Research Article

Statin-induced microRNAome alterations modulating inflammation pathways of peripheral blood mononuclear cells in patients with hypercholesterolemia

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Statins inhibit cholesterol biogenesis and modulate atheroma inflammation to reduce cardiovascular risks. Promoted by immune and non-immune cells, serum C-reactive protein (CRP) might be a biomarker suboptimal to assess inflammation status. Although it has been reported that statins modulated inflammation via microRNAs (miRNAs), evidence remains lacking on comprehensive profiling of statin-induced miRNAome alterations in immune cells. We recruited 19 hypercholesterolemic patients receiving 2 mg/day pitavastatin and 15 ones receiving 10 mg/day atorvastatin treatment for 12 weeks, and performed microarray-based profiling of 1733 human mature miRNAs in peripheral blood mononuclear cells (PBMCs) before and after statin treatment. Differentially expressed miRNAs were determined if their fold changes were >1.50 or <0.67, after validated using quantitative polymerase chain reaction (qPCR). The miRSystem and miTALOS platforms were utilized for pathway analysis. Of the 34 patients aged 63.7 ± 6.2 years, 27 were male and 19 were with coronary artery disease. We discovered that statins induced differential expressions of miR-483-5p, miR-4667-5p, miR-1244, and miR-3609, with qPCR-validated fold changes of 1.74 (95% confidence interval, 1.33–2.15), 1.61 (1.25–1.98), 1.61 (1.01–2.21), and 1.68 (1.19–2.17), respectively. The fold changes of the four miRNAs were not correlated with changes of low-density-lipoprotein cholesterol or CRP, after sex, age, and statin type were adjusted. We also revealed that RhoA and transforming growth factor-β signaling pathways might be regulated by the four miRNAs. Given our findings, miRNAs might be involved in statin-induced inflammation modulation in PBMCs, providing likelihood to assess and reduce inflammation in patients with atherosclerotic cardiovascular diseases.

Introduction

Atherosclerosis is an arterial inflammatory process of atheroma plaque formation initiated by the erroneous accumulation of apolipoprotein B-containing cholesterol, specifically low-density lipoprotein cholesterol (LDL-C), in subendothelial space of arteries [1]. Atherosclerotic cardiovascular disease (ASCVD) is the leading cause of death worldwide, resulting in nearly 12-million deaths from ischemic heart
disease, ischemic stroke, and peripheral arterial disease worldwide [2]. Hypercholesterolemia is one of well-known cardiovascular risk factors, and the risks for coronary heart disease would be raised by 26% and 23% per 1 mmol/l cholesterol increase, in men and women, respectively [3]. Known as an efficacious treatment for hypercholesterolemia, statins lower the risk for major vascular events by 23% per 1 mmol/l LDL-C reduction [4].

Statins have been suggested to exert anti-inflammatory effects on immune cells [5]. The immune cells involved in inflammation progression and resolution of atheroma plaques are mainly from peripheral blood mononuclear cells (PBMCs), including monocytes, lymphocytes, and dendritic cells [6]. Statin could reduce expressions of adhesion molecules in monocyte [7], inhibit proliferation of macrophages in plaques [8], block CD4+ T-cell-mediated apoptosis [9], and suppress miRNA expressions of pro-inflammatory cytokines in PBMCs [10]. Given clinical studies have showed that statins would confer less risk reduction of cardiovascular diseases in individuals with high serum C-reactive protein (CRP) levels [11,12], anti-inflammatory treatment is emerging as a complement to lipid-lowering treatment in ASCVD patients [13]. Accordingly, how to reliably identify ASCVD patients with discernible inflammation in atheroma plaques could be fundamental to implementing anti-inflammatory treatment in clinical setting.

Serum CRP levels might, however, be a biomarker suboptimal to determine inflammation status of atheroma plaques, given that CRP levels could be promoted by immune and non-immune cells. In the Canakinumab Anti-Inflammatory Thrombosis Outcomes Study (CANTOS) recruiting patients with myocardial infarction and high CRP levels of ≥2 mg/dl at baseline, it was revealed that only 55% of patients receiving 150-mg canakinumab achieved CRP levels of <2 mg/dl at 3 months, thereby resulting in significant cardiovascular risk reduction, rather than those with achieved CRP levels of ≥2 mg/dl [14]. A meta-analysis has demonstrated that CVD risk reduction by statin treatment was not associated with achieved level of, or the reduction magnitude of CRP [15]. These findings might be partly because interleukin-6, a potent inducer for CRP production in hepatocytes, is secreted by immune cells [16,17], but also by non-immune cells, such as activated endothelial cells and smooth muscle cells [18,19].

miRNAs (miRNAs) are short non-coding RNAs for post-transcriptional regulations of gene expressions by binding mostly to the 3’ untranslated region of target miRNAs, thus leading to premature degradation and translation suppression of target miRNAs [20]. miRNAs have been reported to mediate statin-induced inflammation modulations [21–23], and, hence, might be feasible biomarkers for assessing inflammation status in immune cells. However, there is a paucity of evidence on comprehensive assessments for statin-induced miRNA alterations in immune cells. We here reported that miRNA alterations in hypercholesterolemic patients before and after receiving statin treatment were discovered using a microarray profiling method in PBMCs, which were an ensemble of immune cells involved in inflammation of atheroma plaques [6].

Materials and methods
Study design and participants
We conducted a sub-study of a multi-center, double-blinded, randomized clinical trial [24], whose study protocol was approved by the research ethics committee of the National Taiwan University Hospital (REC No. 201106075RC). A total of 34 patients, who were recruited in the National Taiwan University Hospital, participated this sub-study, after written informed consents were obtained. Of those, 19 patients received 2 mg pitavastatin daily, and the remaining did 10 mg atorvastatin daily for 12 weeks. Medical history and anthropometric measurements were procured on enrollment.

Blood sampling and isolation of PBMCs
Blood samples were collected after overnight fasting at enrollment and 12 weeks. Laboratory measurements were carried out in a central laboratory, including total cholesterol, triglyceride, high-density-lipoprotein cholesterol (HDL-C), LDL-C, and high-sensitivity CRP. PBMCs were isolated using the Ficoll-Paque PLUS (GE Healthcare) density gradient centrifugation, and were frozen at −20°C for RNA extraction.

RNA extraction and profiling miRNA transcriptome of PBMCs
After PBMCs were properly thawed, total RNA of PBMCs were purified using TRIzol® reagent (Invitrogen). Quantification of RNA was performed using a ND-1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE). We then profiled miRNA alterations using the Affymetrix GeneChip® miRNA 3.0 Array (Affymetrix, Santa Clara, CA), which offered a total of 1733 human mature miRNA probe sets according to the miRBase database version 17. After being polyadenylated, RNA were labeled using the FlashTag™ Biotin HSR RNA Labeling Kit. The biotin-labeled RNAs was hybridized onto microarrays at 48°C and 60 rpm for 16 h. Then, miRNA-hybridized microarrays were stained with phycoerythrín-conjugated streptavidin in a GeneChip® Fluidics Station 450. Finally, the
fluorescence-stained microarrays were scanned for probe signal intensities in a GeneChip® Scanner 3000 7G. The average intensities of the spike-ins and background probe sets were plotted and compared for quality control; and all microarrays achieved the requirements of quality control. The microarray data have been submitted to the Gene Expression Omnibus (accession number: GSE152016).

Selection and validation of miRNAs with stain-induced differential expressions

We applied three methods to identify miRNA candidates with statin-induced differential expression. After microarray intensity signals were pre-processed using the robust multi-array (RMA) method [25], the fold-change method and the Significance Analysis of microarrays (SAM) method were used to select differentially expressed miRNA candidates [26]. The third method was that miRNA signals were fit into linear models for microarray (limma), after pre-processed using the variance stabilizing normalization (VSN) [27,28].

The selected miRNAs candidates were further validated using the quantitative polymerase chain reaction (qPCR) method. Total RNAs were analyzed according to the standard protocol of the miScript PCR System (QIAGEN), and SNORD68 was used as the endogenous control for normalization. miRNAs with differential expressions were defined if the microarray-measured fold changes of $>1.50$ or $<0.67$ were validated by qPCR.

Biological inference of miRNA-modulated signaling pathways

The biological inference of miRNAs was initiated by identifying targeted genes, whose expressions might be modulated by miRNAs. The plausible links between targeted genes and miRNAs could be determined by well-established algorithms or experimental evidences. Based on dedicated database, signaling pathways were fitted by the targeted genes, and were scored and ranked using over-representation enrichment analysis. It was, therefore, inferred that biological pathways with higher ranking were more likely to be modulated by the miRNAs. The web-based miRSystem [29] and miTALOS [30] were platforms designed to incorporate the process of identifying targeted genes and pathway enrichment analysis. Accordingly, we applied the miRNAs with qPCR-validated differential expressions to miRSystem and miTALOS platforms to perform biological inference of miRNA-modulated signaling pathways. We used default settings of the miRSystem, while conducted the leukocyte-confined pathway analysis in the miTALOS platform. Besides, the normalized expressions of targeted genes in PBMCs were adopted from the Blood Atlas of the Human Protein Atlas database [31].

Statistical analysis

Continuous variables were presented as mean and standard deviation; and categorical ones, as number and percentage, unless otherwise specified. Categorical variables were tested using the chi square test or Fisher exact test, where appropriate.

The microarray intensity data were pre-processed using the robust multi-array (RMA) or variance stabilizing normalization (VSN) method [25,32]. The RMA method consists of background correction, quantile normalization, logarithm transformation, and then median polish summarization [25]. Because of the dependence of the variance on the mean intensity, especially in the miRNAs with low signal intensities, the VSN method was proposed to estimate mean-independent variance based on an additive-multiplicative error model fitting generalized logarithm-transformed intensity data after affine calibration [32]. Fold changes of miRNA intensities were used to represent the alterations of miRNA expressions between before-and-after treatment. Given PMBCs were mostly in naïve status, fold changes of miRNAs could be lower in PMBCs than immune cells in atheroma plaques [33]. Accordingly, we defined miRNAs as being differentially expressed if fold change of $>1.50$ or $<0.67$, along with the $q$ value or the false discovery rate of $<0.05$ for the correction of multiple comparisons [34]. Although miRNAs with differential expressions could be selected using the Student’s $t$ test, the stability of $t$ statistic is subject to the small sample size and low expression levels. SAM and limma methods were recommended to address the inappropriateness of the Student’s $t$ test by adjusting variance estimations in a nonparametric or parametric way, respectively. The SAM method calculated a statistic score, whose $q$ value was derived according to permutation estimation [26]. The limma method used the empirical Bayes approach to estimate robust gene-specific variances of linear models for calculating a moderated $t$ statistic, which followed a $t$ distribution under null hypothesis [28]. Benjamini–Hochberg adjustment for multiple comparisons was used to control family-wise error rate in the limma method [35].

Correlations of post-treatment lipid and CRP changes with the SAM-derived fold changes of differentially expressed miRNAs were assessed after sex, age, and statin type were adjusted. To delineate the relationship of differentially expressed miRNAs with baseline CRP levels, we compared log-transformed fold changes among the CRP...


### Results

Among the 34 hypercholesterolemic patients aged 63.7 ± 6.2 years (Table 1), there were 19 (55.9%) patients with stable coronary heart disease, 21 (61.8%) ones with Type 2 diabetes mellitus under oral hypoglycemic treatment, 32 (94.1%) ones with hypertension. And only 5 (14.7%) patients were current smokers (Supplementary Table S1). The proportions of patients with coronary heart disease, diabetes mellitus, hypertension, and smoking habit were not different between the pitavastatin and atorvastatin subgroups ($P = 0.092$, $0.16$, $0.19$, and $0.63$, respectively).

While the reduction percentage of the LDL-C level was 34.5 ± 17.3% after 12-week statin treatment, pitavastatin and atorvastatin treatment achieved similar post-treatment decrease in LDL-C levels (reduction percentage, 32.8 ± 17.5% versus 36.6 ± 17.5%; $P = 0.53$). Among 34 patients, only one had CRP level of greater than 1 mg/dl (Table 1). Statin treatment did not significantly lower serum CRP level (pre-treatment, 0.15 ± 0.24 versus post-treatment, 0.19 ± 0.42 mg/dl; $P = 0.66$) (Supplementary Table S1). Similarly, there was no difference in CRP changes between the pitavastatin or atorvastatin subgroup ($P = 0.67$, 0.34, respectively).
Figure 1. Statin-induced differential expressions of miRNAs in peripheral blood mononuclear cells (PBMC)

miRNAome alterations were profiled using microarrays in 34 hypercholesterolemic patients before and after 12-week statin treatment. Fold changes were defined as the ratios of post-treatment to pre-treatment miRNA signal intensities. miRNAs were considered as having statin-induced differential expressions, if their microarray-profiled fold changes were >1.50 or <0.67 with quantitative polymerase chain reaction (qPCR) validation. * indicated the miRNAs with qPCR-validated fold changes of >1.50 or <0.67. Dotted vertical lines represented as fold changes of 1.50 and 0.67, respectively.

Based on microarray signal intensities, a total of 21 miRNAs candidates with fold changes of >1.50 or <0.67 were selected for qPCR validation. Among them, there were 10 miRNAs having the microarray fold changes in concordance with ones of qPCR validation, in which reference sequences were listed in the Supplementary Table S2. As shown in the Figure 1, we identified that statins induced differential expressions of miR-483-5p, miR-4667-5p, miR-1244, and miR-3609, with qPCR-validated fold changes of 1.74 (95% confidence interval, 1.33–2.15), 1.61 (1.25–1.98), 1.61 (1.01–2.21), and 1.68 (1.19–2.17), respectively.

In the Figures 2 and 3, we explored that, after statin treatment, the fold changes of the four miRNAs would be related to magnitudes of lipid changes and baseline CRP levels, respectively. The Figure 2 showed the age-, sex-, and statin type-adjusted associations between lipids and fold changes of the four differentially expressed miRNAs. While the fold changes of miR-4667 and miR-1244 appeared to correlate with post-treatment changes of total cholesterol and HDL-C (adjusted $r = -0.41$ and 0.38, $P=0.023$ and 0.033, respectively), the associations were not found between the fold changes of miRNAs and LDL-C or triglyceride changes, suggesting that there were greater individual variations in statin-induced miRNA alterations than those in lipid changes. On the other hand, the fold changes of miRNAs were not different among the baseline CRP tertiles (Figure 3), and nor were correlated with post-treatment CRP changes (sex-, age, and statin type-adjusted correlation coefficient, $r = −0.07, 0.001, −0.02, and −0.08; P=0.70, 0.97, 0.91,$ and 0.69, for miR-483-5p, miR-4667-5p, miR-3609, and miR-1244, respectively), probably because serum CRP levels could be affected by the IL-6 production of inflammatory and non-inflammatory cells.

Signaling pathways were determined using miRSystem (Table 2A) and miTALOS (Table 2B) platforms, respectively, in the descending order of likelihood that some or all of the four differentially expressed miRNAs could be involved in modulation of biological signaling.

Among the four differentially expressed miRNAs, only miR-483 was identified by at least three of the miRSystem-integrated prediction programs. The Table 2 outlined the signaling pathways enriched by targeted genes of the differentially expressed miRNAs according to the analysis using the miRSystem and miTALOS platforms. In Table 2A and Supplementary Table S3, it was revealed that RHOA was the mostly expressed in PBMCs among the
Figure 2. Associations between magnitudes of lipid changes and fold changes of the four miRNAs after statin treatment
Correlations of serum lipid changes with fold changes of the four differentially expressed PBMC miRNAs in hypercholesterolemic patients receiving 12-week statin treatment. Lipid changes were defined as (post-treatment minus pre-treatment lipid level)/pre-treatment lipid level. Fold changes of miRNAs were the ratios of post-treatment to pre-treatment microarray-profiled signal intensities. Correlations were adjusted for sex, age, and statin type. Dotted lines represented linear regression lines.

Figure 3. Alterations in statin-induced fold changes of the four PMBC-derived miRNAs across the tertiles of baseline serum CRP levels
Comparing the statin-induced fold changes of the four differentially expressed PBMC miRNAs after 12-week statin treatment according to the tertiles of baseline CRP levels. The fold changes of miRNAs were defined as the ratios of post-treatment to pre-treatment microarray-profiled signal intensities derived from microarray profiling. The comparisons were adjusted for sex, age, and statin type.
Table 2: Identification of potential signaling pathways modulated by the four microRNAs with statin-induced differential expressions according to the miRSystem (A) and the miTALOS (B) platforms dedicated to integrating the process of targeted gene exploration and pathway enrichment analysis

| Database | Signaling pathway | Brief description of signaling pathway | Targeted genes | miRNAs | Ranking score |
|----------|-------------------|----------------------------------------|----------------|--------|---------------|
| **(A) The most relevant ten pathways identified using the miRSystem platform** | | | | | |
| KEGG Pathway interaction database | Leishmaniasis | Disease-specific pathway | ELK1, MARCKSL1 | miR-483 | 0.75 |
| | PDGFR-β signaling pathway | PDGFR-β is a member of PDGF family, and could activate Ras to promote cell proliferation. | RHOA, ELK1 | miR-483 | 0.626 |
| Reactome | NGF signaling via TRKA from the plasma membrane | TRK signaling regulates cell proliferation and neuronal differentiation. | RHOA, ELK1 | miR-483 | 0.609 |
| KEGG | Focal adhesion | The biological processes regulate cell motility, proliferation, differentiation, survival, and gene expressions. | RHOA, ELK1 | miR-483 | 0.527 |
| Reactome | Signaling by neurotrophin receptor | Signaling networks relating to neuronal survival, differentiation, and plasticity. | RHOA, ELK1 | miR-483 | 0.506 |
| Pathway interaction database | Endogenous TLR signaling | A family of pattern recognition receptors are involved in innate and adaptive immunity, and could be triggered by endogenous ligands in inflammation conditions. | RHOA | miR-483 | 0.451 |
| Reactome | Gβγ signaling through PK3Ky | PK3Ky is highly expressed in neutrophils, and is activated by membrane-bound Gβγ. | RHOA | miR-483 | 0.451 |
| Biocarta | BIOCARTA_RAC_7000_PATHWAY | Regulation of G1 to S transition involving Ras, Rac, Rho, and cyclin D1 expression. | RHOA | miR-483 | 0.446 |
| Biocarta | BIOCARTA_7000_PATHWAY | Phospholipids as signaling intermediaries involving regulations of actin assembly, chemotaxis, proliferation and cell survival. | RHOA | miR-483 | 0.442 |
| Biocarta | BIOCARTA_7000_PATHWAY | Erk1/Erk2 MAPK signaling pathway to regulate growth, differentiation, and development. | ELK1 | miR-483 | 0.438 |
| **(B) Pathways identified using the miTALOS platform** | | | | | |
| KEGG | TGF-β signaling pathway | Biological processes regulating cell growth, apoptosis, homeostasis, and differentiation. | TGFBR2, SMAD2, SMAD4, SMAD7, SMURF1, SMURF2, BMP2, ACVR1B, ACVR2A, AP21, EP200, TNF, PPP3CA, MAPK1, RHOA | | 0.008 |
| WikiPathways | Physiological and pathological hypertrophy of the heart | Disease-specific signaling network | RHOA, MAPK1, FOS, ILBSt, PRKCE, PPP3CA, SPAG1 | | 0.016 |
| WikiPathways | Mesodermal commitment pathway | Biological processes involving development | BMP2, SMAD2, SMAD4, ACVR2A, PHF6, NLK, NFE2L2, TOX, C9orf72, WDCP, AEBP2, PARP8, ATPHB2, CCDO1, EMSY, EXT2, LBR5, SETD2, PPP3CA, PBX3 | | 0.042 |

Abbreviations: Gβγ, β and γ subunits of G-protein; KEGG, Kyoto Encyclopedia of Genes and Genomes; MAPK, mitogen-activated protein kinase; NGF, nerve growth factor; PK3Ky, phosphatidylinositol 3-kinase-β; PDGF, platelet-derived growth factor; PDGFR, platelet-derived growth factor receptor; TGF-β, transforming growth factor β; TLR, Toll-like receptor
targeted genes of miR-483, and was the mostly identified among the enriched signaling pathways. As for the most relevant ten signaling identified by the miRSystem, the relevant pathways in the context of PBMCs included focal adhesion, endogenous Toll-like receptor (TLR) signaling, and intracellular signaling networks related to Gβγ protein, cyclin D1, and sphingosine 1-phosphate (Table 2B). While confined to the gene expressions of leukocytes, the mi-TALOS analysis indicated that transforming growth factor-beta (TGF-β) signaling pathway was the most relevant to statin-induced miRNA alterations.

Discussions
In hypercholesterolemic patients receiving 12-week statin treatment, we found that statin treatment could induce miRNA alterations in PBMCs consisting of immune cells involved in the inflammation reactions of atheroma plaques. Four miRNAs of miR-483-5p, miR-4667-5p, miR-3609, and miR-1244 with statin-induced increased expressions were determined using unbiased microarray profiling along with qPCR validations. The four miRNAs were not associated with post-treatment serum LDL-C reduction, nor with the baseline level or post-treatment reduction in serum CRP. Exploring the biological relevance of the four miRNAs with statin-induced differential expressions in PBMCs, we revealed that signaling pathways involving inflammation reactions were relevant to the post-treatment miRNA alterations, including RhoA-related focal adhesion and TGF-β signaling. Our findings indicated that the four miRNAs might be involved in statin-induced inflammation modulations, and thus be feasible biomarkers for assessing inflammation status in response to statin treatment.

Statin-induced miRNA regulations of RhoA GTPase expression
The mechanism for statin-induced anti-inflammatory effects has been proposed that isoprenoid intermediates of cholesterol biosynthesis are collectively suppressed by statins, thus reducing the post-translational prenylation of Rho and Rac GTPases [12]. Prenylation with the isoprenoid intermediates, including farnesylpyrophosphate (FPP) and geranylgeranylpyrophosphate (GGPP) [37], facilitates Rho and Rac GTPases attached onto cell membranes, where these GTPases are normally functional for signaling transductions. RhoA binds the downstream effector, Rho-associated coiled-coil-containing protein kinase (ROCK), and then alters cytoskeleton through myosin light chain or cofilin phosphorylation [38,39], whereby focal adhesion and transendothelial migration of monocytes and T-cells are regulated [40,41].

The relationships between miRNAs and RhoA signaling in circulating immune cells are yet to be elucidated, though it has been evidenced that miRNAs were involved in post-transcriptional regulations of RhoA signaling in non-immune cells. Prior studies have reported that miR-31, miR-125a-3p, miR-133a, miR-155, and miR-185 decreased RhoA expressions in osteoclasts, lung cancer cells, cardiomyocytes, breast cancer cells, and colorectal cancer cells, respectively [42–46]. In the present study, we discovered the association between statin treatment and miR-483-5p expressions. Given that RhoA is essential to signaling of focal adhesion and transendothelial migration of immune cells, statin-induced increase in miR-483-5p expression might repress RhoA-mediated inflammatory reactions.

Statin-induced miRNA regulations of context-dependent TGF-β signaling
TGF-β exerts context-dependent effects on innate and adaptive immune responses via canonical and non-canonical pathways, thereby leading to dual pro-inflammatory and anti-inflammatory roles in the progression and resolution of atheroma plaques. The canonical pathway is mediated in the Smad-dependent way [47], while the non-canonical signaling is mediated in Smad-independent ways, such as inter-connections with RhoA/ROCK signaling pathway [48]. TGF-β is a pro-inflammatory chemoattractant to induce migration of monocytes and dendritic cells [49,50]. On the other hand, the anti-inflammatory effects of TGF-β has been implicated because the disruption of TGF-β signaling in T-cells could lead to atherosclerosis progression [51]. However, when incorporating different interleukin cytokines, TGF-β could facilitate or inhibit CD4+ T-cells differentiation into distinct T-cell subsets, thereby promoting or resolving inflammation in atheroma plaques. In the presence of TGF-β, CD4+ T-cells differentiation are inhibited into Th1 and Th2 cells; but are promoted into Th9, Th17, and Treg when coupled with IL-4, IL-6, or IL-2, respectively [52]. In general, Th1, Th9, and Th17 cells mediate pro-inflammatory reactions, while Th2 and Treg cells do anti-inflammatory ones, though some evidence showed that Th17 cells might have dual effects on plaque stability [53]. Besides, the effects of TGF-β are inhibitory and pro-apoptotic for CD8+ T-cells, NK cells, and B cells [54,55].

Statins have been reported to modulate TGF-β expressions and signaling in immune and non-immune cells. In cultured THP-1 cells, TGF-β expression and production were up-regulated by pravastatin in a dose-dependent manner [56]. A clinical study found that, in hypercholesterolemic patients treated with daily 40 mg pravastatin for 4–6
weeks, post-treatment serum TGF-β levels were elevated, but also that isolated peripheral blood monocytes increased TGF-β expression and production, independently of post-treatment lipid changes [57]. However, there is a paucity of evidence regarding how statins modulate TGF-β signaling. Our findings revealed an in-silico link between TGF-β signaling and statin-induced miRNA alterations in PBMCs. miR-483-5p, miR-4667-5p, miR-3609, and miR-1244 might regulate the expressions of extensive elements in the TGF-β signaling pathway, including the expressions of TGF-β receptors, Smad proteins, Smad inhibitory proteins, transcriptional factors, and non-Smad proteins, such as RhoA. More studies are required to determine the responses of PBMCs when TGF-β signaling is modulated by those miRNAs in diverse pathophysiological settings.

**Study limitations**

Some limitations in our study should be reminded. First, PBMCs are limited to delineate statin-induced miRNAome alterations in distinct immune cell subsets. Interindividual variations on proportions of immune cell types in PBMCs might make it less feasible to detect miRNAs with low expression amount and high variability [33]. Second, miRNA extraction from PBMCs appeared to be more technique-demanding than from plasma, resulting in the concern on the feasibility in clinical practice. However, changes of circulating miRNAs could not be comparable to those of intracellular miRNAs, partly because not all miRNAs in immune cells are secreted in exosomes [58]. Third, as for high-throughput detection of miRNAs, microarray profiling is known to be less specific than RNA sequencing [59]. Given that the purpose of our study was to explore the associations between statin treatment and miRNA-regulated inflammatory responses in immune cells, microarray profiling could be adequate to detect statin-induced miRNA alterations [59]. Fourth, confining pathway analysis to healthy leukocytes in the miTALOS platform might increase the biological relevance at the expense of exploring unexpected connections in hypercholesterolemia-primed PBMCs. Considering that, we combined the results of biological relevance derived from an unconfined and a confined pathway analysis using the miRSystem and the miTALOS platforms, respectively.

**Perspectives**

1. Given that miRNAs mediate in statin-induced anti-inflammation, and that CRP might be suboptimal to assess inflammation status in ASCVD patients, evidence remains lacking on comprehensive profiling of statin-induced miRNAome alterations in circulating immune cells.

2. We profiled miRNAome alterations in hypercholesterolemic patients before and after statin treatment, and discovered that statin treatment led to the increased expressions of miR-483-5p, miR-4667-5p, miR-1244, and miR-3609 in PBMCs. And the four miRNAs with statin-induced differential expressions were involved in RhoA and transforming growth factor-β signaling pathways, probably modulating transendothelium migration and differentiations of immune cells.

3. Our findings implicated that miRNAs of PBMCs might be feasible to assess inflammation status, but also be therapeutic potentials to modulate inflammation in ASCVD patients.

**Competing Interests**

The authors declare that there are no competing interests associated with the manuscript.

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**Author Contribution**

H.J.L. was involved in study design, enrolment and treatment of patients, statistical analysis, and manuscript writing. S.L.Y. participated in the design of the study and carried out microarray experiments. T.C.S., H.C.H., M.F.C., and Y.T.L. conducted the study design, coordinated the study, and interpreted the results. K.L.C. was involved in study design, enrolment and treatment of patients, statistical analysis, and manuscript editing. T.P.L. performed laboratory work and statistical analysis, and edited the manuscript.

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Abbreviations
ASCDV, atherosclerotic cardiovascular disease; CRP, C-reactive protein; LDL-C, low-density lipoprotein cholesterol; PBMC, peripheral blood mononuclear cell; qPCR, quantitative polymerase chain reaction.

References
1 Weber, C. and Noels, H. (2011) Atherosclerosis: current pathogenesis and therapeutic options. Nat. Med. 17, 1410–1422, https://doi.org/10.1038/nm.2538
2 (2018) Global, regional, and national age-sex-specific mortality for 282 causes of death in 195 countries and territories, 1980-2017: a systematic analysis for the Global Burden of Disease Study 2017. Lancet 392, 1736–1788, https://doi.org/10.1016/S0140-6736(18)32203-7
3 Group, W.C.R.C.W. (2019) World Health Organization cardiovascular disease risk charts: revised models to estimate risk in 21 global regions. Lancet Glob. Health 7, e1332-e1345, https://doi.org/10.1016/S2214-109X(19)30318-3
4 Silverman, M.G., Ference, B.A., Im, K., Wiviott, S.D., Giugliano, R.P., Grundy, S.M. et al. (2016) Association Between Lowering LDL-C and Cardiovascular Risk Reduction Among Different Therapeutic Interventions: A Systematic Review and Meta-analysis. JAMA 316, 1289–1297, https://doi.org/10.1001/jama.2016.13985
5 Tousoulis, D., Psarros, C., Demosthenous, M., Patel, R., Antoniades, C. and Stefanadis, C. (2014) In innate and adaptive inflammation as a therapeutic target in vascular disease: the emerging role of statins. J. Am. Coll. Cardiol. 63, 2491–2502, https://doi.org/10.1016/j.jacc.2014.01.054
6 Kleiveland, C.R. (2015) Peripheral Blood Mononuclear Cells. In The Impact of Food Bioactives on Health: in vitro and ex vivo models (Verhoeckx, K., Cotter, P., Lopez-Exposito, I., Kleiveland, C., Lea, T., Mackie, A., Requena, T., Swiatecka, D. and Wichers, H., eds), pp. 161–167. Springer Copyright 2015, Cham (CH)
7 Wojcicki-Stothard, B., Williams, L. and Ridley, A.J. (1999) Monocyte Adhesion and Spreading on Human Endothelial Cells Is Dependent on Rho-regulated Receptor Clustering. J. Cell Biol. 145, 1293–1307, https://doi.org/10.1083/jcb.145.12.1293
8 Tang, J., Lobatto, M.E., Hassing, L., van der Staay, S., van Rijss, S.M., Calagnino, C. et al. (2015) Inhibiting macrophage proliferation suppresses atherosclerotic plaque inflammation. Sci. Adv. 1, e1400223, https://doi.org/10.1126/sciadv.1400223
9 Sato, K., Nuki, T., Gomiya, K., Weyand, C.M. and Hagivara, N. (2010) Statins reduce endothelial cell apoptosis via inhibition of TRAIL expression on activated CD4 T cells in acute coronary syndrome. Atherosclerosis 213, 33–39, https://doi.org/10.1016/j.atherosclerosis.2010.03.034
10 Cerda, A., Rodrigues, A.C., Alves, C., Genovig, F.D., Fajardo, C.M., Nogueira, E.L. et al. (2015) Modulation of adhesion molecules by cholesterol-lowering therapy in mononuclear cells from hypercholesterolemic patients. Cardiovasc. Ther. 33, 168–176, https://doi.org/10.1111/1755-5922.12126
11 Ridker, P.M., Danielson, E., Fonseca, F.A., Genest, J., Gotto, Jr, A.M., Kastelein, J.J. et al. (2008) Rosuvastatin to prevent vascular events in men and women with elevated C-reactive protein. N. Engl. J. Med. 359, 2195–2207, https://doi.org/10.1056/NEJMoa0807646
12 Gesterle, A., Laufs, U. and Liao, J.K. (2017) Pleiotropic Effects of Statins on the Cardiovascular System. Circ. Res. 120, 229–243, https://doi.org/10.1161/CIRCRESAHA.116.308537
13 Ridker, P.M., Everett, B.M., Thuren, T., MacFadyen, J.G., Wang, L., Kang, L.N. et al. (2017) Antiinflammatory Therapy with Canakinumab for ASCVD, atherosclerotic cardiovascular disease; CRP, C-reactive protein; LDL-C, low-density lipoprotein cholesterol; PBMC, peripheral blood mononuclear cell; qPCR, quantitative polymerase chain reaction.
53 Gistera, A., Robertson, A.K., Andersson, J., Ketelhuth, D.F., Ovchinnikova, O., Nilsson, S.K. et al. (2013) Transforming growth factor-beta signaling in T cells promotes stabilization of atherosclerotic plaques through an interleukin-17-dependent pathway. Sci. Transl. Med. 5, 196ra100, https://doi.org/10.1126/scitranslmed.3006133

54 Tamayo, E., Alvarez, P. and Merino, R. (2018) TGFbeta Superfamily Members as Regulators of B Cell Development and Function-Implications for Autoimmunity. Int. J. Mol. Sci. 19, 3928–3945, https://doi.org/10.3390/ijms19123928

55 Viel, S., Marcais, A., Guimaraes, F.S., Loftus, R., Rabilloud, J., Grau, M. et al. (2016) TGF-beta inhibits the activation and functions of NK cells by repressing the mTOR pathway. Sci. Signal 9, ra19, https://doi.org/10.1126/scisignal.aad1884

56 Baccante, G., Mincione, G., Di Marcantonio, M.C., Piccirelli, A., Cuccurullo, F. and Porreca, E. (2004) Pravastatin up-regulates transforming growth factor-beta1 in THP-1 human macrophages: effect on scavenger receptor class A expression. Biochem. Biophys. Res. Commun. 314, 704–710, https://doi.org/10.1016/j.bbrc.2003.12.150

57 Porreca, E., Di Febbo, C., Baccante, G., Di Nisio, M. and Cuccurullo, F. (2002) Increased transforming growth factor-beta(1) circulating levels and production in human monocytes after 3-hydroxy-3-methyl-glutaryl-coenzyme a reductase inhibition with pravastatin. J. Am. Coll. Cardiol. 39, 1752–1757, https://doi.org/10.1016/S0735-1097(02)01857-0

58 Montecalvo, A., Larregina, A.T., Shufesky, W.J., Stolz, D.B., Sullivan, M.L., Karlsson, J.M. et al. (2012) Mechanism of transfer of functional microRNAs between mouse dendritic cells via exosomes. Blood 119, 756–766, https://doi.org/10.1182/blood-2011-02-338004

59 Pritchard, C.C., Cheng, H.H. and Tewari, M. (2012) MicroRNA profiling: approaches and considerations. Nat. Rev. Genet. 13, 358–369, https://doi.org/10.1038/nrg3198