Immune Modulatory Effects of Hypercholesterolemia: Can Atorvastatin Convert the Detrimental Effect of Hypercholesterolemia on the Immune System?

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Abstract

Background and purpose: Many observations showed that hypercholesterolemia can disrupt immune response. Statin drugs that were used for the treatment of hypercholesterolemia patients can interfere in regulation of the immune response and cytokine secretion. The primary aim of the current study was to investigate the immune response among hypercholesterolemia patients, who were treatment-naïve and healthy subjects. The secondary goal of the study was to determine whether atorvastatin can reverse the detrimental effect of hypercholesterolemia on the immune system.

Methods: Peripheral blood mononuclear cells (PBMCs) were isolated from 50 patients afflicted with hypercholesterolemia who were treatment-naïve along with 50 sex- and age-matched hypercholesterolemia patients receiving atorvastatin and 50 sex- and age-matched healthy subjects. Quantitative PCR and ELISA methods were used for gene and protein expression analysis of T helper 1 (Th1) and Th2 related cytokines. Additionally, the expression of the cluster of differentiation (CD) markers on T, B, and NK cells were measured by flow cytometry method.

Results: The results showed that hypercholesterolemia and atorvastatin downregulated the expression of Th1-related cytokines and elevated the levels of Th2-related cytokines. The expression of cell surface markers, CD25 and CD69, was significantly decreased in the treatment-naïve, and atorvastatin groups.

Conclusion: It seems that atorvastatin is not able to repair the deleterious effects of hypercholesterolemia on the immune system, and elevated levels of cholesterol along with the administration of atorvastatin tilt the Th1/Th2 balance in favor of Th2 and reduce T cell activation.

Keywords: Hypercholesterolemia; Immune response; Atorvastatin; Th1/Th2 balance; T cell activation.

Introduction

Hypercholesterolemia is principally defined by high fasting cholesterol that stems from the disturbance in lipid metabolism and genetic defects. It increases the risk of atherosclerosis, cardiovascular diseases, angina, and stroke [1]. Of note, hypercholesterolemia impedes the immune response and cytokine secretion in immune cells. These effects of hypercholesterolemia were due to alteration of cholesterol rates in the cell membrane and cytoplasm of immune cells [2]. Studies indicated that ApoE-/- mice had increased levels of Th2 cells [3] and the decreased number of Th1 cells showing a deficiency in cell-mediated immunity [4-6]. However, the research did not come to a consensus about the association between hypercholesterolemia and the risk of developing cancer [7,8].

Statin, 3-hydroxy-3-methylglutaryl-CoA (HMG-CoA) reductase inhibitors, are a class of compounds used for the treatment of patients with hypercholesterolemia, hyperlipidemia, and cardiovascular diseases [9]. Despite the beneficial effect of atorvastatin on cholesterol lowering, the drug has several side effects including Myalgia [10], rhabdomyolysis, and myopathy [11]. Statins also influence the immune system as they are able to halt T cell homing, migration, and adhesion, the latter is mediated by the downregulation of integrins, adhesins, and matrix metalloprotease proteins (MMPs) [12,13]. It is thought the modulation of T cell response is exerted by the hindrance of antigen presenting cells (APCs) development [14,15] through a reduction in major histocompatibility complex II (MHC II) leaving the T cells inactive [16]. As a result, statins predominantly temper the activity of immune cells monocyte/macrophages and T cells [17]. The ability to avert the activation of Th1 cells upon using statins made them desirable candidates for the alleviation of autoimmune diseases such as rheumatoid arthritis, multiple sclerosis, systemic lupus erythematosus (SLE) [18-20] as well as autoimmune myocarditis [21]. In contrast to animal models, studies performed on human clinical trials showed that atorvastatin did not change any shift in the balance of Th1/Th2 cells [22]. It has been demonstrated that blocking the production of isoprenoids required for prenylation of proteins such as Rho, Rac and Ras superfamily GTPases plays a crucial role in fidelity of statins in amelioration of inflammation [23,24].
There is a few information about the effect of hypercholesterolemia on the immune system in the literature [26-28]. All of the researches dealt with the human studies in terms of statin interaction with the T cell functions were exclusively confined to cultured PBMCs taken from the healthy individuals, patients with autoimmune and cardiovascular diseases [26-28]. Regarding the scarcity of human studies on the role of hypercholesterolemia in T cell activation and differentiation, we aimed to investigate whether:

- hypercholesterolemia can affect the immune system function;
- atorvastatin can oppress the harmful impact of hypercholesterolemia on the immune system.

**Methods and Materials**

**Study population**

In this study, a total of one-hundred fifty individuals participated that were divided into three groups as the following categorization: 50 patients afflicted with hypercholesterolemia who received no medication for cholesterol lowering (treatment-naive group), 50 patients who were sex- and age-matched taking atorvastatin at a dose of 10 mg/kg for at least 4 months (atorvastatin group), and 50 healthy individuals who were sex- and age-matched either (healthy subjects). The participated subjects were chosen from those referring to Hazrat-e-Rasoul Hospital, Tehran, Iran. The diagnosis of hypercholesterolemia was made in accordance with the measurements of cholesterol at the particular time interval confirming that the patients suffer from persistent hypercholesterolemia (Table 1). The exclusion criteria for the volunteers were smoking, neurological disorder, cardiovascular diseases, inflammatory and allergen disorders, and diabetes. The protocol used for the sample collection and patient examinations was confirmed by the ethical committee of the National Institute of Genetic Engineering and Biotechnology (NIGEB), and each volunteer was given informed consent. All methods were performed in accordance to Helsinki Declaration.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Healthy (n=50)</th>
<th>Treatment-naive (n=50)</th>
<th>Atorvastatin (n=50)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex (male/female)</td>
<td>23/27</td>
<td>25/25</td>
<td>26/24</td>
<td>0.9134</td>
</tr>
<tr>
<td>Mean age (range), years</td>
<td>47 (41-55)</td>
<td>49 (44-60)</td>
<td>53 (49-62)</td>
<td>0.3432</td>
</tr>
<tr>
<td>Mean BMI (range), kg/m²</td>
<td>23.2 (20-24)</td>
<td>32.8 (29-35)</td>
<td>28.2 (26-31)</td>
<td>0.0321</td>
</tr>
<tr>
<td>Mean cholesterol level ± SD (mg/dl)</td>
<td>160.2 ± 11</td>
<td>300.4 ± 15</td>
<td>172 ± 19.8</td>
<td>0.0061*</td>
</tr>
<tr>
<td>Mean LDL level ± SD (mg/dl)</td>
<td>102.2 ± 11.8</td>
<td>158.8 ± 14.5</td>
<td>105 ± 15.6</td>
<td>0.0176*</td>
</tr>
<tr>
<td>Mean HDL level ± SD (mg/dl)</td>
<td>40.2 ± 4.3</td>
<td>42.5 ± 10.3</td>
<td>39.5 ± 5.7</td>
<td>0.8971</td>
</tr>
<tr>
<td>Mean Triglyceride level ± SD (mg/dl)</td>
<td>129.8 ± 15.8</td>
<td>124.1 ± 30</td>
<td>122.6 ± 24.3</td>
<td>0.7692</td>
</tr>
</tbody>
</table>

BMI: Body Mass Index; LDL: Low-Density Lipoprotein; HDL: High-Density Lipoproteins

**Table 1: Demographic and clinical characteristics of samples in this study.**

**Cell isolation and serum separation**

Whole blood samples (5 ml) were collected from all individuals and poured in 0.5 M EDTA-containing tubes to avoid coagulation. The peripheral blood mononuclear cells (PBMCs) were separated based on the gradient density centrifugation technique by Ficoll hypaque (Pharmacia, Uppsala, Sweden). For the serum separation, blood samples were centrifuged and stored at -80°C for until use.

**RNA extraction and cDNA synthesis**

Total RNA was extracted from PBMCs using a High Pure RNA Isolation Kit (Roche, Germany) according to the manufacturer’s instructions. The quantity and quality of RNA were determined using NanoDrop 2000 (Wilmington, USA) at 260 nm and agarose gel analysis. For reverse transcription polymerase chain reaction (RT-PCR), 400 ng of total RNA was converted into complementary DNA (cDNA) using Revert Aid First Strand cDNA Synthesis Kit (Thermo Fisher Scientific, Germany) based on manufacturer’s protocols. The produced cDNA was stored at -20°C for spare use.

**PCR and Real-Time PCR analysis**

Primers designing were performed by oligo7 software (version 7.56) using the RefSeq sequences of GenBank database as described in Table 2, the specificity of each primer pair was confirmed by Primer- BLAST online software. A PCR was performed to check for any contaminating DNA and in order to confirm the transcription of interested genes in a final volume of 20 μl with master mix PCR (Cinnagen, Tehran, Iran), gene- specific primers (10 pmol/μl), and cDNA (1 μg) in all samples and then loaded them on gel 1.5%.

Real-time-PCR was performed for relative quantification of the mRNA expression of all interested genes in a final volume of 10 μl, using the SYBR Green Real-time PCR Master Mix (Light Cycler Fast Start DNA Master Plus SYBR Green I, Roche, Germany), and 10 pmol of the previously mentioned primers and 1 μg cDNA. The real time PCR protocol was as follows: pre-denaturation at 95°C for 600 seconds, denaturation at 95°C for 15 seconds, annealing at 60°C for 20 seconds, extension at 72°C for 20 seconds, with 40 cycles. Identification of fluorescent intensity in each sample was performed by Rotor gene (Termocicler Rotor-Gene 6000 Corbett Research/ Australia). Melting curve analysis showed only one peak for each reaction and this was also confirmed by electrophoresis of PCR products that showed only one band of the expected size.
Cytokine measurements

The concentration of IFN-γ, TNF-α and IL-2 cytokines was assayed in the serum of three studied groups. In this section, a total of 60 individuals were recruited that consisted of 20 subjects (11 women+9 men; mean age=47 years) in the treatment-naïve group, 20 (10 women, 10 men; average age=49 years) patients treated with atorvastatin, and 20 (12 women+8 men; mean age=48 years) healthy individuals. The aforementioned factors were all analyzed by the commercial kits (R&D Systems, Minneapolis, Minnesota, USA) at 450 nm.

Immunophenotyping

Cell surface staining was performed for a total of thirty individual consisted of 10 subjects (5 women+5 men; mean age=50 years) in the treatment-naïve group, 10 (6 women+4 men; mean age=48 years) patients in the atorvastatin group, and 10 (5 women+5 men; mean age=45 years) healthy subjects. Immunophenotyping was done using antibodies against CD3 (FITC, Dako Denmark), CD4/CD8 (FITC/PE, BD Biosciences), CD25 (PerCP/Cy5.5, Biolegend), CD69 (PerCP/Cy5.5, Biolegend), CD22 (FITC, Dako, Denmark), CD16 (FITC, Immunostep), CD56 (PE, Dako Denmark). Appropriate isotype controls were used as negative controls to determine the nonspecific staining and background. In the first stage, 100 μl of whole blood was poured into the test tube, and the antibodies were added to the recommended dilutions. Then, the tubes were incubated for 30 minutes at 4°C. After the incubation time, 1 ml of RBC lysis solution was added and incubated for 10-15 minutes at room temperature. In the last step, the sample was run in BD FACS Calibur flow cytometry (BD Biosciences, USA). The obtained data were analyzed by the Flowjo Software (version 7.6.1). Gating was applied to analyze the lymphocytes, and the results were reported as the percentage of CD4- and CD8-positive lymphocytes expressing either CD69 or CD25 surface markers in a total of all T cell population, CD3- and CD22-positive lymphocytes expressing CD45RO marker, and finally, CD16-positive lymphocytes that expressed CD56 surface marker.

Table 2: Primers used in Real-Time PCR for cytokine, transcription, and internal control genes.

<table>
<thead>
<tr>
<th>Genes</th>
<th>Genbank Number</th>
<th>Accession</th>
<th>Forward primer (5’→3’)</th>
<th>Reverse primer (5’→3’)</th>
<th>Amplicon size (base pair)</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-2</td>
<td>NM_000586</td>
<td>ATTACAGAATCCCAAATCTCACAG</td>
<td>GTTGTTTAGATCTCCTTGTTC</td>
<td>217</td>
<td></td>
</tr>
<tr>
<td>IL-4</td>
<td>NM_172348</td>
<td>CTTCCACCTGTCCTTCTCTTG</td>
<td>CTTCTCAATGTGTTTCTTGCTC</td>
<td>130</td>
<td></td>
</tr>
<tr>
<td>IL-5</td>
<td>NM_000879</td>
<td>TGTGATACCAATGAGACTCTTG</td>
<td>GACCTCCAGTGTCCTATTTC</td>
<td>102</td>
<td></td>
</tr>
<tr>
<td>IL-6</td>
<td>NM_001318095</td>
<td>TCTGATTCTAATGGAGAGACCTTCG</td>
<td>CTCACTACTCTCAAATCCTTGCTG</td>
<td>102</td>
<td></td>
</tr>
<tr>
<td>IFN-γ</td>
<td>NM_000619</td>
<td>CGGTAACGTACCTGAATGTCCAAAC</td>
<td>ACCATTACCTGGATGCTTGCAC</td>
<td>141</td>
<td></td>
</tr>
<tr>
<td>TNF-α</td>
<td>NM_000594</td>
<td>CTCCTCTACACACCTCTAACCC</td>
<td>CGAAGTGCTGGTCTTGTTGC</td>
<td>101</td>
<td></td>
</tr>
<tr>
<td>T-bet</td>
<td>NM_013351</td>
<td>CCAAGTTTTAATCAGCAGACAGACAG</td>
<td>CAAGACACGCTACCAACACATC</td>
<td>131</td>
<td></td>
</tr>
<tr>
<td>GATA-3</td>
<td>NM_001002295</td>
<td>CCCTATTAGGCAAGCAGCAGG</td>
<td>CAGGGCTGTTAATATGTTGAG</td>
<td>167</td>
<td></td>
</tr>
<tr>
<td>β-actin</td>
<td>NM_001101</td>
<td>AGACGCGAGGATGGCAGG</td>
<td>GAGACCTCTACAACCCAGCC</td>
<td>161</td>
<td></td>
</tr>
</tbody>
</table>

Statistical analysis

For analyzing the real time PCR data, the PCR efficiency (E) and crossing point deviation (ΔCP) was calculated by LinRegPCR Software (Version 11.0), following the use of Relative Expression Software Tool (REST © 2009, Version 2.0.13) for statistical analysis. Normality of other data was determined using the Kolmogorov–Smirnov test. With respect to the data (parametric vs. non-parametric) analysis of variance (ANOVA) and Kruskal-Wallis tests were applied where appropriate followed by post hoc Tukey’s test. The data were expressed as the mean ± standard deviation (SD). The calculation of the comparison was carried out by the SPSS software (version 16; SPSS, Chicago, ILL). The difference between the variables was statistically significant if the p-value was less than 0.05.

Results

Hypercholesterolemia and atorvastatin decreased the mRNA expression of Th1-related cytokines

The results showed that the treatment-naïve and atorvastatin groups had significantly (p<0.0001) lower the mRNA expression of IL-2, IL-6, IFN-γ, TNF-α cytokines, and T-bet transcription factor in comparison to the healthy individuals. In addition, the mRNA expression of IFN-γ, TNF-α cytokines and T-bet was lower in the atorvastatin group compared with the treatment-naïve group (p<0.01); however, the expression of IL-2 and IL-6 remained unchanged (Figure 1).

Hypercholesterolemia and atorvastatin increased the mRNA expression of Th2-related cytokines

The treatment-naïve group had significantly (p<0.05) higher levels of cholesterol and LDL, as well as the higher expression of IL-5, and GATA-3 in comparison to the healthy subjects; however, the expression of IL-4 did not significant differ between the two groups. Correspondingly, the treatment with atorvastatin caused a significant upregulation in the mRNA expression of IL-5 and GATA-3. The expression of IL-4 was not affected by the administration of atorvastatin in patients with hypercholesterolemia. On one hand, the expression of GATA-3 was significantly (p<0.0001) increased (2 fold)
in atorvastatin group when compared with the treatment-naïve group; yet, the expression of Th2-related cytokines namely IL-4 and IL-5 did not significant differ between two groups (Figure 2).

![Figure 1](image1.png)

**Figure 1:** The bar plot indicates the mRNA expression of Th1-related genes and T-bet transcription factor in PBMC samples. The expression ratios plotted for IL-2, IL-6, IFN-γ, TNF-α, and T-bet genes were significantly decreased in the treatment-naïve and the atorvastatin group patients in comparison to the healthy individuals. The mRNA expression of IFN-γ, TNF-α cytokines, and T-bet was lower significantly in the atorvastatin group compared with the treatment-naïve group. Error bars represent the standard deviation. Number of subjects in each of groups= 50. Statistical significance was designated as: *, p<0.01; ***, p<0.0001.

![Figure 2](image2.png)

**Figure 2:** The bar plot indicates the mRNA relative expression patterns of Th2-related genes and GATA-3 transcription factor in PBMC samples. The treatment-naïve and atorvastatin groups had significantly higher the mRNA expression of IL-5 and GATA-3 genes in comparison to the healthy individuals. The mRNA expression of GATA-3 was increased about 2 folds in atorvastatin group patients in comparison to the healthy individuals. Individuals treated with atorvastatin showed significantly (p<0.01) lower levels of TNF-α than those who were treatment-naïve. On one hand, hypercholesterolemia did not significant effect in the expression of Th2 related cytokines in the treatment-naïve group in comparison to the healthy subjects. Correspondingly, the treatment with atorvastatin caused a significant increase in the expression of IL-5 in comparison to the healthy subjects while the expression of IL-4 was not affected by the administration of atorvastatin in patients with hypercholesterolemia (Figure 3).

![Figure 3](image3.png)

**Figure 3:** The bar plot shows the serum level of IL-2, IFN-γ, TNF-α, IL-4 and IL-5 in all groups. The expression ratios plotted for Th1-related proteins were found to be significantly decreased in treatment-naïve and atorvastatin group patients in comparison to healthy subject. The level of TNF-α was lower in atorvastatin group compared to the treatment-naïve group. The expression ratios plotted for Th2-related proteins did not differ between three groups except IL-5 were found to be significantly increased in atorvastatin group patients in comparison to healthy subject. Error bars represent standard deviation. Number of subjects in each of groups= 20. Statistical significance was designated as: *, p<0.01; ***, p<0.0001.

Correlations of Th1- and Th2-related cytokines protein expression with patient's characteristics

Table 3 shows that no significant correlation between the age, and the expression of all the IL-2, IFN-γ, TNF-α, IL-4 and IL-5 cytokines. A significant negative correlation was found between the level of IL-2 and the BMI of all individuals (r = -0.684; p<0.05). The data also indicated that LDL concentration was significantly negative correlated with the expression of IL-2, IFN-γ, and TNF-α (r = -0.73, r = -0.878, and r = -0.927; p<0.05, respectively), while there are no significant correlation with the expression of IL-4 and IL-5. Accordingly, while there are significant correlations between the levels of IL-2, TNF-α and IL-5 with cholesterol level (r = -0.896, r = -0.724, r = + 0.649; p<0.05, respectively), no significant correlation was found between the cholesterol levels and the expression of IFN-γ and IL-4.

Hypercholesterolemia and atorvastatin decreased expression of T cell activation markers

Flow cytometry analysis of the surface receptors of the T cells showed that the treatment-naïve group had a significant lower (p<0.05) number of CD4+, CD25+, CD4+ CD69+, CD8+, CD25+, and CD8+ CD69+ cells over that of the healthy subjects. Similar results were obtained in patients received atorvastatin in comparison to the

Effect of hypercholesterolemia and atorvastatin on protein levels of Th1- and Th2-related cytokines

The results showed that hypercholesterolemia leads to a significant decrease (p<0.0001) in the expression of Th1 related cytokines including IL-2, IFN-γ, and TNF-α in the treatment-naïve group in comparison to the healthy subjects. In line with this, it was shown that atorvastatin significantly decreased the gene expression of IL-2, IFN-γ, and TNF-α in those patients underwent statin therapy in comparison to the healthy individuals. Individuals treated with atorvastatin had significantly (p<0.01) lower levels of TNF-α than those who were treatment-naïve. On one hand, hypercholesterolemia did not significant effect in the expression of Th2 related cytokines in the treatment-naïve group in comparison to the healthy subjects. Correspondingly, the treatment with atorvastatin caused a significant increase in the expression of IL-5 in comparison to the healthy subjects while the expression of IL-4 was not affected by the administration of atorvastatin in patients with hypercholesterolemia.
healthy individuals except for the number of CD4+ CD25+ cells that were not significantly affected in response to atorvastatin treatment. The number of all T cell groups did not significantly differ between the treatment-naïve and atorvastatin groups (Figure 4).

**Table 3: Association of Th1- and Th2- related cytokines with patient’s characteristics in the treatment-naïve group.**

<table>
<thead>
<tr>
<th>Proteins</th>
<th>Age (years old)</th>
<th>BMI (kg/m²)</th>
<th>Cholesterol (mg/dl)</th>
<th>LDL (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-2</td>
<td>r= + 0.175</td>
<td>p= 0.287</td>
<td>r= -0.684</td>
<td>r= -0.896</td>
</tr>
<tr>
<td></td>
<td>p= 0.017</td>
<td>p= 0.007</td>
<td>p= 0.030</td>
<td></td>
</tr>
<tr>
<td>IFN-γ</td>
<td>r= + 0.214</td>
<td>p= 0.886</td>
<td>r= -0.196</td>
<td>r= -0.215</td>
</tr>
<tr>
<td></td>
<td>p= 0.392</td>
<td>p= 0.171</td>
<td>p= 0.029</td>
<td></td>
</tr>
<tr>
<td>TNF-α</td>
<td>r= + 0.118</td>
<td>p= 0.728</td>
<td>r= -0.216</td>
<td>r= -0.724</td>
</tr>
<tr>
<td></td>
<td>p= 0.534</td>
<td>p= 0.043</td>
<td>p= 0.004</td>
<td></td>
</tr>
<tr>
<td>IL-4</td>
<td>r= + 0.253</td>
<td>p= 0.457</td>
<td>r= + 0.186</td>
<td>r= + 0.189</td>
</tr>
<tr>
<td></td>
<td>p= 0.856</td>
<td>p= 0.256</td>
<td>p= 0.321</td>
<td></td>
</tr>
<tr>
<td>IL-5</td>
<td>r= + 0.196</td>
<td>p= 0.792</td>
<td>r= + 0.227</td>
<td>r= + 0.649</td>
</tr>
<tr>
<td></td>
<td>p= 0.537</td>
<td>p= 0.041</td>
<td>p= 0.085</td>
<td></td>
</tr>
</tbody>
</table>

BMI: Body Mass Index; LDL: Low Density Lipoprotein; *, p<0.05; **, p<0.01; ***, p<0.001, p<0.0001

**Effect of hypercholesterolemia and atorvastatin on the phenotype of T, B memory, and NK cells as well as the CD4/CD8 ratio**

It was shown that 1.9% of B cells in the treatment-naïve group were positive for CD22 and CD45RO that was significantly (p<0.0001) lower than the number of CD22+ CD45RO+ cells in the healthy subjects. However, the number of CD3+ T cells and CD56+ CD16+ NK cells remained unchanged. The results also revealed that the treatment with atorvastatin caused a significant (p<0.001) diminish in the number of cells positive for CD22 and CD45RO compared with the healthy individuals; yet, the number of CD3+ T cells and CD56+ CD16+ NK cells did not significantly differ between the two groups. The ratio of CD4 to CD8 in the treatment-naïve patients and patients received atorvastatin was 3.23% and 3.27%, respectively that shows both hypercholesterolemia and receiving atorvastatin can lead to the decrease (p<0.05) in the ratio of those CD markers in comparison to the healthy individuals. The phenotype of T, B memory, and NK cells as well as the CD4/CD8 ratio did not significantly differ between the treatment-naïve and atorvastatin groups (Figure 5).

The comparison between the number of CD69+ and CD25+ T cells showed that the number of CD69+ T cells was significantly elevated in the healthy group compared with the other groups. Besides, when the number of CD4+ CD25+, CD8+ CD25+, CD4+ CD69+, and CD8+ CD69+ was analyzed in all groups, the percentage of cells expressing CD8 and CD25 was more affected in the treatment-naïve and atorvastatin group compared with the healthy control in which a significant decrease in the number of those cells was reported in the treatment-naïve and the atorvastatin groups in comparison to the healthy subjects (Figure 6).

**Correlation of the CD markers expression with patient characteristics**

In Table 4, there was no correlation between the age and BMI of individuals and the expressions of all CD markers. There was a significant negative correlation between the cholesterol levels and the number of CD4+ CD25+, CD8+ CD25+, and CD8+ CD69+ cells (r= -0.956, r= -0.726 and r= -0.914; p<0.05, respectively). Notably, the ratio of CD4 to CD8 was significantly linked to the concentrations of cholesterol in patients who were treatment-naïve. Furthermore, it was shown that the levels of LDL were negatively correlated with the number of CD4 cells positive for CD25 and CD69 (r= -0.868 and r= -0.674; p<0.05, respectively).

**Discussion**

Hypercholesterolemia is one of the major problems of the developed countries caused by the interaction of the environmental factors and genetic predisposition [29]. Statins are a potent class of chemical compounds that have been shown to inhibit the synthesis of cholesterol and thereby lower the levels of circulating free cholesterol in the blood [30]. Many efforts have been made to illuminate the interaction of hypercholesterolemia and statins on the immune system function. However, those studies were exclusively confined to in vitro
research performed on the cultured PBMCs from healthy individuals and patients who were afflicted with autoimmune and cardiovascular diseases [6,31,32]. To get a better insight on the anti-inflammatory effect of hypercholesterolemia and statins on the functionality of the immune system, we investigated the effect of hypercholesterolemia and atorvastatin on T cell differentiation, Th1/Th2 polarization, and T cell activation in patients who were considered either treatment-naïve or atorvastatin users. We also aim to specify whether atorvastatin could avert the Th2-biased humoral immunity in patients with hypercholesterolemia by lowering cholesterol level in these patients.

In this study, we observed that hypercholesterolemia and atorvastatin shifted the ratio of Th1/Th2 in favour of Th2 resulted in mitigation of the cell-mediated immunity and T cell activity. On one hand, it was shown that hypercholesterolemia and atorvastatin incited the humoral immunity system. Hence, atorvastatin could not reverse the harmful effects of hypercholesterolemia on the immune system. Where the atorvastatin is a more safety and well-tolerability amongst statins family, we chose it for the present experiment [33].

In parallel, in vitro studies have displayed that statin drugs, which used for treatment of hypercholesterolemia patients, impair the proper function of the immune cells and consequently weaken their timely response to infectious agents [41]. The inhibitory effects of statins on Th1 cells caused simvastatin is used for the treatment of various inflammatory diseases such as arthritis in rodent models [26]. In contrast to animal experiments [33], oral administration of simvastatin had no significant effect on T helper polarization [28]. According to our results, it is suggested that atorvastatin decrease the mRNA expression of IFN-γ, TNF-α, IL-2 and IL-6 and T-bet transcription factor. In addition, it increases the expression of IL-5 and GATA3 transcription factor in patients with hypercholesterolemia. One of the possible underlying mechanism in which atorvastatin can drive the Th1/Th2 balance in favour of the overproduction of Th2 is the induction of STAT6 phosphorylation and up-regulation of GATA3 transcription factor [42]. It can accordingly inhibit the differentiation of Th1 cells via the preventing STAT4 phosphorylation and down-regulation of T-bet transcription factor [43]. In this context, atorvastatin can downregulate expression of IFN-γ through disruption of MHC II transcription activation and finally, resulting in the inhibition of antigen presentation on APCs and damping the generation of proinflammatory Th1 cells [44]. We measured the protein levels of Th1- and Th2-related cytokines by the ELISA method in which hypercholesterolemia and atorvastatin both lowered the concentrations of Th1-related cytokines such as IL-2, IFN-γ, and TNF-α in serum and concentrations of IL-5 as Th2-related cytokine increased by atorvastatin. The obtained data from the protein
levels corroborated the mRNA expressions of their cognate genes. It should be noted that the correlation of Th1- and Th2-related cytokines was calculated with the demographic characteristics of patients who were treatment-naïve in which the expression of IL-2, TNF-α and IL-5 was associated with the concentration of cholesterol in the serum of hypercholesterolemia patients. In a suitable manner, the level of LDL was also attributed to the levels of IL-2, IFN-γ and TNF-α in the above patients.

<table>
<thead>
<tr>
<th>CD markers</th>
<th>Age (years old)</th>
<th>BMI (kg/m²)</th>
<th>Cholesterol (mg/dl)</th>
<th>LDL (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4CD25</td>
<td>r= + 0.184</td>
<td>r= - 0.134</td>
<td>r= - 0.956</td>
<td>r= - 0.868</td>
</tr>
<tr>
<td></td>
<td>p= 0.784</td>
<td>p= 0.763</td>
<td>p= 0.002*</td>
<td>p= 0.009*</td>
</tr>
<tr>
<td>CD4CD69</td>
<td>r= + 0.213</td>
<td>r= - 0.214</td>
<td>r= - 0.216</td>
<td>r= - 0.674</td>
</tr>
<tr>
<td></td>
<td>p= 0.897</td>
<td>p= 0.975</td>
<td>p= 0.347</td>
<td>p= 0.023*</td>
</tr>
<tr>
<td>CD8CD25</td>
<td>r= + 0.182</td>
<td>r= - 0.189</td>
<td>r= - 0.726</td>
<td>r= - 0.274</td>
</tr>
<tr>
<td></td>
<td>p= 0.533</td>
<td>p= 0.543</td>
<td>p= 0.044*</td>
<td>p= 0.265</td>
</tr>
<tr>
<td>CD8CD69</td>
<td>r= + 0.138</td>
<td>r= - 0.243</td>
<td>r= - 0.914</td>
<td>r= - 0.194</td>
</tr>
<tr>
<td></td>
<td>p= 0.461</td>
<td>p= 0.586</td>
<td>p= 0.003*</td>
<td>p= 0.785</td>
</tr>
<tr>
<td>CD3CD45</td>
<td>r= + 0.245</td>
<td>r= - 0.194</td>
<td>r= - 0.214</td>
<td>r= - 0.169</td>
</tr>
<tr>
<td></td>
<td>p= 0.364</td>
<td>p= 0.456</td>
<td>p= 0.677</td>
<td>p= 0.885</td>
</tr>
<tr>
<td>CD3CD22</td>
<td>r= + 0.264</td>
<td>r= - 0.149</td>
<td>r= - 0.271</td>
<td>r= - 0.253</td>
</tr>
<tr>
<td></td>
<td>p= 0.802</td>
<td>p= 0.939</td>
<td>p= 0.969</td>
<td>p= 0.353</td>
</tr>
<tr>
<td>CD16CD56</td>
<td>r= + 0.153</td>
<td>r= - 0.215</td>
<td>r= - 0.213</td>
<td>r= - 0.299</td>
</tr>
<tr>
<td></td>
<td>p= 0.602</td>
<td>p= 0.935</td>
<td>p= 0.478</td>
<td>p= 0.863</td>
</tr>
<tr>
<td>CD4CD8</td>
<td>r= + 0.231</td>
<td>r= - 0.167</td>
<td>r= - 0.698</td>
<td>r= - 0.267</td>
</tr>
<tr>
<td></td>
<td>p= 0.593</td>
<td>p= 0.300</td>
<td>p= 0.041*</td>
<td>p= 0.901</td>
</tr>
</tbody>
</table>

Table 4: Correlations of CD markers expression in untreated hypercholesterolemia patients with patient characteristics.

Leung et al. and Weber et al. found that atorvastatin can boost the expression of Th2 cells [26,45]. Link et al. showed that rosuvastatin resulted in a rapid reduction in plasma cytokines such as IFN-γ, IL-6, and TNF-α [46]. Similar to our results, Shimada et al. showed that atorvastatin shifted the balance of Th1/Th2 in favor of Th2 in patients with the acute coronary syndrome (ACS) [27]. Prevention of protein prenylation is one of the bona fide strategies that atorvastatin employs to dampen the signal transduction of pathways involved in inflammation [33,47,48]. IL-2 is required for the initiation and activation of T-cell-mediated immunity playing an important role in combating the cancer cells [49]. Along with IL-2, interferon gamma (IFN-γ) is able to eliminate tumor tissue by means of elevating Th1 cells and T cell activation [50,51]. IFN-γ give rises to the upregulation of MHC II in APCs driving T cell to secret cytolytic compounds to kill the tumor cells [52]. TNF-α has been shown a potent tumor killer agent exerting anti-cancer activity both in vivo and in vitro [53]. The TNF-α is involved in the extrinsic pathway of apoptosis and necrosis in damaged and tumor cells [54]. The IL-6 as the Th1-associated cytokines play a key role in cell-mediated immunity thereby stimulation, proliferation, and differentiation of cytotoxic T cells in the presence of IL-2 [55,56]. IL-5 is also an important interleukin which is under the control of GATA-3 transcription factor playing a pivotal role in the pathogenesis of asthma hyper-eosinophilic syndromes, and eosinophil-dependent inflammatory diseases [57]. Collectively, in the present study, hypercholesterolemia and atorvastatin through down regulation of cytokines with anti-tumor effects might impair cell mediated immunity and probably increase the risk of cancer. Also, these by upregulation of IL-5 cytokine and GATA-3 transcription factor might induce humoral immunity and increase the risk of allergic dependent diseases.

The previous studies showed that increased levels of cholesterol abolish the T cell homeostasis leading to impaired the T cell function [58,59]. A high cholesterol diet can affect the expression and differentiation of T regulatory cells (Treg) [60]. In the present study, it was observed that high level of cholesterol and LDL decreased the expression of CD69 and CD25 activation markers on CD4 and CD8 T cells and reduced memory B cells but had no effect on T memory cells. Furthermore, the correlation analysis revealed that the number of CD4+ CD25+ and CD8+ CD69+ cells is directly associated with the levels of cholesterol and a less extent with the LDL concentration in sera of hypercholesterolemia patients. The ratio of CD4 to CD8 was correlated with the levels of cholesterol as well. It is alleged that cholesterol is needed for stabilization of the cell membrane, especially in lipid raft domain which binds to the chain of TCR and facilitates
dimerization of TCR giving rise to an increase in the affinity of T cell receptors to their cognate antigens [61]. Guo et al. observed that cholesterol can modify receptor signaling through the elevation of different conformational states in membrane receptors and change the composition of T cell membrane resulting to the altered T cell operation [62].

There are contradictory reports about the impact of statins on the expression of CD25 and CD69 on the T cell surface. A study by Blank et al. disclosed that atorvastatin lowered the expression of CD25 and CD69 on the T cells [63], whereas Guo et al. demonstrated that Fluvastatin inhibited CD69 expression in vitro, but it did not affect the expression of CD25 [64]. In this study, we detected decreased expression of CD8+ CD69+, CD8+ CD25+, CD4+ CD69+ cells in response to atorvastatin treatment while the expression of CD4+ CD25+ did not alter. Interestingly, despite a decrease in the number of memory B cells in response to administration of atorvastatin, the number of memory T cells remained unchanged. We also exhibited the decreased number of CD8+ CD25+ cells in the treatment-naive and the atorvastatin groups compared to the healthy controls.

The preventing effect of atorvastatin on the T cell activation is mediated by the disruption of APCs maturation [42]. Additionally, atorvastatin is capable of interfering with the TCR signaling pathways via dysregulation of small Ras-like GTPases and the Rab-dependent pathways that are involved in receptor endocytosis [42]. It also blocks LFA-1–ICAM-1 interaction by binding to LFA-1 [16]. In agreement on our results, Kanda et al. reported that low-dose simvastatin decreased the ratio of CD4 to CD8 T cells when administered in patients with rheumatoid arthritis [65]. We observed a significant decrease in the ratio of CD4/CD8 in peripheral blood of the atorvastatin group. Also, high level of cholesterol and LDL obtained similar results about CD4/CD8 ratios in the treatment-naive group. The low ratio of CD4 to CD8 can raise the risk of cancer because of lacking a sufficient number of Th1 cells [66].

Conclusion

Altogether, regarding our study, it seems that hypercholesterolemia can disrupt immune system through effect of T cell activation and differentiation. On the other hand, besides of beneficial effect of atorvastatin on cholesterol lowering, it diminished the activation of T cells, made a perturbation in inflammatory cytokines, and is not able to repair the deleterious effects of hypercholesterolemia on the immune system.

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References
