Depletion of FOXP3⁺ regulatory T cells promotes hypercholesterolemia and atherosclerosis

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Atherosclerosis is a chronic inflammatory disease promoted by hyperlipidemia. Several studies support FOXP3-positive regulatory T cells (Tregs) as inhibitors of atherosclerosis; however, the mechanism underlying this protection remains elusive. To define the role of FOXP3-expressing Tregs in atherosclerosis, we used the DEREG mouse, which expresses the diphtheria toxin (DT) receptor under control of the Treg-specific Foxp3 promoter, allowing for specific ablation of FOXP3⁺ Tregs. Lethally irradiated, atherosclerosis-prone, low-density lipoprotein receptor–deficient (Ldlr−/−) mice received DEREG bone marrow and were injected with DT to eliminate FOXP3⁺ Tregs. Depletion of Tregs caused a 2.1-fold increase in atherosclerosis without a concomitant increase in vascular inflammation. These mice also exhibited a 1.7-fold increase in plasma cholesterol and an atherogenic lipoprotein profile with increased levels of VLDL. Clearance of VLDL and chylomicron remnants was hampered, leading to accumulation of cholesterol-rich particles in the circulation. Functional and protein analyses complemented by gene expression array identified reduced protein expression of sortilin-1 in liver and increased plasma enzyme activity of lipoprotein lipase, hepatic lipase, and phospholipid transfer protein as mediators of the altered lipid phenotype. These results demonstrate that FOXP3⁺ Tregs inhibit atherosclerosis by modulating lipoprotein metabolism.

Introduction
Cardiovascular disease remains a major cause of death and morbidity despite currently available drug therapies (1). Atherosclerosis constitutes the underlying cause of the clinical manifestations of myocardial infarction, stroke, and gangrene, and is recognized as a chronic inflammatory disease involving a variety of immunological features (2). Subendothelial lipid retention and oxidative modification in the arterial wall elicit a chronic inflammatory process characterized by the formation of an immune cell infiltrate dominated by lipid-laden macrophage foam cells and T cells (2–4). The accumulation and activation of these immunological cells play a pivotal role in the development of atherosclerotic lesions.

Both innate and adaptive immunity are involved in the pathogenesis of atherosclerosis.Pattern recognition receptors of innate immunity account for cholesterol uptake and contribute to the activation of macrophages and endothelial cells. Antigen-specific T cells recognizing LDL in the intima constitute the adaptive immunity component in the pathogenesis of atherosclerosis and elicit proinflammatory stimuli that further exacerbate and propagate this disease (5). Inherent atheroprotective immune mechanisms include natural antibodies to LDL-derived phospholipids and antiinflammatory immune cells such as regulatory T cells (Tregs). Strategies to mobilize and stimulate immunosuppression may provide novel therapeutic approaches to reduce atherosclerotic cardiovascular disease (6, 7).

Tregs were initially characterized as CD4⁺CD25⁺ T cells that could suppress the activity of effector T cells (8). Further studies have identified several varieties of Tregs (9). The forkhead box transcription factor, FOXP3, was identified as the key lineage marker and master switch in the regulation of Treg development and function (10–12). FOXP3 is now accepted as the “gold standard” for defining thymic-derived Tregs (9). Accordingly, selective depletion of FOXP3⁺ Tregs reproduces the hyperinflamed, scurfy phenotype observed in Treg-deficient mice (13). In contrast, depletion of CD4⁺CD25⁺ T cells by administration of an antibody targeting...
Effective depletion of transgenic Treg cells is achieved in chimeric DEREG/Ldlr–/– mice. (A) Proportions of cells expressing eGFP within the CD3+CD4+ population in inguinal lymph nodes from chimeric DEREG/Ldlr–/– mice treated with DT or PBS harvested at 4 and 8 weeks, respectively (n = 6–7 mice per group). (B) Proliferation of lymph node cells from chimeric DEREG/Ldlr–/– mice treated for 8 weeks with DT or PBS and stimulated in vitro with anti-CD3. Data show stimulation index based on 3H-thymidine uptake (n = 4 mice per group). *P < 0.05.

Figure 1

Effective depletion of transgenic Tregs in peripheral blood for more than 3 days after i.p. administration of a single dose (0.125 μg/mouse) of DT to DEREG mice (Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI63891DS1). Based on these findings, a twice-weekly DT administration regimen was chosen for all subsequent experiments. To evaluate the long-term safety in DEREG mice, DT was administered i.p. twice weekly for 8 weeks to DEREG and wild-type mice at 0.125 or 0.25 μg/mouse (20 g body weight). The higher DT dose was found to be associated with signs of toxicity (fatigue, changes in fur appearance, loss of appetite), while no obvious adverse effects were observed at the lower dose in DEREG or wild-type mice. Accordingly, the low DT dose was chosen for long-term administration in the atherosclerosis study.

Long-term DT treatment is tolerated in DEREG mice. We achieved effective depletion of 99% of transgenic eGFP+ Tregs in peripheral blood for more than 3 days after i.p. administration of a single dose (0.125 μg/mouse) of DT to DEREG mice (Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI63891DS1). Based on these findings, a twice-weekly DT administration regimen was chosen for all subsequent experiments. To evaluate the long-term safety in DEREG mice, DT was administered i.p. twice weekly for 8 weeks to DEREG and wild-type mice at 0.125 or 0.25 μg/mouse (20 g body weight). The higher DT dose was found to be associated with signs of toxicity (fatigue, changes in fur appearance, loss of appetite), while no obvious adverse effects were observed at the lower dose in DEREG or wild-type mice. Accordingly, the low DT dose was chosen for long-term administration in the atherosclerosis study.

DT administration eliminates functional Treg in chimeric DEREG/Ldlr–/– mice. DEREG bone marrow was transplanted into lethally irradiated Ldlr–/– mice that received an atherogenic diet to promote hypercholesterolemia and atherosclerosis. One group of the chimeric DEREG/Ldlr–/– mice received DT twice weekly for 8 weeks to eliminate FOXP3+ Tregs. The FOXP3-driven coordinate expression of eGFP and hDTR permitted an identification of FOXP3+ cells in bone marrow chimeras not receiving DT and an evaluation of the efficiency of Treg depletion in the DT-treated mice.

Effective depletion of transgenic (eGFP+) Tregs in inguinal lymph nodes (Figure 1A) and lymphoid organs (not shown) was achieved by DT administration to chimeric DEREG/Ldlr–/– mice. The suppressor function of total Tregs was markedly impaired, as documented by robustly increased proliferation of lymph node cells from mice treated with DT versus PBS (Figure 1B).

In a separate experiment, chimeric DEREG/Ldlr–/– mice were treated with DT versus PBS and bled at different intervals after initiation of treatment. Flow cytometric data on the kinetics of Tregs revealed that over time, depletion of eGFP+ Tregs became less effective. Ten days after initiating DT administration, complete depletion of transgenic Tregs was achieved, followed by a gradual recovery of these GFP+FOXP3+ cells. The total number of FOXP3+ cells was not substantially reduced to a compensatory increase in GFP FOXP3+ cells (Supplemental Figure 1, B–D). Plasma levels of the inflammatory marker, serum amyloid A (SAA), did not differ between groups (Table 1). DT treatment thus led to persistently perturbed immunoregulation with increased T cell activity, though without signs of overt systemic inflammation after 8 weeks of treatment.

To define the source of eGFP– (nontransgenic) FOXP3+ Tregs, we transplanted DEREG bone marrow (on a C57Bl/6 background, thus expressing the CD45.2 isofrom) into Ldlr–/– mice carrying the CD45.1 surface marker. FACS analysis of spleen cells showed that eGFP– (nontransgenic) FOXP3+ Tregs were predominantly derived from the CD45.1+ recipient Ldlr–/– mice (Supplemental Figure 2, A and B). Furthermore, nontransgenic CD45.1+ eGFP– cells pre-
dominated among FOXP3+ cells in the DT-treated mice, whereas CD45.2+ eGFP+ cells constituted the majority of FOXP3+ cells in PBS controls. This suggests a replenishment of the total FOXP3+ Treg pool from this niche that could only partially compensate for the loss of functional FOXP3+ Tregs upon DT treatment (Supplemental Figure 4D). Gel filtration analysis revealed a pronounced elevation of cholesterol in the VLDL/chylomicron remnant (CMR) fraction in DT-treated chimeric DEREG/Ldlr−/− mice (Figure 4B), whereas triglyceride content was lower in the VLDL/CMR particles (Figure 4C). Plasma VLDL/CMR cholesterol levels strongly correlated with lesion size (r = 0.956; P = 0.0006), as did total cholesterol and LDL cholesterol (Figure 4D and Supplemental Figure 4, A and B). Upon DT treatment, plasma apoB concentration was numerically increased, suggesting an increased number of VLDL particles rather than a larger amount of cholesterol per particle (Table 1). DT treatment affected neither body weight nor plasma markers, nor liver or kidney function (data not shown). DT-mediated elimination of Tregs from DEREG/Ldlr−/− mice was associated with markedly increased monocyte counts in peripheral blood (Table 1). No difference was detected in the distribution of monocyte subpopulations (Table 1). Interestingly, total monocyte counts in peripheral blood showed a positive correlation with lesion size (Supplemental Figure 4C), and an inverse correlation was found with GFP+ Tregs (Supplemental Figure 4D).

### Table 1

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Wild-type bone marrow into Ldlr−/−</th>
<th>DEREG bone marrow into Ldlr−/−</th>
<th>P value</th>
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<tr>
<td>Group alias</td>
<td>PBS (A)</td>
<td>DT (B)</td>
<td>PBS (C)</td>
</tr>
<tr>
<td>Body weight (g)</td>
<td>25.3 ± 0.9</td>
<td>25.1 ± 0.9</td>
<td>25.7 ± 0.6</td>
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<td>Total cholesterol (mM)</td>
<td>20.8 ± 1.6</td>
<td>22.8 ± 1.7</td>
<td>15.3 ± 1.5</td>
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<tr>
<td>Triglycerides (mM)</td>
<td>4.2 ± 0.5</td>
<td>3.4 ± 0.7</td>
<td>3.5 ± 0.3</td>
</tr>
<tr>
<td>ApoB (μg/dl)</td>
<td>n.d.</td>
<td>n.d.</td>
<td>328.1 ± 30.7</td>
</tr>
<tr>
<td>Serum amyloid A (pg/ml)</td>
<td>n.d.</td>
<td>n.d.</td>
<td>200 ± 71</td>
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<tr>
<td>Leukocytes (10⁶/μl)</td>
<td>13.9 ± 1.3</td>
<td>16.2 ± 1.3</td>
<td>12.3 ± 0.7</td>
</tr>
<tr>
<td>Monocytes (10⁶/μl)</td>
<td>1.5 ± 0.3</td>
<td>1.2 ± 0.4</td>
<td>0.8 ± 0.2</td>
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<tr>
<td>Ly6Chi/monocytes (%)</td>
<td>n.d.</td>
<td>n.d.</td>
<td>47.2 ± 4.5</td>
</tr>
<tr>
<td>Lymphocytes (10⁶/μl)</td>
<td>10.7 ± 1.0</td>
<td>12.8 ± 1.2</td>
<td>9.1 ± 0.9</td>
</tr>
<tr>
<td>Erythrocytes (10⁶/μl)</td>
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<td>9.4 ± 0.3</td>
<td>9.2 ± 0.3</td>
</tr>
<tr>
<td>Platelets (10³/μl)</td>
<td>1,012 ± 88</td>
<td>1,003 ± 91</td>
<td>1,156 ± 95</td>
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</table>

Mean ± SEM. All groups (Kruskal-Wallis test) followed by pairwise comparisons (Mann-Whitney U test) if P < 0.05 indicated by *, †A vs. C; ‡B vs. D; §C vs. D; n.d., not determined.

The increase in cholesterol was detectable already during the first month of treatment, suggesting that it was causally related to the increase in lesion size observed after 8 weeks (Figure 4A).

Plasma lipids were analyzed in 4 treatment groups of chimeric mice: recipient Ldlr−/− mice receiving wild-type bone marrow and treated with (a) PBS or (b) DT; or Ldlr−/− mice receiving DEREG bone marrow and treated with (c) PBS or (d) DT. Of note, DT treatment had no significant effect on atherosclerotic lesions (Figure 2C), total cholesterol, triglycerides, body weight, or hematological parameters (Table 1). This corroborates previous work showing that lipid levels in Apoe−/− mice are not affected by DT administration (23).

**Treg depletion impairs lipoprotein catabolism.** To determine the effects of Treg depletion on VLDL biosynthesis in the liver, we analyzed plasma lipids after administering Triton WR-1339, which inhibits lipoprotein lipase–dependent (LPL-dependent) lipoprotein catabolism. Cholesterol levels in plasma rose to a similar extent in both groups (Figure 5A), suggesting that the elevated VLDL levels were not due to increased biosynthesis in the liver.

VLDL catabolism was assessed after i.v. injections of FITC-labeled mouse VLDLs. This analysis revealed significantly impaired...
Figure 2
Depletion of transgenic Tregs aggravates atherosclerosis. (A) Representative photomicrographs showing Oil Red O- and H&E-stained sections from the proximal aorta of chimeric DEREG/Ldlr–/– mice treated for 8 weeks with PBS or DT. (B) Quantification of lesion size in cross-sections of the proximal aorta at different levels from the valves; n = 11–12 (PBS) mice and n = 5–8 (DT) mice, respectively. (C) Relative lesion area (lesion area/area inside external elastic lamina × 100) calculated from 4 sections per mouse (300–600 μm) for wild-type/Ldlr–/– and DEREG/Ldlr–/– mice treated for 8 weeks with PBS or DT, respectively. *P < 0.05; **P < 0.01. Scale bars: 100 μm.

Discussion
Regulatory immunity has been implicated in atherosclerosis, but the mechanism and target organ(s) of atheroprotection remain largely...
unknown. We now show in the chimeric DEREG/Ldlr–/– model that:
(a) depletion of FOXP3+ Treg leads to substantially increased atherosclerosis; (b) elimination of FOXP3+ Tregs aggravates hypercholesterolemia; and (c) VLDLs and CMRs are dramatically elevated in plasma due to reduced clearance in the absence of the LDL receptor. Together, these findings establish that FOXP3+ Tregs exert an important atheroprotective effect, and show that plasma lipidprotein metabolism is modulated in part by regulatory immunity.

Previous studies demonstrated that regulatory immunity affects atherosclerosis progression. Elimination of CD4+CD25+ cells increases disease severity, and the transfer of T cells from Cd28–/– mice is associated with reduced numbers of CD4+CD25+ natural Tregs accelerating atherosclerosis in Ldlr–/– mice (16). Similarly, Ldlr–/– mice deficient in ICOS displayed decreased FOXP3+ T cells and increased atherosclerosis (25), while expansion of the circulating Treg pool in Apoe–/– mice by CD31 administration reduced atherosclerosis (26), as did the adoptive transfer of CD4+CD25+ cells from atherosclerotic to disease-prone mice (27). Furthermore, expansion of another regulatory T cell population, the Tr1 cell population, by global or antigen-specific methods, also reduces atherosclerosis in Apoe–/– mice (28, 29). While all these studies highlight the role of regulatory cellular immunity in atherosclerosis, direct evidence for atheroprotective actions of FOXP3-driven immunity is lacking.

We now provide such evidence by using the DEREG model. We had expected that elimination of Tregs would unleash plaque inflammation, causing enhanced production of several proinflammatory mediators. Instead, immune cells and local inflammatory markers in the lesions were either unchanged or reduced when Tregs were depleted, which is likely attributable to the advanced stage of lesions with low cellularity in DT-treated mice. Circulating levels of the inflammatory marker SAA remained unaltered, and while monocyte counts were elevated, no changes were registered between monocyte subclasses. Together, these data suggest that modulation of vascular inflammation may not be the only, and perhaps not even the major, target of Treg control in atherosclerosis.

DT treatment of DEREG/Ldlr–/– mice profoundly reduced the population of GFP-FOX3+ Tregs. This led to a more than 3-fold

**Figure 3**
Depletion of Tregs affects cellular composition of atheroma. (A) Representative fluorescence micrographs depicting eGFP-FOX3+ cells (arrowheads) and eGFP-FOX3+ (arrows) cells in aortic lesions of DEREG/Ldlr–/– mice treated for 8 weeks with PBS or DT. Anti-GFP was labeled with AlexaFluor 488 (green), anti-FOX3 with AlexaFluor 555 (red), and nuclei with DAPI (blue). (B) Quantitation of immunohistochemical staining for T cells (CD3+), Tregs (FOX3+), and I-Aα-expressing (MHCII-expressing) cells (all expressed as stained cells per lesion area) in the proximal aorta of chimeric DEREG/Ldlr–/– mice treated for 8 weeks with DT or PBS; n = 7–8 mice per group. (C) Quantitation of immunohistochemical staining for macrophages (CD68+), dendritic cells (CD11c+), and expression of the adhesion molecule VCAM-1 (all expressed as stained area per lesion area) in the aortic sinus of chimeric DEREG/Ldlr–/– mice treated for 8 weeks with DT or PBS; n = 7–8 mice per group. (D) Cellularity of atherosclerotic lesions (DAPI+ nuclei per lesion area) in the proximal aorta of chimeric DEREG/Ldlr–/– mice treated for 8 weeks with PBS or DT; n = 7–8 mice per group. (E) Necrotic core area relative to total atherosclerotic lesion area in the proximal aorta of chimeric DEREG/Ldlr–/– mice treated for 8 weeks with PBS or DT; n = 7–8 mice per group. *P < 0.05; **P < 0.01.

Scale bars: 50 μm.
increase in effector T cell proliferation due to loss of suppressor activity. However, it did not cause any substantial reduction in the total number of FOXP3+ T cells. Bone marrow transplantation (BMT) experiments using CD45 isotypic markers clarified that this was due to an expansion of FOXP3+ T cells derived from the recipient. This, in turn, may reflect that FOXP3+ cells are less sensitive to irradiation used in the BMT protocol than other T cell types. In line with previous data (22), the robust effect of DT treatment on T effector cell proliferation shows that the recipient-derived FOXP3+ T cells were not effective as suppressor cells, possibly reflecting that they were not fully differentiated Tregs.

In addition to these changes in FOXP3+ cell populations, it was apparent that the depletion of transgenic FOXP3+ Tregs (GFP+ FOXP3+) waned over time despite continued DT treatment. This may reflect genetic alterations in the artificial Foxp3 promoter, an increasingly efficient DT metabolism in the recipients, or immune reactions to DT. At any rate, our data show that selective depletion of Tregs over an extended period is possible by using the DEREG/DT strategy; this permits studies of chronic pathological processes such as atherosclerosis.

The phenotypic screen of DEREG/Ldlr−/− chimeras revealed increased hypercholesterolemia upon Treg depletion, with a 1.7-fold increase in total plasma cholesterol. This was due to a dramatic increase in large, cholesterol-rich, VLDL/CMR-sized particles. Since such particles are highly atherogenic in Ldr−/− mice (30), it is likely that their accumulation in blood crucially contributed to the accelerated atherosclerosis observed when Tregs were depleted. This notion is supported by the strong correlation between VLDL cholesterol concentration in plasma and lesion size. This, in turn, suggests that Tregs under normal conditions control VLDL cholesterol levels and that this contributes to their atheroprotective role.

The clearance of injected VLDLs and CMs was reduced in Treg-ablated DEREG/Ldr−/− mice. In contrast, neither lipoprotein secretion nor mRNA expression of enzymes involved in the VLDL biosynthesis pathway was affected by Treg depletion. Therefore, Tregs modulate blood lipid levels by regulating lipoprotein catabolism.

VLDLs and CM particles are catabolized in a complex process that involves hydrolysis of triglycerides by several lipases includ-
Figure 5
VLDL/CMR lipoprotein catabolism is impaired in Treg-depleted mice. (A) Biosynthesis of apoB-containing plasma lipoproteins. Data show plasma cholesterol levels in chimeric DEREG/Ldlr<sup>−/−</sup> mice treated for 8 weeks with DT or PBS followed by i.v. administration of Triton WR-1339 to inhibit lipoprotein lipase–dependent VLDL catabolism; n = 9 (PBS) and n = 7 (DT), respectively. (B) Clearance of injected FITC-VLDL in chimeric DEREG/Ldlr<sup>−/−</sup> mice treated for 8 weeks with DT or PBS. FITC-derived fluorescence was analyzed in plasma samples at the indicated time points. Data for each individual were normalized to the fluorescence of plasma taken 1 minute after injection; n = 4 per group. (C) In vivo turnover of CM particles injected into chimeric DEREG/Ldlr<sup>−/−</sup> mice treated for 8 weeks with DT or PBS. Data show kinetics of the CM [14C] retinol core particle clearance from blood and are expressed as radiolabeled moieties corrected for weight; n = 5 (PBS); n = 7 (DT). (D) CM [14C] retinol uptake in the liver. Data expressed as radiolabeled moieties corrected for weight; n = 5 (PBS); n = 7 (DT). *P < 0.05.

In summary, we provide evidence that Treg depletion results in significantly accelerated atherosclerosis and an atherogenic lipoprotein profile that is mediated by reduced clearance of large lipoprotein particles. Our findings highlight that immune mechanisms have an impact on metabolic events and show that chronic inflammation can promote cardiovascular disease by instigating metabolic disturbances.

Methods

Animals and treatment protocols. DEREG mice on a C57BL/6 background were previously generated (13) and were maintained and used as hemizygous transgenics. Offspring were typed by flow cytometry for the presence of an eGFP<sup>+</sup> cell population within the CD3<sup>+</sup>CD4<sup>+</sup> population of peripheral blood cells. Bone marrow cells, isolated from the femur and tibia of male 8- to 10-week-old DEREG or DEREG<sup>−/−</sup> littersmates were i.v. injected (7.5 × 10<sup>6</sup> cells per mouse) into lethally irradiated (14 Gy) 6- to 8-week-old Ldlr<sup>−/−</sup> mice (The Jackson Laboratory). After recovery for 6 weeks, Ldlr<sup>−/−</sup> mice were fed a high-cholesterol diet (1.25% cholesterol, 0% cholate; Research Diets). Mice were randomly assigned to treatment with DT (6.25 g/kg body weight, i.p. injection 2 times per week; Merck, no. 322326) or control (100 μl of endotoxin-free PBS containing 0.1% BSA). Following 8 weeks of treatment, the mice were anesthetized by CO<sub>2</sub> asphyxiation and their organs were harvested. The treatment groups comprised:

Group A: Wild-type bone marrow injected into Ldlr<sup>−/−</sup> mice treated for 8 weeks with PBS; i.p. injection.
Group B: Wild-type bone marrow injected into Ldlr<sup>−/−</sup> mice treated for 8 weeks with DT; i.p. injection.
Group C: DEREG bone marrow injected into Ldlr<sup>−/−</sup> mice treated for 8 weeks with PBS; i.p. injection.
Group D: DEREG bone marrow injected into Ldlr<sup>−/−</sup> mice treated for 8 weeks with DT; i.p. injection.

For certain studies regarding the origin of FOXP3<sup>+</sup> cells, Ldlr<sup>−/−</sup> mice were crossed with CD45<sup>1/1</sup> mice (B6.SJ-Ly5.1<sup>−/−</sup> mice, stock-nr: 002014, The Jackson Laboratory) to generate Ldlr<sup>−/−</sup>/CD45<sup>1/1</sup> mice which then received bone marrow from DEREG mice on a regular C57BL/6 background (CD45<sup>2/2</sup>). Staining for CD45.1/CD45.2 was performed as with other cell surface markers using antibodies directed against CD45.1 and CD45.2 (eBioscience).

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Figure 6
Gene expression array analysis shows changes in liver transcriptomes upon Treg depletion. Shown here are gene expression heat maps of global Affymetrix gene expression array analysis of liver RNA from DEREG/Ldlr−/− mice. The rows correspond to genes and the columns to individual mice treated with DT or PBS, respectively. Relative gene expression is shown based on normalization for each gene in the DT and PBS-treated groups. Color coding indicates increased gene expression in red, and decreased expression in blue compared with the other group, respectively. Sets of genes involved in functional annotation clusters as defined by the DAVID annotation analysis are grouped accordingly. Arbitrary titles that summarize the functional role of displayed genes in a cluster are shown in parentheses. Clusters show differentially expressed transcripts based on an FDR less than 0.1 and an absolute linear fold change greater than 1.5.
Tissue preparation and histological analysis. To evaluate atherosclerotic lesion size, a total of 8 cryosections (10 μm) extending cranially from the aortic valves at 100-μm intervals were prepared, stained with Oil Red O and hematoxylin, and analyzed using Leica Q500MC image analysis software (19). Lesion area per cross-section and relative area of lesions were quantified and the results expressed as the mean of 4 consecutive sections per mouse available for all individual mice. In brief, relative lesion area was calculated for each section as F% = (100 × L/A), where L is lesion area and A is area inside the external elastic lamina. F% is averaged over all levels analyzed (300–600 μm² above aortic cusps), and the mean is calculated for each treatment group. This method eliminates artifacts caused by oblique sections. Necrotic core was expressed as acellular area per lesion area.

Immunolocalization studies. Primary antibodies used for immunohistochemistry comprised anti-CD68 (AbD Serotec), anti–VCAM-1 and anti–I-Ab (both BD Biosciences), anti-CD3 (Southern Biotechnology), and anti-FOXP3 (eBioscience); all of them were rat IgG to mouse antigens. Isotype-matched antibodies were used as controls. Antibodies were titrated to optimum performance and applied to acetone-fixed cryosections followed by detection using an ABC peroxidase kit and DAB substrate (both from Vector Laboratories). Immunofluorescence was performed on aortic sections as described (13). Quantification of stainings was documented as the ratio of thresholded chromogen to fluorogen area or number of stained cells per intimal lesion area and nucleated-positive (DAPI) area. Quantitative image analysis was performed independently by 2 investigators blinded to the study protocol using either Quantimed (Leica) or analySIS (Olympus) software.

Flow cytometry. Flow cytometry was performed on leukocytes isolated from blood, spleen, or inguinal lymph nodes. Primary labeled antibodies

Figure 7
Treg depletion modulates expression of genes controlling inflammation and lipid metabolism in the liver. Quantitative real-time RT-PCR analysis of liver mRNA from DEREG/Ldlr⁻/⁻ mice treated for 8 weeks with DT or PBS. Signals were normalized to Hprt. *P < 0.05; n = 6 per group. (A) Ifng, (B) Tnfa, (C) Cd3, (D) Apoc3, (E) Apob, (F) Dgat, (G) Lpl, (H) Lipc, (I) Vldlr, (J) Ptp, (K) Sort1, and (L) Atf3.
were used for extracellular staining against CD3, CD4, and CD45 isoforms CD45.1 and CD45.2 (all from BD Biosciences) and CD25 (eBioscience). Intracellular staining was performed using the anti–mouse/rat FOXP3 Staining Set (eBioscience), rabbit anti-GFP (Invitrogen), and anti-FOXP3 (eBioscience). The staining procedure to detect monococyte subclases was performed essentially as described (39). Briefly, cells were incubated for 15 minutes at room temperature with FcBlock (anti–CD16/CD32; eBioscience) and fixable Aqua LIVE/DEAD stain (Invitrogen). Subsequently, the cells were washed and incubated for 45 minutes at room temperature with the following antibodies: anti-Ly6G, anti-CD49b, anti-CD19, anti-CD3, anti NK1.1 (all PE-conjugated), anti–CD11b-PECy7, and anti–Ly6C-FITC (all from eBioscience). All samples were washed, fixed in 1% formalin, and analyzed on a CyAn ADP Analyzer using Summit version 4.3 software (both Beckman Coulter).

**Functional immune assay.** Inguinal lymph node cells were isolated by meshing lymph nodes from individual mice through a 100 μm nylon mesh followed by osmotic lysis of red blood cells (EL buffer; QIAGEN). Cells were washed in PBS, cultured in duplicate in a 96-well plate at 5 × 10^5 cells per well in IMDM containing 5% FCS, and incubated in the absence or presence of rat anti-mouse CD3 antibody (1 μg/ml; SouthernBiotech) for 72 hours. During the last 18 hours, [3H]-thymidine (1 μCi/ml) was added for each milligram of protein in VLDLs and incubated at room temperature for 2 hours. Conjugates were separated from free fluorochrome by gel filtration using a PD10 column (GE Healthcare) eluted with PBS. FITC conjugation was evaluated by absorption spectroscopy at 495 nm against an FITC standard curve. Protein concentration was determined by the Bradford assay (Bio-Rad) and adjusted to 2 mg/ml. Of this preparation, 100 μg of FITC (Sigma-Aldrich) was dissolved in DMSO (1 mg/ml) was added for each milligram of protein in VLDLs and incubated at room temperature for 2 hours.

**Hematology and plasma biochemistry.** Whole blood (EDTA) was analyzed by a Scil Vet abc hemocounter (Scil). Clinical biochemistry profiling of plasma was made on a Vitros DT60II Chemistry System (Ortho Clinical Diagnostics, Johnson & Johnson Co.) as described previously (40).

**Lipoprotein analysis.** Lipoprotein profiles were analyzed using a micro-FPLC column (30 × 0.32 cm Superose 6B; GE Healthcare) coupled to a system for online separation and subsequent detection of cholesterol (41). Plasma cholesterol and triglycerides were measured using enzymatic colorimetric kits (Randox Laboratories) according to the manufacturer’s protocol. ApoB concentrations were measured by a commercial ELISA following the manufacturer’s instructions (AlerCHEK).

**Analysis of lipoprotein catabolism and biosynthesis.** For in vivo clearance, VLDLs (d = 1.006 – 1.019 g/ml) were isolated by ultracentrifugation from pooled plasma of Apoe–/– mice in the presence of 2 mM benzamidine, 0.5 mM PMSF, and 0.1 U/ml aprotinin. After isolation, VLDLs were dialyzed against PBS containing 1 mM EDTA and labeled using a modified version of a previously described method (42). In brief, 2 mg/ml VLDL was dialyzed overnight against 0.5 M NaHCO3 pH 9.5. Next, 50 μg of FITC (Sigma-Aldrich) dissolved in DMSO (1 mg/ml) was added for each milligram of protein in VLDLs and incubated at room temperature for 2 hours. Conjugates were separated from free fluorochrome by gel filtration using a PD10 column (GE Healthcare) eluted with PBS. FITC conjugation was evaluated by absorption spectroscopy at 495 nm against an FITC standard curve. Protein concentration was determined by the Bradford assay (Bio-Rad) and adjusted to 2 mg/ml. Of this preparation, 100 μl was injected into the tail vein, and blood samples were collected into EDTA-coated tubes 1, 5, 15, 30, and 60 minutes after injection. After centrifugation (1,500 g for 5 minutes), plasma samples were diluted 1:20 in PBS and fluorescence (excitation 485 nm, emission 535 nm) was analyzed using a Fluoroskan Ascent FL plate fluorosence reader (Thermo Scientific). Data presented are normalized to the fluorescence measured in samples obtained at 1 minute (100%).

VLDL synthesis was assessed after irreversibly blocking VLDL catabolism (43). Briefly, fasted mice received Triton WR-1339 (500 mg/kg i.v. tyloxapol; Sigma-Aldrich) using a 10% (w/v) solution and were bled from the tail into EDTA-coated tubes at the indicated time points. Triglycerides and cholesterol were measured in plasma as described above.

For in vivo measurements of CM clearance, uncirculated double-labeled ([3H]-retinol ester and [14C]-triacylglycerol) CMs were isolated from rat intestinal lymph through cannulation of the thoracic duct and handled.
as described elsewhere (44). Chimeric DEREG/Ldlr−/− mice treated for 8 weeks with DT or PBS and fed a high-cholesterol diet were injected with CMs corresponding to 2 mg triglycerides through the tail vein, and blood was collected after 1, 3, 7, and 15 minutes from the retro-orbital vein plexus in temporarily isoflurane-anesthetized animals. After the final bleeding, the mice were sacrificed and their tissues were collected and rinsed in PBS. Blood samples were extracted for esterified lipids as previously described (44), traced for the radioabeled moieties, and corrected for weight. Tissue samples were homogenized in chloroform/methanol (2:1) using tubes with ceramic beads, total lipids were extracted, and the tissues were adjusted for weight.

LPL and HL were released into the plasma with heparin injections (50 U/kg), and postheparin blood was collected from the tail vein after 15 minutes. Plasma was collected by ultracentrifugation and stored at -80°C. LPL activity was assessed in a system using [1H]-labeled triolein in an emulsion with Intralipid 10% composition as the enzyme substrate. Rat serum was added as apoCII donor (45). HL was inhibited using antibodies against rat HL (46). Specific HL activity was measured using a gum arabic–stabilized emulsion (47). Postheparin plasma lipase activity was determined as the mean of triplicate readings. Lipase activity is expressed as mU/ml, where 1 mU corresponds to 1 nmol of fatty acid released per minute at 25°C (48). Phospholipid transfer activity of PLTP was measured with a radiometric assay as described by Jauhiainen et al. (49) using an exogenous, lipoprotein-independent phospholipid transfer assay with labeled phospholipid vesicles as a substrate.

**RNA isolation.** mRNA was isolated from liver, intestine, heart, white adipose tissue, skeletal muscle, spleen, and lung using the RNeasy kit from QIAGEN. Total RNA quality was analyzed by an Agilent 2100 Bioanalyzer (Agilent Technologies) and quantified using a Nanodrop 1000 (Thermo Fisher Scientific).

**Gene expression array.** mRNA of liver and intestine was submitted to a core facility (Affymetrix Core Facility, Novum, Huddinge, Sweden and Uppsala Array Platform, Uppsala University, Sweden) for hybridization to GeneChip Mouse Gene 1.0 ST Arrays (Affymetrix). The microarray expression data were preprocessed with R statistical software (50) using the RMA method as implemented in the Affymetrix package. Differential expression analysis was performed with the moderated t statistic and false discovery rate (FDR) multiple test correction methods from the limma package. Transcripts with an FDR less than 0.1 and an absolute linear fold change greater than 1.5 were considered significantly differentially expressed. The raw microarray expression data have been deposited in NCBI’s Gene Expression Omnibus (Gene Expression Omnibus accession numbers GSE6279 for liver and GSE39351 for intestine; http://www.ncbi.nlm.nih.gov/geo). The differentially expressed genes were clustered according to their annotation using the DAVID Bioinformatics Resource to identify significantly enriched functional annotation terms in the categories of gene ontology and pathways (51).

**Real-time PCR.** Superscript-II and random hexamers (both from Invitrogen) were used for reverse transcription. Real-time PCR used TaqMan Universal Master Mix and premanufactured primers and probes (Assay-on-Demand) for genes of interest, with hypoxanthine guanine ribonucleosyltransferase (Hprt) used as the housekeeping gene, and was run on an ABI 7900 HT Fast RT-PCR system (all Applied Biosystems). Data were analyzed on the basis of the relative expression method with the formula: 2−ΔΔCT, where ΔΔCT = ΔCT (sample) − ΔCT (calibrator = average C(T) values of all samples), and ΔCT is the CT of the Hprt housekeeping gene subtracted from the C(T) of the target gene (52).

**Western blot analysis.** Total proteins were isolated from a 25-mg liver sample using T-PER (Thermo Scientific) and homogenized with glass homogenizers. Protein concentration was determined by the Bradford assay and adjusted to 1.5 mg/ml. Thirty-microgram samples were diluted with 2× Laemmli sample buffer, boiled, separated by electrophoresis using 4%–15% Mini-PROTEAN TGX Precast Gels, and transferred to a PVDF membrane (all Bio-Rad). Sortilin-1 was detected using a rabbit polyclonal anti-sortilin antibody (ab16640, Abcam), followed by peroxidase-labeled goat anti-rabbit IgG (Dako), and visualized using an ECL system (GE Healthcare). Rabbit anti-tubulin antibody (ab4074, Abcam) was used as a loading control. Quantification was performed using the ImageJ software (NIH) gel analyzer tool.

**Statistics.** Values are expressed as mean ± SEM unless otherwise indicated. Groupwise comparisons (n = 2) were performed using a nonparametric Kruskal-Wallis test followed by a Mann-Whitney U test as appropriate. Pairwise comparisons were performed using a nonparametric Mann-Whitney test unless otherwise indicated. Spearman’s rank correlation test was performed to assess correlations. Statistics performed for the arrays are described in that section. A P value less than 0.05 was considered significant. All experiments reported here were repeated at least twice with similar results.

**Study approval.** All animal experiments were approved by the Stockholm Regional Board for Animal Ethics.

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