graft and were treated as indicated. (A) Anti–IL-7Rα mAb given from D–21 to either PTD90 or PTD30 induced indefinite graft survival in most or all of mice. (B and C) MLR with 3H-thymidine incorporation showed that the treatment of islet graft recipients with anti–IL-7R from D–21 significantly reduced T cell proliferation in response to stimulation by donor splenocytes (B), but did not alter T cell responses to third party splenocytes (C). (D) IFN-γ ELISPOT showed that mice treated with anti–IL-7R from D–21 had significantly fewer IFN-γ–secreting T cells in response to stimulation by donor splenocytes. (E) Mice treated with anti–IL-7R from D–21 had lower levels of DSAs. (F) BALB/c nude mice underwent islet grafts from C57BL/6 donors and then received 5 × 10^6 of either T cells from BALB/c mice that had been treated with anti–IL-7R from D–21 and become tolerant to C57BL/6 islet grafts or T cells from naive BALB/c mice (\( n = 4 \) for each groups). T cells from tolerant mice were less effective in inducing islet graft rejection than those from naive mice. *\( P < 0.05 \); **\( P < 0.01 \); ***\( P < 0.001 \).
Lymphocyte phenotyping performed on PTD180 showed that lymphocyte reconstitution had occurred in tolerant mice (Supplemental Figure 2). In order to test whether tolerance can be transmissible, T cells purified from the spleen of tolerant mice sacrificed on PTD180 were adoptively transferred to T cell–deficient BALB/c nude mice that had received islet grafts from C57BL/6 donors. Without T cell transfer, BALB/c nude mice accepted allograft indefinitely. The transfer of T cells from naive wild-type BALB/c to BALB/c nude mice that had previously received C57BL/6 islet graft 2 weeks earlier led to graft rejection in 3 of 4 mice, whereas the same experiment using T cells from tolerant mice only led to graft rejection in 1 of 4 mice. However, the small sample size (n = 4 for each group) precluded statistical analysis (Figure 1P).

**IL-7R blockade following T cell depletion strongly prolongs skin allograft survival.** We next tested the effects of IL-7R blockade in a much more stringent model of fully allogeneic skin allograft from C57BL/6 donors to BALB/c recipients. Anti–IL-7R mAb given alone either from D0 or D–21 until rejection did not result in a meaningful prolongation of graft MST compared with no treatment (10 vs. 9.5 days; P = 0.37 and 12 vs. 9.5 days; P = 0.0006, respectively) (Figure 2). Because IL-7R blockade is expected to inhibit T cell reconstitution after T cell depletion therapy, we then tested whether the addition of an anti–IL-7R mAb to a T cell depletion protocol could prolong skin graft survival. We treated recipient mice first with a combination of 2 depleting mAbs, anti–CD4 and anti–CD8 in order to rapidly deplete T cells, followed by either isotype control or anti–IL-7R mAb 400 μg qod from PTD1 until rejection. Mice having received anti–CD4 and anti–CD8 followed by isotype control (referred to as “depletion alone”) had graft MST of 30 days (range: 19–35 days) (P < 0.0001 compared with untreated mice). Mice having received anti–CD4 and anti–CD8 followed by anti–IL-7R mAb (referred to as “depletion plus anti–IL-7R”) had MST of 58 days (range: 36 to >90 days) which were significantly longer than that of the “depletion alone” group (P < 0.0001) (Figure 2).

**IL-7R blockade following T cell depletion delays T cell reconstitution and favors immune regulation.** First of all, we compared the total T cell and T cell subset numbers among 4 groups of skin graft recipients having received no treatment, anti–IL-7R alone from D0, anti–IL-7R alone from D–21, and T cell depletion by anti–CD4 and anti–CD8, followed by anti–IL-7R and found that a profound reduction of T cells, including CD4+FOXP3+ effector T cells such as achieved with the last regimen was necessary for the prolongation of skin graft survival (Supplemental Figure 3).

We next focused our analyses on the 2 groups of mice treated by depletion alone and depletion plus anti–IL-7R and compared the lymphocyte subset numbers and frequencies between these 2 groups on PTD35. At this time point, mice that received depletion alone had rejected their grafts, whereas grafts were well preserved in mice under the cover of depletion plus anti–IL-7R. We found that IL-7R blockade following T cell depletion profoundly inhibited T cell reconstitution. Mice with depletion plus anti–IL-7R had a 10- to 20-fold reduction in the number of total T cells, CD4+ and CD8+ T cells in the LNs, spleen, and peripheral blood compared with mice with depletion alone (all P < 0.05) (Figure 3, A and B, and Supplemental Figure 4, A and B). B cell numbers were also reduced but to a lesser extent. The effect of IL-7R blockade on the thymus was even more pronounced (Supplemental Figure 4C). The fact that anti–CD4 and anti–CD8 mAbs did not deplete B cells (Supplemental Figure 4D) whereas IL-7R blockade in mice reduced both T and B cell numbers explains why B cell number was reduced in the group with depletion plus anti–IL-7R compared with depletion alone.

Despite a decrease in their absolute numbers, the relative frequencies of CD4+FOXP3+ Tregs were increased in the depletion plus anti–IL-7R group compared with the depletion alone group (22.3% ± 7.7% vs. 10.5% ± 0.9% and 27.4% ± 3.5% vs. 14.4% ± 0.8% of CD4+ T cells in the LNs and spleen, respectively; both P < 0.05). As a result, the Treg/T effector ratio defined by CD3+CD4+FOXP3+/CD3+CD4+FOXP3− was increased (0.30 ± 0.13 vs. 0.12 ± 0.02 and 0.38 ± 0.07 vs. 0.17 ± 0.01 in the LNs and spleen, respectively; both P < 0.05). Of note, CD4+FOXP3+ Tregs comprised CD4+CD25+FOXP3+ and CD4+CD25−FOXP3+ T cells; the percentage of both populations was increased by IL-7R blockade (Figure 3A). Similarly, the percentage of memory T cells defined as CD3+CD62L−CD44hi among total T cells was increased by IL-7R blockade (Figure 3B). These data suggest that this immunoregulatory mechanism is not specific to Treg, but rather affects all memory T cells.

**Blockade of PD-L1/PD-1 pathway enhances IL-7R blockade.** To test whether the addition of an anti–PD-L1 mAb to the depletion plus IL-7R blockade regimen would enhance the overall effects of this treatment, we treated mice with depletion alone, depletion plus anti–IL-7R, and depletion plus IL-7R and anti–PD-L1 mAbs. Without PD-L1 treatment, the addition of IL-7R alone prolonged graft survival compared with no treatment (30 vs. 9.5 days; P < 0.0001). The addition of IL-7R and anti–PD-L1 mAbs prolonged graft survival compared with depletion alone (58 vs. 30 days; P < 0.0001).
anti–IL-7R treatment in an autoimmune diabetes model (21, 22) and to accelerate rejection in several transplant models (26). Surprisingly, the addition of anti–PD-L1 to our treatment protocol did not alter graft survival (Figure 4), suggesting that this pathway is not crucial for the prolongation of skin graft survival by IL-7R blockade. An alternative explanation is that, under the cover of IL-7R blockade following T cell depletion, a treatment with multiple mechanisms of action, a contributive role of the PD-1 signaling pathway may become less visible. On the other hand, Treg depletion shortened skin graft MST from 58 days to 48 days (Figure 4), although the difference only had a borderline statistical significance ($P = 0.06$) perhaps due to small sample size ($n = 5$). This trend suggests that the increase in Treg relative frequency may be one of the underlying mechanisms of IL-7R blockade that favors skin graft survival. 

**IL-7R blockade following T cell depletion abrogates both cellular and humoral alloimmune responses.** IFN-$\gamma$ ELISPOT revealed significantly fewer donor-specific IFN-$\gamma$-secreting T cells in skin graft recipients treated with depletion plus anti–IL-7R compared with those treated with depletion alone (29 ± 26 vs. 203 ± 76 spots/10$^5$ T cells; $P < 0.0005$) (Figure 5A). Similarly, MLR with 3H-thymidine incorporation showed a very weak donor-specific T cell response in mice treated with depletion plus anti–IL-7R, whereas there was...
a strong T cell proliferation in mice treated with depletion only (245 ± 55 vs. 11410 ± 6456 counts per minute; P < 0.0001) (Figure 5B). Of note, unlike in the islet graft model, T cell proliferation in response to third party splenocytes was also inhibited in mice treated with depletion plus anti–IL-7R (data not shown). Cytokine measurement in the MLR supernatant by ELISA also showed a complete abrogation of IL-2, TNF, and IL-17 secretion by T cells from mice with depletion plus anti–IL-7R compared with mice with depletion only (8.0 ± 4.4 vs. 79.8 ± 38.1, 16.8 ± 8.3 vs. 140.1 ± 25.5, and 2.5 ± 2.2 vs. 62.0 ± 33.8 pg/ml, respectively; all P < 0.01) (Figure 5C). Together, these data show that T cells that have escaped depletion may become anergic to donor APCs and suggest that the effects of IL-7R blockade are not restricted to depletion. The humoral alloimmune response was also abrogated, as evidenced by a significant reduction in DSA levels in mice treated with depletion plus anti–IL-7R compared with mice treated with depletion only (616 ± 31 vs. 4101 ± 1877; P < 0.0001) (Figure 5D).

Because IL-7R blockade following T cell depletion leads to a decrease in memory T cell numbers and an inhibition of alloimmune responses, we wonder whether this treatment is effective in alloimmunized mice and whether it would also affect the antiviral immune responses acquired through previous infections or vaccinations. To address the first question, we performed skin grafts from C57BL/6 donors to BALB/c recipients without treatment; as expected, all grafts were rejected within 10 days. One month after the first graft, recipients were rechallenged with a second C57BL/6 skin graft and divided into 3 groups: depletion plus anti–IL-7R, depletion alone, and no treatment. Although we also observed a statistically significant prolongation of graft survival in the group treated by depletion plus anti–IL-7R compared with depletion alone or no treatment (MST = 15, 11, and 6 days, respectively; P < 0.05) (Supplementary Figure 5), graft survival was far shorter than that obtained in nonpreimmunized recipients (Figure 2), suggesting that IL-7R blockade following T cell depletion is less effective when recipients have been primed by a first graft.

To address the second question, we used a model of viral infection in which naïve BALB/c mice received 2 i.m. injections of 1010 infectious particles of an adenovirus vector serotype 5 (AdV5) at 4 weeks apart. Two weeks after the second injection, mice received PBS for 5 weeks, anti–IL-7Rα mAb for 5 weeks, or T cell depletion by 2 injections of anti-CD4 and anti-CD8 mAbs followed by anti–IL-7Rα mAb for 7 weeks and then were sacrificed for analyses. ELISA showed no difference in anti-adenovirus antibody titers among the 3 groups of mice (Supplemental Figure 6A). IFN-γ ELISPOT also revealed no difference in the frequency of antianenval memory T cells between the control group and the group that received anti–IL-7R alone for 5 weeks (Supplemental Figure 6B), suggesting that a short course of IL-7R blockade by itself does not impair previously acquired antiviral immunity. However, T cell depletion by anti-CD4 and anti-CD8 followed by anti–IL-7R for 7 weeks led to a decrease in the frequency of antianenvalo memory T cells, as expected in the case of a strong immunosuppressive regimen. Therefore, IL-7R blockade following T cell depletion is likely to be more suitable for an induction rather than a long-term maintenance therapy in transplantation.

T cell targeting is critical for the efficacy of anti–IL-7R therapy in the prolongation of skin graft survival. Our aforementioned data demonstrated that IL-7R blockade exerts its effects through targeting T cells; however, at least in mice, B cells are also affected by IL-7R blockade. Therefore, we further clarify the role B cells in our treatment protocols. We performed skin grafts from BALB/c donors to C57BL/6 recipients, which were treated with T cell depletion by a combination of anti-CD4 and anti-CD8 mAbs, followed by anti–IL-7Rα mAb (DEP + a–a–IL-7R). Some recipients were also treated with either an anti–PD-L1 mAb to block the PD-1/PD-L1 signaling pathway or an anti–CD25 mAb to deplete Tregs. As also shown in Figure 2, treatment with DEP + a–IL-7R resulted in graft MST of 58 days; the addition of anti–PD-L1 did not alter graft survival (MST = 57 days; P = 0.89 compared with DEP + a–IL-7R), whereas the addition of anti–CD25 shortened graft MST to 48 days (P = 0.06 compared with DEP + a–IL-7R).

![Figure 4](http://www.jci.org)
there was little inflammatory infiltration in the skin grafts from mice with depletion plus anti–IL-7R (Figure 7A). Concordantly, gene expression measurement using TaqMan Low Density Array (TLDA) revealed several-fold decreases in the expression of proinflammatory genes in the skin grafts of mice treated with depletion plus anti–IL-7R compared with those of mice treated with depletion alone. Among these genes were genes coding Th1 cytokine (IFN-γ and TNF), Th2 cytokines (IL-4, IL-5, and IL-13), chemokines (CCL-2, CCL-3, CCL-5, CXCL-10, and CXCL-11), costimulatory molecules (CD40, CD86, and ICOS), granzyme B, and perforin (Figure 7B and Table 1).

By PTD70, however, most skin grafts under treatment by depletion plus anti–IL-7R had also been rejected. Lymphocyte phenotyping at the time of graft rejection revealed that all lymphocyte subset numbers in all lymphoid organs were still very low, comparable to those observed earlier on PTD35 (Supplemental Figure 7). The skin is well known for its immunogenicity (reviewed in ref. 28) and contains abundant Langerhans cells, whose development is not IL-7 dependent (29). Therefore, skin grafts may be a preferred site for T cell homing, explaining why they are still rejected despite the persistence of lymphopenia.

We next performed immunohistology of skin grafts of mice treated by depletion plus anti–IL-7R at the time of rejection to see which types of cells participate in the rejection process. For comparison, we also did immunohistological staining of accepted skin grafts under treatment by depletion plus anti–IL-7R on PTD35 as well as rejecting skin grafts of untreated control mice. As shown in Supplemental Figure 8, accepted skin grafts of treated mice on PTD35 had little inflammatory infiltration. When rejection occurred, skin grafts of treated mice were characterized by an influx of leukocytes including macrophages, granulocytes, and T cells. However, the leukocyte infiltration in the rejecting skin grafts of treated mice was still less intense than that in untreated mice. We also quantified the expression of inflammatory genes in rejecting and accepted skin grafts and found a similar tendency. In mice treated by IL-7R blockade following T cell depletion, skin grafts undergoing rejection displayed an increase in the expression of inflammatory gene transcripts compared with accepted skin grafts on PTD35 (Supplemental Figure 9A). However, the levels of inflammatory gene expression in rejecting skin grafts of treated mice were still significantly lower than those of untreated control mice (Supplemental Figure 9B).

IL-7R blockade synergizes with treatment protocols combining T cell depletion with either CTLA-4Ig or low-dose tacrolimus. In order to further prolong skin graft survival, we added either CTLA-4Ig (abatacept) or low-dose tacrolimus to the treatment protocols. CTLA-4Ig alone or low-dose tacrolimus alone only slightly prolonged graft MST (11 days for either treatment compared with 9.5 days for no treatment; \( P < 0.01 \)). CTLA-4Ig combined with T cell depletion by anti-CD4 and anti-CD8 prolonged MST to 28.5 days, and the addition of anti–IL-7R mAb to this treatment protocol further increased MST to 52.5 days (\( P < 0.05 \) ) (Figure 8A).

The synergistic effect was more pronounced when low-dose tacrolimus was used instead of CTLA4-Ig. Two-thirds of mice treated with a combination of T cell depletion, low-dose tacrolimus from the day of graft, and anti–IL-7R had skin graft survival of at least 90 days, whereas mice treated with the same protocol but without IL-7R blockade had MST of only 32.5 days (\( P < 0.001 \) ) (Figure 8B). We next investigated whether we could shorten the duration of anti–IL-7R treatment to facilitate immune reconstitution. At the same time, we also started low-dose tacrolimus from PTD12 instead of D0 to reduce immunosuppression in the early phase of lymphodepletion. We found that 80% of mice treated with IL-7R blockade following T cell depletion and low-dose tacrolimus from PTD12 had graft survival of at least 90 days whether anti–IL-7R mAb was stopped on
PTD70 or PTD35, indicating that a short course of anti–IL-7Rα mAb is sufficient to prolong graft survival in this drug combination. On the contrary, mice treated with T cell depletion and low-dose tacrolimus from PTD12 but without anti–IL-7Rα mAb had skin graft MST of only 33 days ($P < 0.005$) (Figure 8B). Importantly, lymphocyte phenotyping on PTD90 showed that the shorter course of anti–IL-7R treatment was associated with a better recovery of thymic function and a higher number of T and B cells in the LNs (Figure 8C), thereby limiting the potential overimmunosuppression associated with prolonged lymphopenia.

**Discussion**  
Inhibition of IL-7 or IL-7R by specific blocking mAbs has been successfully tested in models of autoimmune diseases (21, 22, 30–32) and graft-versus-host disease following bone marrow transplantation (33). In the context of organ transplantation, there is a single report in which an anti-mouse IL-7 polyclonal Ab had no effect when given alone, but prolonged heart graft survival when combined with CD40/CD40L costimulatory blockade (34).

Using an islet allograft model, we demonstrated that tolerance could be achieved when anti–IL-7Rα mAb was given alone from 3 weeks before graft (Figure 1A). Because this blocking mAb does not have a rapid lytic effect, it must take a few weeks to obtain a significant reduction in lymphocyte numbers (Supplemental Figure 1). However, once established, tolerance persisted despite lymphocyte reconstitution after the end of anti–IL-7R treatment (Supplemental Figure 2) and tolerant mice developed donor-specific hyporesponsiveness (Figure 1, B–D). Furthermore, T cells from tolerant mice did not induce islet graft rejection in T cell–deficient nude mice as effectively as T cells from naive mice (Figure 1F). Taken together, these data indicate that immunoregulation was established after a short course of IL-7R blockade. However, this model might overestimate the efficacy of IL-7R blockade because STZ has recently been demonstrated to induce lymphopenia associated with an increase in Treg frequency and produce immunosuppressive effects (35).

Therefore, we next used a more stringent model of fully mismatched skin allograft and demonstrated that the addition of IL-7R blockade to T cell depletion therapy inhibited T cell reconstitution and doubled graft MST (Figure 2). The mechanisms underlying the prolongation of skin graft survival using this treatment protocol are multifactorial. The most evident effect of IL-7R blockade to T cell depletion therapy inhibited T cell reconstitution on allospecific memory T cell formation and activation as demonstrated by IFN-γ ELISPOT and MLR (Figure 5, A and B). Moreover, T cells from mice that accepted their skin graft under IL-7R blockade secreted much less Th1 and Th17 cytokines (Figure 5C), a finding in concordance with previous studies using models of autoimmune diseases (21, 22, 30). Similarly, accepted skin grafts under IL-7R blockade expressed much lower levels of proinflammatory cytokines (Figure 7B and Table 1).

The increase in Treg/T effector ratio associated with IL-7R blockade is likely to contribute to graft acceptance because Treg depletion by anti–CD25 mAb tended to shorten skin graft survival (Figure 4). The role of Tregs in the prolongation of graft survival under lymphopenic conditions is well known; for example, the adoptive transfer of exogenous Tregs to mice treated with costimulatory blockade after having undergone subtotal T cell depletion restored heart graft tolerance (5). IL-7–deprived Tregs were also shown to maintain their suppressive functions (22).

Since IL-7R blockade in mice affects both T and B cells, whereas in human, B cell number is not decreased by IL-7Rα mutations, we further investigated the role of B cells in our treatment protocols. Using B cell–deficient muMT mice as skin allograft recipients, we demonstrated that IL-7R blockade following T cell depletion prolonged graft survival whether B cells were present or absent (Figure 2 and Figure 6A). On the other hand, B cell depletion by anti–CD20 mAb in wild-type mice, although not sufficient to prolong skin graft survival, led to a significant decrease in DSA levels (Figure 6D).

Taken together, our data show that IL-7R blockade after T cell depletion prolongs skin allograft survival through different mechanisms, including inhibition of T cell reconstitution, inhibition of the formation and activation of allospecific memory T cells, and induction of immunoregulation associated with an increase in Treg/T effector ratio. On the other hand, in mice, the reduction in B cell

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**Table 1**  
List of genes significantly suppressed in the skin grafts of mice treated with depletion plus anti–IL-7R (A) compared with depletion + isotype Ig (I)

<table>
<thead>
<tr>
<th>Genes</th>
<th>Log2 FC (A/I)</th>
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<td>Il13</td>
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<tr>
<td>Il4</td>
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</tr>
<tr>
<td>Il1b</td>
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<tr>
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<td>H2-Ea</td>
<td>-1.6</td>
</tr>
<tr>
<td>H2-Eb1</td>
<td>-1.5</td>
</tr>
<tr>
<td>Stat1</td>
<td>-1.2</td>
</tr>
<tr>
<td>Cd86</td>
<td>-1.2</td>
</tr>
</tbody>
</table>

This table reports the relative expression of the genes shown in Figure 7B. FC, fold change.
number caused by IL-7R blockade may contribute to the decrease in DSA and attenuate the B cell–mediated component of rejection. Finally, combination treatment with other immunosuppressants further enhances graft survival in this stringent skin allograft model. IL-7R blockade following T cell depletion synergized with either CTLA-4Ig or suboptimal doses of tacrolimus (Figure 8, A and B). The latter protocol is likely more effective because tacrolimus is more potent than other drugs in inhibiting memory T cell activation (7), whereas memory T cells have been shown to be less dependent on costimulation for activation (36, 37).

However, lymphopoiesis in human and in mice has several differences. Under normal conditions, the maintenance of the T cell pool in adult humans and in mice is sustained almost exclusively by peripheral T cell division and by thymic output, respectively (38). But after T cell depletion therapy, the increased thymic output does contribute to immune reconstitution in adult transplant patients (1). We and others found that T cell numbers returned to normal about 3 months after a brief course of T cell depletion by anti-CD4 and anti-CD8 mAbs in mice (39), whereas it may take up to 3 years for lymphocyte numbers in the blood of kidney transplant patients treated by alemtuzumab or thymoglobulin induction to return to baseline levels (40, 41).

Nevertheless, mouse skin allograft is a predictive preclinical model frequently used in the early phase of development of new immunosuppressive drugs in transplantion (42–44). The skin graft survival obtained in our study is remarkable because, without bone marrow transplantation, not many treatment protocols could result in similar graft outcomes, especially when fully allogeneic BALB/c and C57BL/6 mice are used for donors and recipients (43, 44). Therefore, our results suggest that IL-7R blockade may present as a promising therapeutic candidate in organ transplantation. Thoroughly designed treatment protocols combining an anti–IL-7R mAb with other immunosuppressants should harness the beneficial effects of IL-7R blockade to promote allograft acceptance.

Methods

Mice and allograft models. Eight-week-old male C57BL/6 (H-2b), BALB/c (H-2d), and SJL (H-2d) mice were purchased from Centre d’Elevage Janvier. Six-week-old male BALB/c nude (H-2d) and B cell–deficient muMT mice (B6.129S2-Ighmtm1Cgn/J) on a C57BL/6 genetic background were purchased from Charles River France. To prepare for islet transplantation, BALB/c or BALB/c nude recipients received 1 i.p. injection of 250 mg/kg of STZ 5 to 10 days before graft to induce diabetes, which is defined as a nonfasting blood glucose greater than 17 mmol/l (300 mg/dl) on at least 2 occasions. Pancreases were harvested from C57BL/6 donors, digested by collagenase, and islets purified by Ficoll gradient centrifugation. Each diabetic BALB/c mouse received about 500 islets under the left renal capsule. Graft functioning and rejection diagnosis were determined when blood glucose decreased to less than 10 mmol/l (180 mg/dl) and increased again to greater than 17 mmol/l, respectively. For the skin allograft model, tail skin from donors was grafted to the dorsal trunk of recipients and protected by bandage for 6 days. The skin graft was then inspected every day and rejection was defined as graft necrosis of more than 80% of the surface.

mAbs and treatment protocols. Some mAbs were produced in our lab using the following hybridomas: rat anti-mouse IL-7R (A7R34), CD4 (GK1.5), CD8 (2.43), and CD25 (PC61). A7R34 was a gift from Shin-Ichi Nishikawa (Laboratory for Stem Cell Biology, RIKEN Center for Developmental Biology, Kobe, Japan), and other hybridomas were purchased from LGC Standards. Rat anti-mouse PD-L1 (10F9G2) was purchased from Bio X Cell, and rat anti-mouse CD20 (5D2) was provided by Genentech. A rat anti-dinitrophenol (DNP) mAb (LO-DNP-61) was purchased from the Laboratory of Experimental Surgery, Universite Catholique de Louvain (Brussels, Belgium) and used as rat IgG2a isotype control for the anti–IL-7R mAB. Human CTLA-4Ig (Abatacept; Bristol Myers Squibb) and tacrolimus (Astellas) were purchased from the pharmacy of CHU de Nantes. All treatments were given i.p. Anti-CD4 and anti-CD8 were given in combination on days –3 and –1 before skin graft at 200 μg per injection. CTLA-4Ig was given at 500 μg once per week from the day of skin graft (D0) to the day of rejection or to PTD70 if there was no rejection. Low-dose tacrolimus was given at 0.5 mg/kg/d from the day of skin graft or 0.7 mg/kg/d from...
PD12; in either case, it was continued to the day of rejection or to the end of the experiment if there was no rejection. Anti-IL-7R α mAb was given at 400 µg qod; the time and duration of treatment is specified for each experiment in Results. Of note, A7R34 has been effectively used for a relatively long period of time (31), suggesting that the anti-drug Abs that might develop in recipients do not abrogate its effects.

Lymphocyte phenotyping by flow cytometry. Mice were euthanized under general anesthesia. Blood was drawn from cardiac puncture; spleen, thymus, and mesenteric LNs were harvested. Whole-blood leukocyte phenotyping was performed using BD TruCOUNT Tubes (BD Biosciences) according to the manufacturer’s instructions. Cells from LNs, spleen, and thymus were isolated using BD Falcon Cell Strainers (BD Biosciences). The following anti-mouse antibodies were used for flow cytometry: anti-CD3, anti-CD4, anti-CD8, anti-CD19, anti-CD44, anti-CD45-2, anti-CD45R (B220), anti-CD62L, anti-CD127, anti–Gr-1 (Ly-6G and Ly-6C), anti-NKG2D, anti–PD-1 and anti-FOXP3; all were purchased from BD Biosciences except anti-CD8, anti-CD19, anti-CD44, anti–CD45-2, anti-CD45R (B220), anti-CD62L, which was purchased from eBiocience, and anti-CD127 (A7R34), which was prepared in our lab. Flow cytometry was performed using a BD LSRII flow cytometer.

Measurement of DSA. Blood of recipient mice was obtained either by retroorbital sinus sampling or by cardiac puncture in the case of sacrifice. Plasma of recipients was diluted at 1/20 and incubated for 30 minutes with an FITC-conjugated goat anti-mouse Ig (11 to 14) or anti–IL-7Rα mAb (A1 to A4). (A) Representative photomicrographs of skin graft histology on PTD21 (H&E). Original magnification, ×200. (B) Unsupervised clustering of differentially expressed genes in skin grafts on PTD21 using mouse immune panel TLDA. The heat map represents normalized and color-coded relative expression values (2−ΔΔCq). Red values indicate overexpression, and green values indicate underexpression.
II and autoMACS Pro Separator (Miltenyi Biotec). IFN-γ ELISPOT was performed in which each well contained 10^5 T cells, 10^5 non-T cells, and 0.3 nmol/peptide/ml of a pool of overlapping 15-mer peptides covering the complete sequence of the hexon protein of adenovirus 5 (Peptivator – AdV5, Miltenyi Biotec). To quantify antiadenoviral antibodies, ELISA plates were coated overnight with 10^9 virus particles per well, washed, and incubated consecutively with serial dilutions of mouse sera (1/500 to 1/10,000), peroxidase-conjugated donkey anti-mouse IgG (Jackson ImmunoResearch), TMB substrate, and H2SO4 to stop the reaction, and then read using an ELISA plate reader.

Statistics. Statistics were done using the GraphPad Prism 5.0 software (GraphPad Software). Data were presented as mean ± SD unless otherwise stated. Graft survivals were reported as MST and analyzed by the Kaplan-Meier method using the log-rank test. Other values were compared using the Mann Whitney test. P < 0.05 was considered significant.

Study approval. All animal studies were approved by the institutional review board of INSERM Unit 1064.

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