Circulating Irisin Levels Are Not Affected by Coffee Intake: A Randomized Controlled Trial

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Introduction

Irisin, a novel myokine thought to play an important role in energy expenditure by mediating the exercise-induced browning of fat [1–3], is hypothesized to play an important role in modulating energy expenditure, obesity and metabolism. Coffee consumption also increases energy expenditure and leads to positive metabolic effects, but whether these effects are mediated by irisin remains unknown. The objective of this study was to determine the association between baseline irisin levels and the metabolic profile in humans and to investigate whether consumption of caffeinated coffee alters irisin levels. To this end, a secondary analysis was performed investigating irisin levels at baseline and after eight weeks in 32 healthy, overweight coffee drinkers who were randomized to consumption of 5 cups per day of instant caffeinated coffee, decaffeinated coffee, or water. Spearman correlation and analysis of covariance analyses were performed to identify possible associations. Irisin levels were positively correlated with waist circumference (r = 0.41, p = 0.02), fat mass (r = 0.44, p = 0.01) and CRP (r = 0.47, p = 0.007). Though there was a trend towards increased levels of irisin over time in the caffeinated coffee group (+1.8%) when compared to the placebo group (−4%) this did not reach statistical significance (p = 0.75 for the trend). This first randomized trial failed to reveal any effects of coffee consumption on irisin levels, but a larger trial, appropriately sized on the basis of data provided by this study, is needed to conclusively investigate such a relationship.

Trial Registration: ClinicalTrials.gov NCT00305097
Caffeine increases energy expenditure and positively affects metabolism potentially via interaction with the skeletal muscle and whether irisin is a potential mediator in this process is as yet unknown.

The purpose of this study is twofold: to identify associations between irisin and markers of metabolism in humans and to determine whether coffee consumption affects irisin levels. As caffeine increases energy expenditure and irisin levels appear to rise with increased energy expenditure, we hypothesize that irisin levels will increase with coffee consumption. To this end, we have performed a secondary analysis of the serum levels of irisin in overweight coffee drinkers who were randomly assigned to consumption of caffeinated coffee, decaffeinated coffee, or water for eight weeks. This study aims to shed further light on the possible mechanisms by which irisin and coffee consumption can lead to improved health outcomes.

Materials and Methods

Subjects
Forty-one overweight (BMI 25–35 kg/m²) but otherwise healthy adults who were regular coffee drinkers (≥2 cups/day) were recruited between 2006 and 2008 with inclusion and exclusion criteria that have previously been described [20]. Participants were randomized via the PROC PLAN procedure in the Statistical Analysis System 9.1 (SAS Institute Inc., NC, US) to three different groups (caffeinated coffee, decaffeinated coffee, no coffee) in block sizes of six. Thirty-two subjects were analyzed in this study as the remaining samples were unavailable for assay (Figure 1). The protocol for this trial and supporting CONSORT checklist are available as supporting information (see Checklist S1 and Protocol S1).
and to abstain from caffeine-containing foods throughout the study. Participants returned after eight weeks for adherence questionnaires and repeat blood draws.

**Intervention**

Each day, participants in the two coffee arms of the study consumed five two-gram portions of instant coffee (caffeinated or decaffeinated Nestlé’s Taster’s Choice) prepared with 6 ounces of boiling water. Participants randomized to the no coffee arm drank instead five 6-ounce glasses of water. As was previously reported, the five daily cups of caffeinated coffee provided 345 mg caffeine and 302 mg chlorogenic acid while five cups of decaffeinated coffee provided 216 mg chlorogenic acid [20]. At six weeks, serum caffeine levels were measured in all participants during a non-fasting blood draw to assess for compliance [20].

**Measurements**

At each study visit, weight, height, and waist circumference were measured by a trained investigator according to standard definitions. Body composition was measured using Tanita (model Quantum II, Lean Body software, RJL Systems, Clinton Township, MI, US) single frequency bioelectrical impedance analysis [23,24]. Glucose was measured by glucose hexokinase method using an autoanalyzer in the central clinical laboratory at the Beth Israel Deaconess Medical Center. The homeostasis model assessment for insulin resistance (HOMA-IR) was calculated as [(fasting glucose × fasting insulin)/405] from the OGTT [20]. All samples were stored in a liquid nitrogen freezer at <-130°C until assayed. Samples were assayed in duplicate with all samples for the same participant included in the same batch. The investigator performing the assays was blinded to the intervention group that the samples belonged in. Double-antibody radioimmunoassay (Immulite Chemiluminescence, Siemens Co., New York, NY, US) was used to measure insulin [intra-assay coefficient of variation (CV) 2.1%], C-reactive protein (CRP) (CV 3.6%), and IL-6 (CV 4.7%). Adiponectin (CV 3.9%) and irisin (CV 6.3%) levels were measured by enzyme linked immunosorbent assay kits (Millipore Corporation, Billerica, MA, US, for adiponectin; Phoenix Pharmaceuticals, Inc, Burlingame, CA, US, Cat. No. EK-067-53 for irisin) as previously described [8].

### Statistical Analysis

Comparisons of the baseline characteristics for each group were performed using $\chi^2$ test for categorical variables and analysis of variance for the continuous variables with normal distribution (see Figures S1–S5 to see scatter plots of the variables analyzed). Normality test was done by Shapiro-Wilk test and Kruskal-Wallis test was performed for non-normally distributed variables. Spearman correlation coefficients were calculated to determine associations between changes in irisin and the changes in the biomarkers and anthropometric data (see Tables S1–S4 for complete results).

For the interventional arm of this study, we conducted a secondary analysis of the trial data, comparing the change in irisin from baseline to the 8-week visit between the treatment groups using an analysis of covariance model. The model included the percentage change from baseline as the dependent variable with adjustment for changes in fat mass and CRP over the eight week period. Percentage change from baseline was calculated as the difference between the eight week and baseline irisin levels divided by the baseline level. The model was also examined with 8-week irisin level as the dependent variable adjusted for the baseline irisin level, change in fat mass and change in CRP. Statistical significance was evaluated at an alpha level of 0.05 unless otherwise indicated. All statistical analyses were performed using SPSS Version 21 (SPSS, Inc., Armonk, NY). All data used in these analyses can be made available on request.

**Table 1.** Baseline characteristics of subjects by coffee group.

<table>
<thead>
<tr>
<th></th>
<th>Decaffeinated Coffee (n = 10)</th>
<th>Caffeinated Coffee (n = 12)</th>
<th>Placebo (n = 10)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (% female)</td>
<td>3 (30)</td>
<td>5 (41.7)</td>
<td>5 (50)</td>
<td>0.70</td>
</tr>
<tr>
<td>Age (years)</td>
<td>40.3 (15.9)</td>
<td>36.7 (6.4)</td>
<td>48.3 (16.5)</td>
<td>0.14</td>
</tr>
<tr>
<td>Ethnicity (% non-Hispanic white)</td>
<td>8 (80)</td>
<td>6 (50)</td>
<td>7 (70)</td>
<td>1.00</td>
</tr>
<tr>
<td>BMI</td>
<td>30.1 (1.9)</td>
<td>29.7 (2.0)</td>
<td>30.6 (1.8)</td>
<td>0.56</td>
</tr>
<tr>
<td>Irisin (ng/mL)</td>
<td>55.4 (8.0)</td>
<td>47.8 (5.5)</td>
<td>50.6 (8.7)</td>
<td>0.07</td>
</tr>
<tr>
<td>Hip circumference (cm)</td>
<td>110.2 (4.2)</td>
<td>106.1 (6.4)</td>
<td>108.7 (7.4)</td>
<td>0.29</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>105.5 (6.4)</td>
<td>96.9 (8.1)</td>
<td>101.1 (8.3)</td>
<td>0.05</td>
</tr>
<tr>
<td>Fat mass (kg)</td>
<td>29.5 (6.3)</td>
<td>25.1 (6.5)</td>
<td>29.3 (6.0)</td>
<td>0.19</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>3.3 (1.4–12.7)</td>
<td>1.0 (0.33–1.4)</td>
<td>1.8 (0.91–6.4)</td>
<td>0.04</td>
</tr>
<tr>
<td>IL6 (pg/mL)</td>
<td>2.0 (1.0)</td>
<td>1.1 (0.6)</td>
<td>2.3 (1.4)</td>
<td>0.04</td>
</tr>
<tr>
<td>Adiponectin (µg/mL)</td>
<td>6.6 (3.9)</td>
<td>6.2 (2.7)</td>
<td>7.2 (2.6)</td>
<td>0.75</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td>85.9 (12.9)</td>
<td>87.5 (9.6)</td>
<td>82.3 (11.7)</td>
<td>0.64</td>
</tr>
<tr>
<td>HOMA-IR</td>
<td>2.7 (1.4–4.2)</td>
<td>2.0 (1.1–2.8)</td>
<td>2.1 (1.5–2.4)</td>
<td>0.05</td>
</tr>
</tbody>
</table>

Significant difference between groups at $p<0.05$.

Data displayed as means (standard deviation) for continuous variables and number (percentage) for categorical variables.

CRP and HOMA-IR are shown as median (interquartile range).

Abbreviations: BMI, body mass index; CRP, C-reactive protein; IL-6, interleukin 6; FPG, fasting plasma glucose; HOMA-IR, the homeostasis model assessment for insulin resistance.
Table 2.

Table 2a. Spearman correlation coefficients between irisin levels and baseline anthropometric data

<table>
<thead>
<tr>
<th>Baseline irisin (ng/mL)</th>
<th>% change in irisin</th>
<th>Absolute change in irisin (ng/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>r</td>
<td>p</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>0.25</td>
<td>0.17</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>0.41</td>
<td>0.02</td>
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<tr>
<td>Waist:Hip ratio</td>
<td>0.06</td>
<td>0.75</td>
</tr>
<tr>
<td>Fat mass (kg)</td>
<td>0.44</td>
<td>0.01</td>
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</tbody>
</table>

Table 2b. Spearman correlation coefficients between irisin levels and changes in biomarkers

<table>
<thead>
<tr>
<th>Baseline irisin (ng/mL)</th>
<th>% change in irisin</th>
<th>Absolute change in irisin (ng/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>r</td>
<td>p</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>−0.12</td>
<td>0.51</td>
</tr>
<tr>
<td>IL-6 (pg/mL)</td>
<td>−0.04</td>
<td>0.82</td>
</tr>
<tr>
<td>Adiponectin (µg/mL)</td>
<td>0.12</td>
<td>0.52</td>
</tr>
<tr>
<td>HOMA-IR</td>
<td>0.35</td>
<td>0.06</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td>0.11</td>
<td>0.56</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; CRP, C-reactive protein; IL-6, interleukin 6; HOMA-IR, the homeostasis model assessment for insulin resistance; FPG, fasting plasma glucose.

doi:10.1371/journal.pone.0094463.t002
Results

Baseline Characteristics

The average age (±SD) of these 32 subjects was 41.4 (±13.8) with a BMI of 30.1 (±1.9). Other baseline characteristics of the study population are shown in Table 1. The three groups were not significantly different from each other aside from lower baseline CRP and IL-6 levels in the caffeinated coffee group. Of note, though age and gender distribution were similar in the 32 included and 9 excluded samples, the average BMI of the subjects analyzed was higher than that of the excluded samples (31.1 vs 27.9, p < 0.001).

Correlation between Irisin Levels and Anthropometric Data

Correlations between irisin levels and baseline anthropometric data are shown in Table 2a. Baseline irisin levels were positively and significantly correlated with waist circumference (r = 0.41, p = 0.02) and fat mass (r = 0.44, p = 0.01) and this relationship remained even after adjustment for age and sex (r = 0.38, p = 0.04 for waist circumference and r = 0.43, p = 0.03 for fat mass, see table S1). Though there was a positive association between irisin and BMI (r = 0.25, p = 0.17), this did not reach significance.

Correlation between Irisin Levels and Other Biomarkers of Metabolism

Correlations between irisin levels and changes in other biomarkers over the study period are shown in Table 2b. Change in irisin positively correlated with change in CRP (r = 0.47, p = 0.007) though this was lost after adjustment for age, sex, and BMI (r = 0.31, p = 0.1, see table S4). Though these did not reach significance, irisin exhibited a direct relationship with change in IL6 (r = 0.29, p = 0.1) and a negative relationship with change in adiponectin (r = −0.12, p = 0.53) in the unadjusted model.

Effect of Coffee Consumption on Irisin Levels

Results of the ANCOVA model calculating percentage change in irisin levels over eight weeks by coffee group are graphically depicted in Figure 2. Over eight weeks, irisin levels decreased by 5.5% in the decaffeinated coffee group (95% confidence interval of −20.1 to 9.1) and by 4% in the control group (−18.7 to 10.7) while levels increased by 1.8% in the caffeinated coffee group (−11.9 to 15.5). Though irisin levels appeared to increase with caffeinated coffee consumption, this did not reach statistical significance (p = 0.75). The ANCOVA model using eight-week irisin level as the dependent variable with changes in CRP and fat mass as covariates also found no significant differences between the three groups (p = 0.83).

Discussion

In this study, irisin levels demonstrated positive correlations with markers of adiposity such as fat mass and waist circumference and CRP, a marker of inflammation. While we had hypothesized that irisin levels would rise with increased caffeine consumption, this did not reach statistical significance.

The positive associations between irisin and markers of obesity in this study support the findings of several recent studies and are consistent with our prior hypothesis that irisin levels are increased in obesity as a means of counteracting rising insulin resistance [5,8,9,11,25]. A recent study found that irisin is secreted by adipose tissue in addition to skeletal muscle providing a mechanism that could explain the association between rising BMI and irisin levels [26]. In further support of this theory, though these associations failed to reach significance likely due to our
small sample size, irisin tended to be positively associated with markers of inflammation and HOMA-IR and inversely related to the insulin-sensitizing hormone adiponectin. These findings add to the small but growing body of literature which suggests that irisin is associated with obesity-linked insulin resistance in humans. This continues to be an active area of research and more studies in humans are necessary to better understand the complex role of irisin in human metabolism.

Coffee intake increases energy expenditure and has been associated with a decreased incidence of diabetes and the metabolic syndrome in multiple studies [15,16,20–22]. The mechanism of this effect is not yet clear but could be a result of improved glucose uptake by skeletal muscle or up-regulation of insulin sensitizing hormones such as adiponectin [17,20]. Though irisin is secreted in response to exercise, another state of increased energy expenditure, we did not detect a statistically significant change in irisin levels after eight weeks of coffee consumption [8]. Using these initial data, power calculations were performed using GPower 3.1.6 (Franz Faul, Universita¨t Kiel, Germany) which showed that 53 samples would be required in each group to investigate this relationship, approximately five times the number analyzed in this cohort. Prior to this study there was no data with which to perform such a power analysis. Thus ultimately, a larger scale clinical trial would be needed to conclusively demonstrate whether the trend towards increased irisin levels in those who consume caffeine is indeed significant.

The key limitation of this study is the small and fixed sample size, limiting the power of the study to identify significant relationships and increasing the likelihood of extreme values affecting the results; however, prior to this study no data were available to determine the appropriate sample size. In terms of anthropometric data, the association between irisin and BMI might have been attenuated by the small range of BMI in this study given that all participants were overweight at baseline. The small size and relative homogeneity of this study population also limits generalizability of our findings.

Supporting Information

Figure S1 Scatter plot depicting irisin levels versus BMI and waist circumference. (TIF)

Figure S2 Scatter plot depicting irisin levels versus waist: hip ratio and fat mass. (TIF)

Figure S3 Scatter plot depicting irisin levels versus CRP and adiponectin. (TIF)

Figure S4 Scatter plot depicting irisin levels versus fasting plasma glucose and HOMA-IR. (TIF)

Figure S5 Scatter plot depicting irisin levels versus IL6. (TIF)

Table S1 Spearman correlation coefficients between irisin levels and baseline anthropometric data. (TIF)

Table S2 Spearman correlation coefficients between irisin levels and changes in anthropometric data. (TIF)

Table S3 Spearman correlation coefficients between irisin levels and baseline biomarker levels. (TIF)

Table S4 Spearman correlation coefficients between irisin levels and changes in biomarkers. (TIF)

Checklist S1 CONSORT Checklist. (DOC)

Protocol S1 Study Protocol. (DOC)

Acknowledgments

We would like to thank Dr. Rob M. van Dam for his work in the design and implementation of the original pilot study and for allowing us to use these samples.

Author Contributions

Conceived and designed the experiments: CSM PRP. Performed the experiments: PRP. Analyzed the data: PRP CSM KHP. Contributed reagents/materials/analysis tools: JYH NMW. Wrote the paper: PRP.

References


