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TAP1-Deficiency Does Not Alter Atherosclerosis Development in Apoe \(^{-/-}\) Mice

Daniel Kolbus\(^{1}\), Irena Ljungcrantz\(^{1}\), Ingrid Söderberg\(^{1}\), Ragnar Alm\(^{1}\), Harry Björkbacka\(^{1}\), Jan Nilsson\(^{1}\), Gunilla Nordin Fredrikson\(^{1,2,}\)

1 Department of Clinical Sciences, Skane University Hospital Malmö, Lund University, Malmö, Sweden
2 Faculty of Health and Society, Malmö University, Malmö, Sweden

Abstract

Antigen presenting cells (APC) have the ability to present both extra-cellular and intra-cellular antigens via MHC class I molecules to CD8\(^{+}\) T cells. The cross presentation of extra-cellular antigens is reduced in mice with deficient Antigen Peptide Transporter 1 (TAP1)-dependent MHC class I antigen presentation, and these mice are characterized by a diminished CD8\(^{+}\) T cell population. We have recently reported an increased activation of CD8\(^{+}\) T cells in hypercholesterolemic Apoe \(^{-/-}\) mice. Therefore, this study included TAP1-deficient Apoe \(^{-/-}\) mice (Apoe \(^{-/-}\)/Tap1 \(^{-/-}\)) to test the atherogenicity of CD8\(^{+}\) T cells and TAP1-dependent cross presentation in a hypercholesterolemic environment. As expected the CD8\(^{+}\) T cell numbers were low in Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) mice in comparison to Apoe \(^{-/-}\) mice, constituting ~1% of the lymphocyte population. In spite of this there were no differences in the extent of atherosclerosis as assessed by en face Oil Red O staining of the aorta and cross-sections of the aortic root between Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) and Apoe \(^{-/-}\) mice. Moreover, no differences were detected in lesion infiltration of macrophages or CD3\(^{+}\) T cells in Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) compared to Apoe \(^{-/-}\) mice. The CD3\(^{+}\)CD4\(^{+}\) T cell fraction was increased in Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) mice, suggesting a compensation for the decreased CD8\(^{+}\) T cell population. Interestingly, the fraction of CD8\(^{+}\) effector memory T cells was increased but this appeared to have little impact on the atherosclerosis development. In conclusion, Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) mice develop atherosclerosis equal to Apoe \(^{-/-}\) mice, indicating a minor role for CD8\(^{+}\) T cells and TAP1-dependent antigen presentation in the disease process.

Introduction

According to the current view, oxidized LDL (oxLDL) within the vascular wall initiates an immune response leading to influx of phagocytes and T cells to the inflammatory site, which ultimately leads to a chronic inflammation and growth of atherosclerotic lesions [1,2]. Hypercholesterolemia promotes atherosclerosis and leads to an imbalance in pro- and anti-atherogenic T cell populations. Pro-atherogenic T cells specific for oxLDL are found in atherosclerotic lesions and circulation [3,4] and mice deficient in CD4\(^{+}\) T cells displayed decreased lesion size [5,6]. In contrast, Ellhage et al. reported increased lesion development in cd8\(^{-/-}\) mice [7] whereas Dansky et al. stated that T and B cells had a minor impact in atherosclerosis [8]. The balance between pro- and anti-atherogenic signals within the T cell population may determine disease development. The induction of regulatory CD4\(^{+}\) T cells is anti-atherogenic [9] while depletion leads to increased lesion development [10], highlighting the dual role of CD4\(^{+}\) T cells and that imbalance may accelerate atherosclerosis. Less focus has been on the role of CD8\(^{+}\) T cells in atherosclerosis. Ellhage et al. reported that aortic lesions of Apoe \(^{-/-}\) mice deficient in CD8\(^{+}\) T cells receiving chow diet did not differ in size compared to Apoe \(^{-/-}\) mice with a normal CD8\(^{+}\) T cell population [7]. In contrast CD8\(^{+}\) T cells were found to comprise up to 50% of the lymphocyte population in advanced human atherosclerotic lesions [11] and Apoe \(^{-/-}\) mice expressing β-galactosidase (β-gal) in aortic smooth muscle cells (SMCs) developed CD8\(^{+}\) T cell driven arteritis and atherosclerosis upon immunization with β-gal specific dendritic cells (DC) [12]. We recently conducted a study in hypercholesterolemic Apoe \(^{-/-}\) mice comparing the CD4\(^{+}\) and CD8\(^{+}\) T cell responses early in the disease process. There was an induction of CD8\(^{+}\)IL-4 and CD8\(^{+}\) IFN-γ cells in heart draining lymph nodes and spleen and increased proliferation of CD8\(^{+}\) splenocytes compared to CD4\(^{+}\) counterparts [13]. This prompted us to formulate the hypothesis that a hyperlipidemic diet leads to activation of effector CD8\(^{+}\) T cells that drive development of atherosclerotic plaques in a CD4\(^{+}\)-independent way. To test this, Tap1 \(^{-/-}\) mice [14] were crossed with Apoe \(^{-/-}\) mice to generate a hypercholesterolemic mouse model with low CD8\(^{+}\) T cell numbers. The antigen peptide transporter 1 (TAP1) together with TAP2 constitutes a MHC class I transporter maintaining the major pathway of MHC class I cross presentation [15], diminishing this pathway would possibly affect pro-atherogenic CD8\(^{+}\) T cell responses. Surprisingly, the Apoe \(^{-/-}\)/Tap1 \(^{-/-}\) mice developed lesions of the same size as the Apoe \(^{-/-}\)
mice, indicating that TAP1-deficiency has no or minor effect on atherosclerosis.

**Materials and Methods**

**Ethics Statement**

The Local Animal Care and Use Committee at Lund University approved (Permit numbers M159-07 and M153-10) the experimental protocol used in the study. All surgery was performed under anesthesia, and all efforts were made to minimize suffering.

**Animals**

Female apolipoprotein E deficient- and TAP1-deficient mice on a C57BL/6 background were purchased from Jackson Laboratories, USA and C57BL/6 (wild type, WT) mice were obtained from in house breeding. To generate double deficient mice, Apoe−/− and Tap1−/− mice were crossed to obtain parental genotypes. The number of animals used was 10–13 and 7–10 per experimental group for mice sacrificed at 26 and 14 weeks of age, respectively. All animals were controlled for respective genetical background by PCR assay. Tap1−/− and Apoe−/− Tap1−/− mice did not have detectable levels of Tap1. The animals were kept under controlled laboratory conditions in individually ventilated cages (IVC) and food and water were provided ad libitum. The Apoe−/− Tap1−/− mice did not display any obvious phenotypically differences in comparison to Apoe−/− mice such as body weight and blood lipids (table 1 and 2). Further, we recorded no premature death or abnormal behavior. Thus, they appeared healthy, but Tap1−/− mice had bald, non-fur covered areas in the back, which was not apparent in Apoe−/− Tap1−/−, Apoe−/− or WT mice. At the age of 6 weeks, diet was shifted from chow to high fat diet (HFD; 0.15% cholesterol and 21% fat (Lantmännen, Sweden)). The mice were 6 weeks, diet was shifted from chow to high fat diet (HFD; 0.15% cholesterol and 21% fat (Lantmännen, Sweden)). The mice were sacrificed at 8 or 20 weeks after diet change by intraperitoneal injection of 0.1% sodium pentobarbital (Argent, Solon, Ohio). The heart was collected and stored in Histochoice at −80°C until processing. Plasma was collected from cardiac puncture and stored at −80°C over night. The slides were incubated with 0.5% Triton-X 100 followed by incubation with H2O2 according to the MOMA-2 protocol. Slides were incubated for 30 minutes in 80% methanol (Merck Chemicals, Darmstadt, Germany) diluted in 5 ml PBS (pH 7.4), followed by removal of spleen and lymph nodes and thereafter immersion of 5 ml of the fixative Histochoice (Arnesco, Solon, Ohio). The aorta was then dissected free of connective tissue and fat, cut longitudinally, mounted en face and stored in Histochoice. The heart was collected and stored in Histochoice at 4°C until processing. Plasma was collected from cardiac puncture and stored at −80°C until assayed. In a second set of experiments female mice (6 Apoe−/− mice and 6 Apoe−/− Tap1−/− mice) were sacrificed 22 weeks after diet change. However, histochoice was not used, lymphoid organs were not assayed and the heart was snap frozen in liquid nitrogen and stored at −80°C.

**Analysis of plaque macrophage, T cell and neutral lipid content**

The heart was embedded in OCT (Optimal Cutting Temperature; Tissue-TekZoeterwoude, The Netherlands) and frozen sections of 10 μm were collected. All staining experiments were done at room temperature if not stated otherwise. Macrophage stained sections were fixed in ice-cold acetone for 10 minutes, incubated with 0.5% Triton-X100 (Merck Chemicals, Darmstadt, Germany) and 3% H2O2 (Apoteket AB, Sweden) for 5 minutes, each step separated by PBS rinsing. Thereafter the sections were blocked with 10% mouse serum in PBS for 30 minutes and quickly dipped in PBS before staining with a rat anti-mouse MOMA-2 antibody (monocyte/macrophage, BMA Biomedicals, Augst, Switzerland) diluted in 10% rat serum in PBS and incubated at +4°C over night. The slides were incubated with biotinylated rabbit anti-rat IgG (BA-4001, Vector Laboratories, Burlingame, CA) for 50 minutes followed by a 30-minute incubation with ABC solution (ABC elite, Vector Laboratories) and color development using the DAB detection kit (Vector Laboratories). For T cell detection, slides were fixed in acetone, washed and permeabilized using Triton-X100 followed by incubation with H2O2 according to the MOMA-2 protocol. Slides were incubated for 30 minutes in 10% goat serum followed by incubation with rabbit anti-human CD3 (cross-reacts with mouse CD3 [17]; A0452, Dako Cytomation, Fort Collins, CO) diluted in 2% goat serum at 4°C overnight. This was followed by a 50-minute incubation with biotinylated goat anti-rabbit IgG (BA-1000, Vector Laboratories) and the ABC/DAB procedure according to the MOMA-2 protocol. Omissions of the primary or secondary antibodies were used as negative controls. To analyze T cell subsets in aortic root lesions sections of snap frozen heart tissue were fixed in acetone for 15 minutes, rehydrated in PBS for 10 minutes and incubated with 0.3% H2O2 for 2 minutes followed by PBS wash. The sections were blocked with 10% mouse serum for 30 minutes followed by a 15 minute block with avidin (Thermo Scientific) and a 15 minute block with biotin (Thermo Scientific). The slides were washed with PBS and incubated with rat anti-mouse CD4 (553044, clone RM4-5, BD Pharmingen) or rat anti-mouse CD8a (553029, clone RM4-5, BD Pharmingen) diluted in 10% mouse serum for 30 minutes. Omission of the antibody was used as negative control.

**Table 1. Lesion size, weight, plasma cholesterol and triglycerides at 14 weeks of age.**

<table>
<thead>
<tr>
<th>Lesion size (mm² × 10^3)</th>
<th>WT</th>
<th>Tap1−/−</th>
<th>Apoe−/−</th>
<th>Apoe−/− Tap1−/−</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n/a</td>
<td>n/a</td>
<td>318.7 ± 93.9</td>
<td>350.1 ± 119.8</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Weight (g)</th>
<th>25.5 ± 4.4</th>
<th>20.6 ± 1.4</th>
<th>22.9 ± 2.4</th>
<th>21.3 ± 1.9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>86.8 ± 26.8</td>
<td>71.9 ± 15.3</td>
<td>604.6 ± 87.3</td>
<td>650.0 ± 92.7</td>
</tr>
<tr>
<td>Triglycerides (mg/dl)</td>
<td>25.7 ± 15.7</td>
<td>24.7 ± 7.1</td>
<td>42.6 ± 5.4</td>
<td>50.7 ± 9.4</td>
</tr>
</tbody>
</table>

| n/a, not applicable; |
| P<0.01 vs. WT; |
| P<0.001 vs. WT and Tap1−/−; |
| P<0.01 vs. Tap1−/−; |
| P<0.001 vs. WT and Tap1−/−. The number of animals in respective group was 7 (WT), 10 (Tap1−/−), 10 (Apoe−/−) and 7 (Apoe−/− Tap1−/−). In the triglyceride assay, one Apoe−/− Tap1−/− mouse displayed 3 times higher values than group mean and was therefore excluded from analysis. |

doi:10.1371/journal.pone.0033932.t001

**Table 2. Plasma cholesterol, triglycerides and weight at 26 weeks of age.**

<table>
<thead>
<tr>
<th>Cholesterol</th>
<th>WT</th>
<th>Tap1−/−</th>
<th>Apoe−/−</th>
<th>Apoe−/− Tap1−/−</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>104.3 ± 21.0</td>
<td>85.1 ± 11.6</td>
<td>792.6 ± 101.3</td>
<td>740.1 ± 134.6</td>
</tr>
<tr>
<td>Triglycerides</td>
<td>24.5 ± 18.8</td>
<td>33.8 ± 23.8</td>
<td>47.6 ± 17.7</td>
<td>54.0 ± 20.1</td>
</tr>
<tr>
<td>Weight</td>
<td>33.5 ± 6.6</td>
<td>22.2 ± 3.6</td>
<td>24.4 ± 3.1</td>
<td>22.5 ± 1.6</td>
</tr>
</tbody>
</table>

| P<0.01 vs. WT; |
| P<0.001 vs. Tap1−/−; |
| P<0.01 vs. Tap1−/−; |
| P<0.001 vs. Tap1−/−, Apoe−/− and Apoe−/− Tap1−/−. The number of animals in respective group was 12 (WT), 13 (Tap1−/−), 11 (Apoe−/−) and 13 (Apoe−/− Tap1−/−). |

doi:10.1371/journal.pone.0033932.t002
control. The color development procedure was as reported for the MOMA-2 protocol. For lipid staining, the sections were put in 0.24% Oil Red O diluted in 60% isopropanol for 10 minutes, rinsed with tap water to remove redundant Oil Red O, and dipped in 60% isopropanol followed by washing with dH2O before nuclear staining with hematoxylin for 15 seconds. En face (BioPix IQ 2.0, Göteborg, Sweden). copy and computer aided morphometry by a blinded observer.

Chemical and histochemical analyses were quantified by microscopy and computer aided morphometry by a blinded observer (BioPix IQ 2.0, Göteborg, Sweden).

**Cell preparation and flow cytometry**

**A. Mice analyzed at 26 weeks of age.** The spleen and mediastinal lymph nodes (MeLN) were meshed through a cell strainer (70 µm, BD Bioscience). The single cell suspension of the lymph nodes was washed in RPMI medium (Gibco, USA) and resuspended in complete medium (RPMI 1640 supplemented with 10% FCS, 1% Sodium pyruvate, 1% Hepes, 1% Penicillin/Streptomycin, 1% L-Glutamine and 0.1% β mercaptoethanol [Gibco]). Splenocytes were pelleted and resuspended in red blood cell lysis buffer (Sigma) for two minutes at room temperature to remove erythrocytes. Cells were washed and resuspended in complete medium. Spleen and MeLN cells were distributed in 96-well round bottom plates (Sarstedt, Landskrona, Sweden) at a density of 1.5 × 10⁶ cells/ml for antibody staining. Cells were then divided into groups being antibody stained directly (day 0) or after Concanavallina A (ConA) incubation (day 2). Directly stained cells were incubated with a Fc-receptor blocking antibody (FcR; CD16/32; clone 93, Biologic) for 5 minutes followed by incubation with either of two antibody panels. For panel one; CD80-PE (clone 16-10A1), CD11c-PE/Cy7 (N4-18), CD86-PB (GL-1) and I-A/E-A700 (M5-114.15.2; all Biologic, San Diego, CA, USA except CD80-PE; eBioscience, San Diego, CA, USA) and panel two CD44-AF488 (IM7), CD122-PE (5H4), CD3-PE/Cy7 (145-2C11), CD4-PB (GK1.5), CD2L2-APC (MEL-14) and CD8-APC/Cy7 (53-6.7) at 4°C for 30 minutes. 7AAD (Sigma; 1 µl/sample) was included in both panels. Further, cells were washed with FC buffer (0.5% bovine serum albumin (Sigma) and 0.5 mM EDTA in phosphate buffered saline) and incubated with 4% PFA (paraformaldehyde; Sigma) for 20 minutes followed by wash and resuspension in FC buffer. Cells in the day 2-group were first incubated in presence or absence of 0.625 µg/ml ConA (Sigma) and incubated for 2 days at 37°C and 5% CO₂. The antibody staining procedure was identical to the procedure for the day 0-group and both groups were acquired in the same flow cytometry run. Measurements were performed using a Cytometric ADP (Beckman Coulter, Brea, CA, USA), analyses were performed using FlowJo (Treestar inc., Ashland, OR, USA, version 7.6.1) and gating was adjusted using fluorescence minus one (FMO) control staining.

**B. Mice analyzed at 14 weeks of age.** The MeLN were meshed through a cell strainer (70 µm, BD Bioscience). The single cell suspension of the lymph nodes was washed in cell medium (PBS (Gibco), 2% FCS, 2 mM EDTA) and resuspended in RPMI-1640 (Gibco). The spleens were minced into pieces and incubated in spleen dissociation buffer (Stemcell Technologies, Vancouver, Canada) according to manufacturer’s instructions. Cells were resuspended in RPMI-1640 and distributed in 96-well round bottom plates (Sarstedt) at a density of 1 × 10⁶ cells/ml. Thereafter cells were incubated with a Fc-receptor blocking antibody (FcR; CD16/32; clone 93, Biologic) for 5 minutes followed by incubation with CD3-PE/Cy7 (145-2C11), CD4-PB (GK1.5) and CD25-APC (PC61; all Biologic) for 30 minutes at 4°C. The cells were resuspended in Fix/Perm solution (eBioscience), washed with permeabilization buffer (eBioscience) and blocked with FcR for 5 minutes prior to incubation with FoxP3-PE (MF-14, Biologic) for 30 minutes at 4°C. Cells were washed with permeabilization buffer, resuspended in FC buffer (1% fetal calf serum (Gibco) and 0.5 mM EDTA in PBS) and acquired as described for the 26 weeks old mice.

**Proliferation of splenocytes from mice at 14 weeks of age**

Spleen CD11c⁺ cells were isolated using Easy sep CD11c⁺ positive selection kit (Stemcell Technologies) with a purity of 71%. Subsequently CD4⁺ cells were isolated using Easy sep CD4⁺ positive selection kit (Stemcell Technologies) with a purity of 70% followed by CD8⁺ cell isolation using Easy sep mouse CD8⁺ T cell enrichment kit (Stemcell Technologies) with a purity of 89%. CD11c⁺, CD4⁺ and CD8⁺ cells (each 5 × 10⁵/well) were cultured in medium (RPMI-1640 medium containing 10% heat-inactivated fetal calf serum, 1 mMol/L sodium pyruvate, 10 mMol/L Hepes, 50 U penicillin, 50 µg/ml streptomycin, 0.05 mMol/L β-mercaptoethanol and 2 mMol/L L-glutamine, Gibco, Invitrogen) in 96-well round bottom plates (Sarstedt) in presence or absence of 0.625 µg/ml ConA (Sigma) for 72 hours at 37°C and 5% CO₂. To measure cell proliferation, [methyl-³H] thymidine (PerkinElmer, Waltham, MA, USA) was added (20 µl/well diluted 1:20) and incubated 16-20 hours. Macromolecular material was then harvested on glass filter filters using a Printed Filtermat A (1450-421, Wallac Oy, Turku, Finland). Filters were air-dried and the bound radioactivity was measured in a beta-counter (Wallac 1450 MicroBeta, Ramsey, MN, USA).

**Analysis of cytokines in plasma**

Cytokine concentrations in plasma were analyzed using a mouse Th1/Th2 9-Plex (IFN-γ, IL-1β, TNF-α, IL-2, IL-12, IL-4, IL-5, IL-10 and KC/GRO/CINC (CXCL1) Ultra-Sensitive Kit (Meso Scale Discovery, Gaithersburg, MO, USA), following the instructions of the manufacturer. Samples were analyzed separately for each individual animal. The lower detection limit in this assay was 1.02–9.50 pg/ml depending on the cytokine assayed.

**Analysis of cholesterol and triglyceride content**

Total plasma cholesterol and plasma triglycerides were quantified with colorimetric assays, Infinity™ Cholesterol and Triglyceride (Thermo Electron, Waltham, MA, USA).

**Statistical analysis**

Analysis of data was performed using unpaired t-test or Mann Whitney test for skewed data. Data are presented as mean± standard deviation. Kruskal-Wallis test and Dunn’s post hoc test was used to analyze plasma triglyceride data. Analysis was performed using GraphPad Prism 3.01 (Graphpad software, La Jolla, CA, USA) and a level of P<0.05 was considered significant.

**Results**

Characterization of the mouse model and quantification of atherosclerosis

We first quantified CD8⁺ T cells in *ApoE−/−* mice and counterparts deficient in the MHC class I transporter TAP1. As expected the CD8⁺ T cell fraction and numbers were low in...
Figure 1. CD8$^+$ T cells in spleen and mediastinal lymph nodes (MeLN). Flow cytometry graphs showing the cell populations in spleen from one representative mouse from each group (A). Numbers given in the graphs are per cent cells out of all lymphocytes. The fraction (B) and total cell count (C) of CD8$^+$ T cells in spleen and MeLN of Apoe$^{-/-}$, Apoe$^{-/-}$ Tap1$^{-/-}$ and Tap1$^{-/-}$ mice. The cells were isolated from respective tissue, stained with fluorescent antibodies and analyzed by flow cytometry. As expected, the CD8$^+$ T cell fraction was depressed in the mice lacking the Tap1 gene. Each dot in the figure represents one mouse. ***P<0.001.

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Thus, TAP1 deficiency does not induce severe alterations in the intra-
difference in CD3
MeLN was almost doubled (figure 3C), which may explain the
mately eight times larger in MeLN and spleen of
+ Tap1−/− mice did not develop quantifiable lesions and were
excluded from analysis. The same pattern was found in mice given
high fat diet for 8 weeks (table 1). Since TAP1 deficiency could
influence inflammation and lipid accumulation in the lesions we
analyzed the T cell-, macrophage- and lipid content in the old
mice as indicators of lesion stability. No differences were detected
in lesion infiltration of macrophages (20.8±5.8% vs. 21.3±3.8%) or
CD3+ T cells (30.7±10.2% vs. 25.6±3.6%) in Apoe−/− Tap1−/−
compared to Apoe−/− mice. There were no detectable amounts of
CD8+ T cells in lesions of mice given HFD for 22 weeks (data not
shown). Further, no differences were found in CD4+ T cell lesion
infiltration in Apoe−/− Tap1−/− compared to Apoe−/− mice (data
not shown). The neutral lipid accumulation in the lesions was not
affected (11.2±4.5% in Apoe−/− Tap1−/− vs. 13.8±4.3% in Apoe−/−).
Thus, TAP1 deficiency does not induce severe alterations in the intraleisional milieu. Systemically, plasma cholesterol and triglyceride
levels were not different between groups on Apoe−/− background but
as expected levels were increased in these groups compared to mice
on an Apoe−/− background (table 1 and 2). The weights did not differ
between Apoe−/− and Apoe−/− Tap1−/− mice (table 1 and 2).

Cell characterization

The T cell and dendritic cell (DC) populations of spleen and
MeLN were analyzed to characterize the response to hypercholesterolemia systemically and in conjunction to lesions. While
viable CD3+ T cells were less abundant in spleen of Apoe−/−
Tap1−/− mice (figure 3A and 3B), there was no difference in
MeLN compared to Apoe−/− mice (figure 3B). The fraction of
CD3+CD4+ T cells was higher in Apoe−/− Tap1−/− mice and
Tap1−/− mice compared to Apoe−/− mice in both spleen and
MeLN (figure 3A and 3C and data not shown). While the increase
in spleen was moderate, the CD3+CD4+ T cell compartment in
MeLN was almost doubled (figure 3C), which may explain the
difference in CD3+ T cells between the groups in spleen compared
to the indifference between groups in MeLN (figure 3B); however,
since the number of CD4+ T cells did not differ in MeLN
(figure 3D) the relative rise in the CD4+ T cell population is likely a
result of the diminished CD8+ T cell population. Interestingly, the
number of CD4+ T cells in spleen increased in the Apoe−/− Tap1−/−
mice (figure 3D). Since this population could mediate inflammation
we analyzed the fraction of memory effector cells. There was no
difference in CD4+CD44+CD62L+ T cells in Apoe−/− Tap1−/−
mice compared to Apoe−/− mice in MeLN or spleen (figure S1A).
Surprisingly, the corresponding CD8+ population was approximately eight times larger in MeLN and spleen of Apoe−/− Tap1−/−
mice compared to Apoe−/− mice (figure S1B). However, the
CD8+CD44+CD62L+CD122+ regulatory T cell [19] population in
the small CD8+ T cell population was also higher in Apoe−/−
Tap1−/− mice (figure S2), which could compensate rise in effector
cells. In contrast, within the CD4+ population the CD4+CD25FoxP3+
regulatory T cell fraction was decreased in MeLN but not in spleen of Apoe−/− Tap1−/− mice given HFD for 8
weeks (14 weeks old at death) compared to equivalent Apoe−/− mice
(figure S3). Since the major activation pathway of CD8+ T cells
occur via DCs we analyzed abundance and activation of CD11c+
cells in spleen and MeLN of 26 weeks old mice. The fraction of
CD11c+ cells in spleen, but not in MeLN, was lower in Apoe−/−
Tap1−/− mice compared to Apoe−/− mice (figure 4B). Further,
CD11c+ cells in both organs expressed lower levels of CD80, CD86 and a trend towards decreased MHC class II levels (figure 4A, 4C and 4D).

Cell characterization of proliferated lymphoid cells

In order to elucidate the proliferative response of CD4+ and CD8+ T cells, lymphocyte suspensions from spleen and MeLN were incubated with ConA for two days. The fraction of respective cell type was assessed at day 0 and day 2 using flow cytometry analyses. The cell pattern at day 0 is presented in figure 1 and 3. However, incubation with ConA resulted in a robust rise in CD3+CD8+ T cells at day 2 in the Apoe2/2 Tap12/2 mice in both spleen and MeLN (figure S4). Interestingly, the CD3+CD8+ T cell population at day 2 was also increased in Apoe2/2 mice in comparison to CD3+CD4+ T cells (figure S4). This may indicate that the CD3+CD8+ T cells respond strongly to the ConA incubation while the CD3+CD4+ T cells are weak responders.

Proliferation of splenocytes

To test the impact of antigen presenting cells (APCs) on the proliferation of separated CD4+ and CD8+ T cells we performed analysis in mice given high fat diet for 8 weeks (14 weeks old at death). The CD4+ and CD8+ T cells from Apoe2/2 Tap12/2 mice and Apoe−/− mice were magnetically separated and stimulated for three days with ConA and an equal amount of CD11c+ cells. In this setting the CD8+ T cells proliferate poorly in both Apoe2/2 Tap12/2 mice and Apoe−/− mice (figure 5). In contrast, CD4+ T cells proliferate more than the CD8+ T cells in both groups. Interestingly, CD4+ T cells had an increased proliferation in Apoe−/− Tap1−/− mice compared to Apoe−/− mice (figure 5). Stimulation of T cells with ConA is considered to give an estimation of the T cell priming status from the in vivo environment. Thus, cells in the CD4+ co-culture seems to be more pre-primed compared to cells in the CD8+ co-culture and more in TAP deficient animals.

Plasma cytokines

The plasma level of IL-12, IL-1β, KC, IFN-γ, TNF-α, IL-2, IL-10, IL-4 and IL-5 was analyzed but no differences between the groups of mice were found (data not shown).

Discussion

T cell immunity plays an important role in the development of atherosclerosis, but the participation of CD8+ T cells and MHC class I antigen presentation has not been fully investigated. Previous reports on the role of CD8+ T cells in the development of atherosclerosis are few and describe a diverse impact on disease indicating a need for further characterization. We recently evaluated the CD8+ T cell response to hypercholesterolemia in an Apoe−/− mouse model leading to the conclusion that initial atherosclerosis is characterized by a CD8+ T cell response which was more rapid and stronger than the corresponding CD4+ T cell response [13]. The hypothesis of the current study was therefore
that an Apoe<sup>−/−</sup> mouse model with few CD8<sup>+</sup> T cells (Apoe<sup>−/−</sup>Tap1<sup>−/−</sup>) would have a decreased lesion development in response to a high fat diet compared to an Apoe<sup>−/−</sup> mouse model with normal levels of CD8<sup>+</sup> T cells. However, lesion size was found to be equal in the Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice and Apoe<sup>−/−</sup> mice. As lesion size in younger mice display the same pattern, the lesion severity may be of less importance in this system. However, as the diet composition may impact the presence of antigens implicated in autoimmune lymphocyte response, experimental groups given chow diet could have been included in the study. Interestingly, a study in which chow-fed Apoe<sup>−/−</sup> mice with a mutation in the CD8-encoding gene Lyt-2 developed equivalent atherosclerosis as Apoe<sup>−/−</sup> mice with functional CD8<sup>+</sup> T cells [7]. Accordingly, neither loss of function in CD8<sup>+</sup> T cells nor the TAP1-dependent CD8<sup>+</sup> T cell activation is essentially affecting the lesion development. The result raises the question if 1) the CD4<sup>+</sup> T cell population compensates pro-inflammatory effects of the normal CD8<sup>+</sup> T cell population or 2) the CD8<sup>+</sup> T cell population is not involved in the disease process. The CD4<sup>+</sup> fraction of Tap1<sup>−/−</sup> and Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice was enlarged compared to TAP1-expressing mice in contrast to studies in Tap1<sup>−/−</sup> mice [14] and Tap1<sup>−/−</sup> humans [20] having normal CD4<sup>+</sup> levels. Although the effector memory CD4<sup>+</sup> T cell population was equal in both groups, the actual number of cells was larger in Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice due to the enlarged CD4<sup>+</sup> T cell population. Moreover, the CD4<sup>+</sup>CD25<sup>+</sup>FoxP3<sup>+</sup> regulatory T cell population was decreased in MeLN of the group given HFD for 8 weeks supporting a disease-driving property of this T cell subset. Hence, pro-atherogenic cells within this population may maintain disease by compensating for the loss of CD8<sup>+</sup> T cells. Apart from this, diminishment of the CD8<sup>+</sup> T cell population may reduce the impact of both pro- and anti-inflammatory CD8<sup>+</sup> T cells on atherosclerosis. This could be a transient process or affect either of the cell populations more than the other. Interestingly, incubation of lymphocytes with ConA resulted in an increased fraction of CD8<sup>+</sup> T cells while the CD4<sup>+</sup> T cells were weaker responders and displayed a decrease in the fraction of lymphocytes. This supports the data presented in our previous study [13] and results by Aldrich et al. describing a strong antigen-specific response in CD8<sup>+</sup> T cells from Tap1<sup>−/−</sup> mice [21]. However, when splenic CD4<sup>+</sup> or CD8<sup>+</sup> T cells were mixed with CD11c<sup>+</sup> cells and stimulated with the polyclonal T cell activator ConA, there was an increased proliferation in CD4<sup>+</sup> cultures from Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice compared to corresponding CD8<sup>+</sup> cultures. Furthermore, the CD4<sup>+</sup> cultures from Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice proliferated more than corresponding cultures in Apoe<sup>−/−</sup> mice. Since CD8<sup>+</sup> T cells benefit from CD4<sup>+</sup> T cell assistance [22] the proliferation of the isolated CD8<sup>+</sup> T cells may be negatively regulated. In contrast, the CD4<sup>+</sup> T cells in the Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice seem to be more pre-primed than in the Apoe<sup>−/−</sup> mice. Taken together, the data may indicate that CD8<sup>+</sup> T cells need CD4<sup>+</sup> T cell help to proliferate properly and CD4<sup>+</sup> T cell proliferation may be inhibited by CD8<sup>+</sup> T cell proliferation in a co-culture. Since the results of this in vivo testing seem to have little impact on the disease mechanism it may not be reflected in vivo. The in vivo system may promote other functional mechanisms and/or the CD8<sup>+</sup> T cells may not respond to endogenous antigens present in the in vivo system as strong as they respond to ConA. Thus, the CD8<sup>+</sup> T cell population in Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice was only 12% of that in Apoe<sup>−/−</sup> mice but it contained relatively more effector memory- and regulatory T cells. Although CD8<sup>+</sup> T cells in TAP1-deficient mice are known to be functional [23], the effector cell- and regulatory T cell population here was still smaller than in Apoe<sup>−/−</sup> mice, giving them a limited impact on atherosclerosis.

Since about one third of all MHC class I-peptide complexes are presented via the TAP1-independent pathway [24] it was expected that abundance as well as activation of CD11c<sup>+</sup> DCs would be lower in Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice than Apoe<sup>−/−</sup> mice as was demonstrated in the present study. Considering the size of the CD8<sup>+</sup> T cell population the remaining CD11c<sup>+</sup> cell population may be sufficient to increase the CD8<sup>+</sup> effector memory cell population. Consequently, the larger CD4<sup>+</sup> T cell population may not be adequately stimulated to achieve a similar rise in effector memory cells. However, activation of CD4<sup>+</sup> and CD8<sup>+</sup> T cells involve not only APC derived MHc class I/II-TCR contact but also stimulation from CD4<sup>+</sup> T cells. They can acquire MHC class I/II-peptide complexes from APCs and together with co-stimulatory molecules activate CD4<sup>+</sup> or CD8<sup>+</sup> T cells and increase the APC derived stimulation [22]. Thus, the decreased DC activation in Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice may be compensated by antigen presentation mediated by the increased CD4<sup>+</sup> T cell population, potentially activating both pro-inflammatory CD4<sup>+</sup> and CD8<sup>+</sup> T cell subsets. However, activation of CD4<sup>+</sup> and/or CD8<sup>+</sup> T cell subsets targeting antigens present in atherosclerotic lesions would result in increased lesional T cell infiltration. As the

**Figure 4.** CD11c<sup>+</sup> cells and expression of CD80<sup>+</sup>, CD86<sup>+</sup> and MHC class II<sup>+</sup> on CD11c<sup>+</sup> cells in spleen and MeLN. Histograms corresponding to the cell populations in the spleen from one representative mouse from each group (A). Gate boundaries were set by fluorescence minus one controls (solid grey). The fraction of CD11c<sup>+</sup> cells (B) and CD11c/CD80<sup>+</sup>, CD11c/CD86<sup>+</sup> and CD11c/MHCII<sup>+</sup> cells in spleen (C) and MeLN (D) of Apoe<sup>−/−</sup> and Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice. The cells were isolated from respective tissue, stained with fluorescent antibodies and analyzed by flow cytometry. Each dot in the figure represents one mouse. *P<0.05, **P<0.01.

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**Figure 5.** Proliferation of splenic CD4<sup>+</sup> and CD8<sup>+</sup> co-cultures of young mice. Proliferation were analysed in ConA stimulated (90 hrs) CD4<sup>+</sup>/CD11c<sup>+</sup> (CD4<sup>+</sup>) and CD4<sup>+</sup>/CD11c<sup>−</sup> (CD8<sup>+</sup>) cell co-cultures in Apoe<sup>−/−</sup> and Apoe<sup>−/−</sup>Tap1<sup>−/−</sup> mice with presence of radioactive labelled thymidine during the last 16–20 hrs. The cells were isolated from spleens of mice given high fat diet for 8 weeks and magnetically separated into CD11c<sup>+</sup>, CD4<sup>+</sup> and CD8<sup>+</sup> cells. CPM denotes counts per minute. Each dot in the figure represents one mouse. *P<0.05.

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amount of T cell subsets was similar or below detection limit the impact of T cells for lesion development may be questioned in this model. Moreover, the macrophage lesion infiltration was not affected indicating that TAP1 deficiency does not exert major APC-driven effects on atherosclerosis progression. Thus, the similar lesion size in Apoe−/− mice and Apoe−/− Tap1−/− mice reflects a minor role of TAP1 for atherosclerosis development. However, the mice of this study were housed in an environment with low exposure to infections. Several pathogens, including Chlamydia pneumoniae, herpes simplex virus and cytomegalovirus have been associated with atherosclerosis [25]. As CD8+ T cells are activated by bacterial/viral antigens via TAP dependent- and independent pathways [16] it is conceivable that the model of this study contains limitations. Thus, assuming a low infectious burden in the mice of the present study, a reduced amount of autoreactive CD8+ T cells targeting lesion specific antigens may favor other atherogenic cell types.

In conclusion, hypercholesterolemic Apoe−/− mice that lack TAP1-dependent antigen presentation develop atherosclerotic lesions, inflammatory cell infiltration and lipid accumulation equal to Apoe−/− mice. The relative CD4+ T cell population was larger in both local and systemic lymphoid tissues in Apoe−/− Tap1−/− mice compared to Apoe−/− mice, whereas CD4+ T cell numbers were increased in spleen but not in MeLN. In addition, the CD11c+ population had a lower expression of activation markers in Apoe−/− Tap1−/− mice. Although the effector memory T cell population was larger in the small CD8+ T cell population of Apoe−/− Tap1−/− mice the impact on disease is likely to be limited.

Taken together, the present study provides novel information indicating a limited role of TAP1 and CD8+ T cells in atherosclerosis development.

Supporting Information

Figure S1 Effector memory CD4+ T cells and CD8+ T cells in spleen and MeLN. The fraction of (A) CD3+CD4+CD62L− and (B) CD3+CD8+ CD44+CD62L− T cells in spleen and MeLN of Apoe−/− and Apoe−/− Tap1−/− mice. The cells were isolated from respective tissue, stained with fluorescent antibodies and analyzed by flow cytometry. Each dot in the figure represents one mouse. ***P<0.001.

Figure S2 Regulatory CD8+ T cells in spleen and MeLN. Analysis of CD3+CD8+CD44−CD62L+ CD122+ T cells in Apoe−/− and Apoe−/− Tap1−/− mice in spleen and MeLN. The cells were isolated from respective tissue, stained with fluorescent antibodies and analyzed by flow cytometry. Each dot in the figure represents one mouse. **P<0.01, ***P<0.001.

Figure S3 Regulatory CD4+ T cells in spleen and MeLN of young mice. Analysis of CD3+ CD4+CD25+FoxP3+ T cells in spleen and MeLN of Apoe−/− and Apoe−/− Tap1−/− mice given HFD for 8 weeks. The cells were isolated from respective tissue, stained with fluorescent antibodies and analyzed by flow cytometry. Each dot in the figure represents one mouse. **P<0.01.

Figure S4 ConA-stimulation of lymphocytes. Lymphocytes from spleen and MeLN were incubated with ConA for 2 days and the relative fraction of CD4+ and CD8+ T cells was assessed by flow cytometry analysis. The CD8+ T cell fraction was larger than the CD4+ T cell fraction in spleen and MeLN of both Apoe−/− and Apoe−/− Tap1−/− mice. **P<0.01, ***P<0.001.

Author Contributions

Conceived and designed the experiments: DK HB JN GNF. Performed the experiments: DK IL IS RA. Analyzed the data: DK. Wrote the paper: DK HB JN GNF.

References


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