Rheumatoid arthritis (RA) patients are at a 2-fold risk for cardiovascular (CV) disease compared with the general population. This elevated risk is largely attributed to inflammation rather than to an increased prevalence of traditional CV risk factors such as hyperlipidemia. In fact, RA patients have lower total cholesterol (TC) and low-density lipoprotein cholesterol (LDL-C) levels than those in the general population. This apparent paradox may be explained by favorable changes in other lipid measurements. The objective of this study was to determine the longitudinal association between changes in inflammation with advanced lipoprotein measurements and high-density lipoprotein (HDL) cholesterol efflux capacity.

**Methods and Results**—We conducted this study in a longitudinal RA cohort from a large academic center, including subjects with high-sensitivity C-reactive protein (hs-CRP) reduction ≥ 10 mg/L at 2 time points 1 year apart. Subjects receiving statins during the study period or preceding 6 months were excluded. We compared total cholesterol, LDL cholesterol, HDL cholesterol, apolipoprotein B, and apolipoprotein A1 levels and HDL cholesterol efflux capacity at baseline and 1-year follow-up by using the paired t test. We also assessed the correlations between reductions in hs-CRP with percentage change in lipid parameters. We studied 90 RA subjects (mean age 57 years, 89% female), all of whom were receiving disease-modifying antirheumatic drugs. We observed a 7.2% increase in LDL cholesterol levels (P = 0.02) and improvement in efflux capacity by 5.7% (P = 0.002) between baseline and follow-up, with a median hs-CRP reduction of 23.5 mg/L. We observed significant correlations between reductions in hs-CRP with increases in apolipoprotein A1 (r = 0.27, P = 0.01) and HDL cholesterol efflux capacity (r = 0.24, P = 0.02).

**Conclusion**—Among RA subjects experiencing reductions in hs-CRP, we observed increased LDL cholesterol levels and concomitant improvements in HDL cholesterol efflux capacity. These findings provide further insight into lipid modulation and the beneficial effect of reduction in inflammation on lipids in vivo. (J Am Heart Assoc. 2015;4:e001588 doi: 10.1161/JAHA.114.001588)

**Key Words:** inflammation • lipids • rheumatoid arthritis
cholesterol efflux capacity, which is the ability of HDL particles to extract cholesterol from lipid-laden macrophages, may be better correlated. Few studies have evaluated inflammation with longitudinal changes in HDL efflux capacity, because both inflammation and HDL efflux capacity are considered relatively stable in the general population in the absence of treatment. In RA, there is evidence that level of inflammation is associated with HDL cholesterol efflux capacity. A cross-sectional study (n=25) found that RA patients with a high level of disease activity had impaired HDL function compared with subjects with low disease activity. HDL cholesterol efflux capacity is associated with an increased risk of CV disease independent of HDL-cholesterol (HDL-C) levels and may be a more useful marker for CV risk prediction for patients with RA and other inflammatory diseases compared with the use of LDL-C as a marker.

Current CV risk estimators such as the Framingham Risk and Reynolds’ Risk Score underestimate CV risk in RA by 2-fold in women with RA. Understanding the relationship between inflammation and changes in lipid parameters is a key step for informing CV risk management in RA. The objective of this study was to provide an overview of changes in routine lipids including TC, LDL-C, and HDL-C, as well as advanced lipoprotein measurements including apolipoprotein (apo)B, apoA1, and HDL cholesterol efflux capacity in RA patients who experience a significant reduction in inflammation as measured with high-sensitivity CRP. Our second objective was to quantify the correlation between longitudinal changes in CRP with changes in these lipid parameters. We hypothesized that a reduction in CRP would be associated with increases in LDL-C and concomitant improvements in HDL cholesterol efflux capacity.

Methods
We conducted this study in the Brigham and Women’s Hospital Rheumatoid Arthritis Sequential Study (BRASS), a prospective observational cohort study. All subjects were age 18 or older and had a rheumatologist diagnosis of RA. Subjects in BRASS had annual clinical assessments including measurements of RA disease activity, assessments of CV risk factors, and CRP. Details of this cohort were previously reported. We included subjects in BRASS who experienced a CRP reduction ≥10 mg/L at 2 consecutive time points, 1 year apart, and with blood samples available for analysis. Samples were measured for TC, LDL-C, HDL-C, apoA1, apoB, and HDL cholesterol efflux capacity. Because statins are potent LDL-C–lowering agents, we included only patients who did not receive statins 6 months prior to the baseline blood collection date and during the 1-year follow-up period.

High-sensitivity CRP was measured by using a standard immunoturbidimetric assay on the Roche P Modular system (Roche Diagnostics) with reagents and calibrators from Roche. In this assay, an antigen–antibody reaction occurs between CRP in the sample and an anti-CRP antibody sensitized to latex particles resulting in agglutination. The antigen–antibody complex is detected spectrophotometrically with the magnitude of change proportional to the concentration of CRP in the sample. TC, LDL-C, HDL-C, apoA1, and apoB measurements were performed according to standardized techniques in the clinical laboratories. We performed HDL cholesterol efflux capacity per published methods by using J774 cells derived from a murine macrophage cell line. Briefly, the J774 cells were plated and radiolabeled with 2 μCi of 3H-cholesterol/mL. ATP-binding cassette transporter A1 (ABCA1) was up-regulated by means of a 16-hour incubation with 0.3 mmol/L 8-(4-chlorophenylthio)-cAMP. We added 2.8% apoB-depleted serum to the efflux medium for 4 hours. To quantify the efflux of radioactive cholesterol from the cells, we used liquid scintillation counting. Efflux was calculated by using the following formula: ([microcuries of 3H-cholesterol in media containing 2.8% apoB-depleted serum-microcuries of 3H-cholesterol in serum-free media]/[microcuries of 3H-cholesterol in cells extracted before the efflux step]). All assays were performed in duplicate.

We tested for significant differences in TC, LDL-C, HDL-C, apoA1, apoB, and HDL cholesterol efflux capacity at baseline and 1-year follow-up by using the paired t test. For the primary analyses, we determined the correlations between changes in CRP and the percentage change in each lipid parameter [(lipidfollow-up−lipidbaseline)/lipidbaseline] by using the Pearson correlation test. Because of the non-normal distribution of CRP, we performed all correlation and association studies by using the natural log of the change (reduction) in CRP (CRPbaseline−CRPfollow-up). We performed a sensitivity analysis by using a Spearman rank correlation test between the change in CRP (CRPbaseline−CRPfollow-up) and percentage change in the lipid parameters.

As a secondary analysis, we tested for confounding of the association between changes in CRP and lipids by age and sex. Additionally, we tested the association between the change in CRP and HDL cholesterol efflux capacity adjusted by change in HDL-C levels in one model and change in apoA1 levels in a second model. To determine whether the changes in the lipid parameters may be more specific to serum markers of inflammation compared with other clinical measures of RA disease activity, we tested the association between changes in the RA Disease Activity Score 28 (DAS28-CRP) and changes in HDL cholesterol efflux capacity. The DAS28 is an index score that includes objective and subjective clinical signs and symptoms of RA: the swollen and tender joint count of 28 joints, CRP level, and the patient

DOI: 10.1161/JAHA.114.001588
global score of arthritis disease activity over the past week (scale of 0 to 10, with 0 being no disease activity and 10 being the worst disease activity). High disease activity as defined by DAS28-CRP is a score of >5.1, moderate disease activity is >3.2 and ≤5.1, low disease activity is ≥2.6 to ≤3.2, and remission is <2.6.

All aspects of this study were approved by the Partners Healthcare Institutional Review Board, and the subjects gave informed consent. Analyses were performed by using SAS 9.2 (SAS Institute).

Results

We studied 90 RA cases with a mean age of 57.0; 89% were female, 78% were anti-CCP positive, and they had a mean RA disease duration of 16.5 years (Table 1). The median CRP at baseline was 28.6 mg/L, and the mean DAS28 was 5.0 (moderate disease activity). The median CRP at 1-year follow-up was 4.3 (IQR 5.6), an absolute reduction of 23.5 mg/L (IQR 54.0, P<0.0001), and the mean reduction in DAS28-CRP was 1.6 (SD 1.5, P<0.0001). This represented an overall 85% reduction in median CRP and a significant change in the DAS28-CRP from the higher end of moderate to low disease activity. All subjects were receiving ≥1 DMARD with the majority of subjects receiving methotrexate (53%) or a TNFi (48%); 27% were receiving combination MTX and TNFi at baseline.

Compared with baseline, we observed a significant increase in LDL-C by 7.2% (P=0.02) and improvements in HDL cholesterol efflux capacity by 5.7% (P=0.002) (Table 2). There was a trend toward increased apoA1 of 4.1% (P=0.05). We observed no significant changes in HDL-C, apoB, and 2 measures of the atherogenic index TC/HDL-C or apoB/apoA1 between baseline and 1-year follow-up.

We observed significant correlations between a reduction in CRP with increases in LDL-C, HDL-C, and apoA1 and improvements in HDL cholesterol efflux capacity (Figure). No significant correlation was observed between change in CRP and apoB levels (r=0.14, P=0.20) and the atherogenic indices (TC/HDL-C, r=−0.21, P=0.05 and apoB/apoA1, r=−0.10, P=0.33). In the sensitivity analysis, the significant correlation

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>N=90</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean (SD) y</td>
<td>57.0 (12.0)</td>
</tr>
<tr>
<td>Female, %</td>
<td>88.9</td>
</tr>
<tr>
<td>RA disease duration, mean (SD) y</td>
<td>16.5 (11.5)</td>
</tr>
<tr>
<td>Anti-CCP positive, n (%)</td>
<td>70 (77.8)</td>
</tr>
<tr>
<td>RF positive, n (%)</td>
<td>67 (74.4)</td>
</tr>
<tr>
<td>BMI, mean (SD) kg/m²</td>
<td>27.0 (5.5)</td>
</tr>
<tr>
<td>Diabetes mellitus, n (%)</td>
<td>11 (12.2)</td>
</tr>
<tr>
<td>Hypertension, n (%)</td>
<td>19 (21.0)</td>
</tr>
<tr>
<td>History of ischemic heart disease, n (%)</td>
<td>31 (34.4)</td>
</tr>
<tr>
<td>Hs-CRP, median (IQR), mg/L</td>
<td>28.6 (21.7)</td>
</tr>
<tr>
<td>DAS28, mean (SD)</td>
<td>5.0 (1.6)</td>
</tr>
<tr>
<td>RA treatment, n (%)</td>
<td>Methotrexate 44 (53.1)</td>
</tr>
<tr>
<td>Tumor necrosis factor inhibitor</td>
<td>40 (48.2)</td>
</tr>
<tr>
<td>Other DMARDs</td>
<td>13 (14.4)</td>
</tr>
</tbody>
</table>

Anti-CCP indicates anti-cyclic citrullinated peptide; RA, rheumatoid arthritis; RF, rheumatoid factor; BMI, body mass index; hs-CRP, high-sensitivity C-reactive protein; DAS28, Disease Activity Score 28; DMARD, disease-modifying anti-rheumatic drug.

Table 2. Mean Change in Lipids, Lipoproteins, and HDL Cholesterol Efflux Capacity Between Baseline and 1-Year Follow-up (N=90)

<table>
<thead>
<tr>
<th>Measurement</th>
<th>Baseline (SD)</th>
<th>Follow-up (SD)</th>
<th>Mean Δ (SD)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cholesterol, mg/dL</td>
<td>187.2 (43.1)</td>
<td>183.2 (37.0)</td>
<td>−4.4 (35.2)</td>
<td>0.24</td>
</tr>
<tr>
<td>LDL-C, mg/dL</td>
<td>102.0 (32.4)</td>
<td>109.0 (34.6)</td>
<td>+7.0 (3.0)</td>
<td>0.02*</td>
</tr>
<tr>
<td>HDL-C, mg/dL</td>
<td>65.3 (20.0)</td>
<td>66.3 (20.5)</td>
<td>+0.99 (12.8)</td>
<td>0.50</td>
</tr>
<tr>
<td>apoB, mg/dL</td>
<td>92.1 (25.1)</td>
<td>95.1 (26.2)</td>
<td>+3.1 (19.3)</td>
<td>0.14</td>
</tr>
<tr>
<td>apoA1, mg/dL</td>
<td>183.8 (45.0)</td>
<td>191.4 (44.0)</td>
<td>+7.6 (35.8)</td>
<td>0.05</td>
</tr>
<tr>
<td>HDL cholesterol efflux capacity</td>
<td>1.05 (0.17)</td>
<td>1.11 (0.16)</td>
<td>+0.06 (0.16)</td>
<td>0.002*</td>
</tr>
</tbody>
</table>

Atherogenic indices

| Total cholesterol/HDL-C | 3.2 (1.2) | 3.0 (1.2) | −0.11 (0.88) | 0.21 |
| apoB/apoA1 | 0.53 (0.19) | 0.52 (0.2) | −0.006 (0.10) | 0.72 |

Median change in CRP was a reduction of 23.5 mg/L (IQR 54.0) between baseline and follow-up. HDL-C indicates high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; apoB, apolipoprotein B; apoA1, apolipoprotein A1; CRP, C-reactive protein.

*Significant, P<0.05
†N=87 for LDL-C, HDL-C, apoB, and apoA1 due to insufficient sample volumes in 3 subjects.
between a reduction in CRP and apoA1 and HDL cholesterol efflux capacity remained \((\text{apoA1}, r=0.27, P=0.009; \text{HDL cholesterol efflux capacity}, r=0.24, P=0.03)\). However, the correlations between reduction in CRP with increases in LDL-C \((r=0.18, P=0.10)\) and HDL-C \((r=0.17, P=0.10)\) had the same trend as the primary analysis but did not reach statistical significance.

We examined the association between change in CRP with the lipid parameters adjusted by age and sex and found that neither age or sex was significant in the models. The reduction in CRP remained significantly associated with all lipid measurements after adjustment for age and sex: LDL-C \((P=0.02)\), HDL-C \((P=0.04)\), apoA1 \((P=0.02)\), and HDL cholesterol efflux capacity \((P=0.03)\).

In a secondary analysis, we investigated the association between reduction in CRP and HDL cholesterol efflux when adjusting for changes in HDL-C and apoA1 levels. The association was attenuated after adjusting for changes in HDL-C levels \((P=0.06)\) and apoA1 levels \((P=0.14)\). While we found no significant correlation between changes in HDL-C levels and HDL cholesterol efflux capacity \((r=0.17, P=0.12)\), we observed a significant correlation between changes in apoA1 levels with improvements in HDL cholesterol efflux capacity \((r=0.33, P=0.002)\).

None of the changes in DAS28-CRP were significantly correlated with lipid parameters; however, the trends were similar to those observed with changes in CRP and lipid parameters (Table 3).

**Discussion**

Among RA subjects who experienced a reduction in CRP, we observed a significant increase in LDL-C levels and concomitant improvement in HDL cholesterol efflux capacity. The magnitude of CRP reduction was significantly correlated with larger reductions in CRP were correlated with larger increases in ApoA1 and larger improvements in HDL cholesterol efflux capacity.

---

**Figure.** Correlations between magnitude of reduction in CRP (natural log transformed) and the percentage change in (A) LDL-C, (B) HDL-C, (C) apoA1, and (D) HDL cholesterol efflux capacity between baseline and 1-year follow-up. apoA1 indicates apolipoprotein A1; CRP, C-reactive protein; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol.
capacity. A trend for correlation also existed between the magnitude of CRP reduction and increases in LDL-C and HDL-C. These findings provide a potential explanation for the apparent paradox observed in the associations between lipid levels and CV risk in RA. Concurrent with increases in LDL-C, were favorable lipid changes that may counterbalance the elevated CV risk associated with higher LDL-C levels. These changes included improvements in the ability of HDL to efflux cholesterol from lipid laden plaques, alongside stable apoB levels, considered a more direct measure of atherogenic non-HDL-C particles than LDL-C.

The relation between a reduction in CRP and changes in lipid parameters was present despite patients receiving various RA treatment modalities, suggesting that the association is not treatment specific. The trend for an inverse relationship between inflammation and HDL-C levels observed in this present study corroborate findings from a cross-sectional study of 11,437 subjects with simultaneous CRP and HDL-C levels from 4 large biochemistry laboratories. In this heterogeneous population, higher levels of CRP, defined as levels >5 mg/L, were associated with lower levels of HDL-C. While we believe our findings support a link between reduction in inflammation and reduction in CV risk, our results cannot account for whether specific RA treatments may have additional beneficial or deleterious influences on CV risk beyond modifying inflammation.

Changes in lipid levels were more closely associated with changes in CRP than the DAS28, a composite measure of RA disease activity that includes subjective (eg, swollen joint count) and objective (eg, patient reported global health) measures. Although the study subjects experienced significant improvements in DAS28, suggesting a response to therapy, we did not observe a significant correlation between DAS28 reduction and changes in the lipid parameters. This lack of correlation was not surprising. The DAS28 is a composite index that includes subjective measures such as the patient’s assessment of arthritis activity over the past week. DAS28 scores can remain high in the absence of inflammation, particularly in subjects with persistent pain secondary to structural joint damage from RA. Thus, CRP and potentially other serum inflammatory markers may be more optimal biomarkers to interpret lipid changes than global RA disease activity indices.

We believe this study provides convincing evidence showing that reduction in inflammation is associated with improvements in HDL cholesterol efflux capacity, consistent with the limited human studies to date. In cross-sectional studies, HDL cholesterol efflux capacity is impaired in RA subjects with high disease activity compared with those with low disease activity, as well as other inflammatory diseases such as systemic lupus erythematosus and psoriasis, compared with controls. Evidence for a longitudinal relationship between inflammation and HDL cholesterol efflux capacity was demonstrated in a study that induced low-grade endotoxemia, a state of heightened inflammation, in 20 healthy adults and found impairment in HDL cholesterol efflux function when comparing HDL function before and after endotoxemia. A study of 15 subjects found that anti-psoriatic treatment was associated with a significant improvement in HDL cholesterol efflux capacity. In addition, a recent study in mice linked myeloperoxidase-mediated oxidation of apoA1 to impairment of reverse cholesterol transport in vivo and the ability of apoA1 to reduce lipid and macrophage content in atherosclerotic lesions. Increased myeloperoxidase activity is associated with higher CRP levels in RA.

In RA, knowledge of the inflammatory state may have additional importance for interpretation of lipid levels and CV risk. An RA patient with a low LDL-C and high CRP may be at elevated CV risk, based on other lipid parameters such as impaired HDL cholesterol efflux function, compared with a patient with a higher LDL-C level but well-controlled disease and normal HDL cholesterol efflux function. Whether the effect of reducing inflammation in the general population will have a significant impact on HDL cholesterol efflux capacity is not clear. The CV risk reduction associated with lowering inflammation as measured by CRP in the general population was demonstrated in several randomized controlled trials by using statin therapy. However, the magnitude of inflammation and, by definition, the amount of reduction that can occur in an individual without an underlying inflammatory disease are smaller compared with an individual with RA. As an example, an individual with high-sensitivity CRP >3 mg/L in the general population is considered to have elevated CV risk, while the mean high-sensitivity CRP in a treated cohort of RA patients is 9.7 mg/L.

Table 3. Correlation Between Reduction in DAS28 and Percentage Change in Lipid and Lipoprotein Parameters

<table>
<thead>
<tr>
<th>Measurements</th>
<th>Correlation</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cholesterol</td>
<td>0.06</td>
<td>0.56</td>
</tr>
<tr>
<td>LDL-C</td>
<td>0.10</td>
<td>0.35</td>
</tr>
<tr>
<td>HDL-C</td>
<td>0.09</td>
<td>0.42</td>
</tr>
<tr>
<td>apoB</td>
<td>0.03</td>
<td>0.76</td>
</tr>
<tr>
<td>apoA1</td>
<td>0.01</td>
<td>0.88</td>
</tr>
<tr>
<td>HDL cholesterol efflux capacity</td>
<td>0.12</td>
<td>0.29</td>
</tr>
</tbody>
</table>

LDL-C indicates low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; apoB, apolipoprotein B; apoA1, apolipoprotein A1; DAS28, Disease Activity Score 28; TC, total cholesterol.
Interestingly, while LDL-C levels increased with reduction in inflammation, the levels of apoB that typically mirror LDL-C levels in the general population remained stable. The mechanism underlying discrepant changes in LDL-C and apoB in this study is unclear. The Friedewald equation used to calculate LDL-C may be inaccurate when triglyceride levels are high, but would not explain the consistent increases in LDL-C observed across different studies associated with RA biologic therapy. DMARDs are not known to be associated with significant increases in triglyceride levels. This finding requires careful follow-up studies using LDL-particle number and size to provide further mechanistic insight.

Concurrent with a reduction in CRP and increases in LDL-C, we observed increases in apoA1 and a significant improvement in HDL cholesterol efflux capacity, pointing toward an improved lipid profile and reduced CV risk. While we did not observe a significant improvement in the atherogenic index, as has been observed in some RA treatment studies, we observed a trend toward reduced atherogenicity of the lipid profile as measured with TC/HDL-C and apoB/apoA1, suggesting larger studies are needed to evaluate this relationship.

We acknowledge limitations to this study. Lipid measurements were not performed in the fasting state. However, there are data to suggest that the variation between fasting and nonfasting states for TC, LDL-C, and HDL-C would not significantly impact interpretation of CV risk. In a population-based study, mean levels varied by <2% for TC and HDL-C and <10% for LDL-C. We could not account for patients who underwent lifestyle modifications that could result in lipid changes between baseline and 1-year follow-up. While we reviewed medical charts for statin use before and during the 1-year follow-up, it is possible that additional patients on statins were missed. Due to the methods used for sample collection and storage in the RA cohort, which were incompatible with NMR analyses, we were unable to further explore whether the discrepancy between the changes in LDL-C and apoB levels were true changes or artifacts in assay sensitivity.

In summary, RA patients experiencing a reduction in inflammation may have an overall improved lipid profile despite increased LDL-C levels. Reduction in CRP was correlated with increases in apoA1 and improvements in the HDL cholesterol efflux capacity. These findings highlight the importance of incorporating levels and changes in inflammation when considering lipid management and CV risk assessment in patients with RA and potentially other inflammatory diseases. Further, these data may provide insight into pathways for elevated CV risk in patients in the general population with higher levels of CRP. Future studies are planned to evaluate lipid composition by using NMR, which can provide information on the size and number of LDL and HDL particles with changes in levels of inflammation. Moreover, studies are needed to investigate the relationship between changes in inflammation and lipid levels and the impact of these changes on CV risk in RA.

Acknowledgments

We acknowledge Gary Bradwin and Dr Nader Rifai, Children’s Hospital Boston, for their valuable suggestions on lipoprotein assays and expertise on methods for measurement.

Sources of Funding

Dr Liao is supported by National Institutes of Health (NIH) grant K08 AR060257 and the Harold and Duval Bowen Fund. Dr Mehta is funded by grant HL-Z-0000 from the Division of Intramural Research at the NIH.

Disclosures

None of the authors have conflicts of interest relevant to the topic of the manuscript. Dr Weinblatt receives research grant funding from Bristol Myers Squibb, UCB, Crescendo Biosience; Dr Shadick receives research grant funding from Crescendo Bioscience, Amgen, UCB, Abbvie, BMS, and Genentech.

References


The Association Between Reduction in Inflammation and Changes in Lipoprotein Levels and HDL Cholesterol Efflux Capacity in Rheumatoid Arthritis
Katherine P. Liao, Martin P. Playford, Michelle Frits, Jonathan S. Coblyn, Christine Iannaccone, Michael E. Weinblatt, Nancy S. Shadick and Nehal N. Mehta

*J Am Heart Assoc.* 2015;4:e001588; originally published January 30, 2015;
doi: 10.1161/JAHA.114.001588

The *Journal of the American Heart Association* is published by the American Heart Association, 7272 Greenville Avenue,
Dallas, TX 75231
Online ISSN: 2047-9980

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://jaha.ahajournals.org/content/4/2/e001588