Effects of weight loss via high fat vs. low fat alternate day fasting diets on free fatty acid profiles

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Cardiovascular disease risk is associated with excess body weight and elevated plasma free fatty acid (FFA) concentrations. This study examines how an alternate-day fasting (ADF) diet high (HF) or low (LF) in fat affects plasma FFA profiles in the context of weight loss, and changes in body composition and lipid profiles. After a 2-week weight maintenance period, 29 women (BMI 30–39.9 kg/m²) 25–65 years old were randomized to an 8-week ADF-HF (45% fat) diet or an ADF-LF (25% fat) diet with 25% energy intake on fast days and ad libitum intake on feed days. Body weight, BMI and waist circumference were assessed weekly and body composition was measured using dual x-ray absorptiometry (DXA). Total and individual FFA and plasma lipid concentrations were measured before and after weight loss. Body weight, BMI, fat mass, total cholesterol, LDL-C and triglyceride concentrations decreased (P < 0.05) in both groups. Total FFA concentrations also decreased (P < 0.001). In the ADF-LF group, decreases were found in several more FFAs than in the ADF-HF group. In the ADF-HF group, FFA concentrations were positively correlated with waist circumference. Depending on the macronutrient composition of a diet, weight loss with an ADF diet decreases FFA concentrations through potentially different mechanisms.

Results

Subjects. Of the 35 women who started the study, 29 completed the trial. One subject was excluded due to scheduling conflicts and five subjects were excluded due to not having lost any weight during the weight loss period or noncompliance to the diet regime (n = 2 in the ADF-HF group and n = 3 in the ADF-LF group). Baseline characteristics of the ADF-HF and ADF-LF groups are reported in Table 1. The groups were well matched for age, BMI and body composition.
Weight loss, body composition and plasma lipid concentrations. During the initial weight maintenance period (week 1–2), no between-group differences were found in any of the baseline characteristics. Over the 8-week ADF diet period, both the ADF-HF and the ADF-LF groups decreased in weight, BMI, fat mass, and waist circumference (Table 1). There were no differences in the extent of weight loss, changes in BMI, and WC between groups. There was a trend (P = 0.05) in weight, BMI, fat mass, and waist circumference in the ADF-HF group. Significant decreases were observed in LNA (P = 0.01), MA (P = 0.049), AA (P = 0.007), LA (P = 0.004), PA (P = 0.001) and OA (P = 0.001) in the ADF-LF group (Table 2). In contrast, in the ADF-HF group there were significant reductions in FFA concentrations of LNA (P = 0.02) and LA (P = 0.048).

Relationship between changes in body composition and FFA concentrations. Overall, a decrease in weight correlated with a decrease in MA (r = 0.391; P = 0.039). In the ADF-LF group weight loss correlated with decreases in DHA (r = 0.642; P = 0.013). In the ADF-HF group, lower fat mass correlated with decreases in LA (r = 0.635; P = 0.015). Changes in waist circumference in the ADF-HF group correlated with changes in LNA (r = 0.554; P = 0.040), LA (r = 0.700; P = 0.005), PA (r = 0.623; P = 0.017), OA (r = 0.610; P = 0.027), and total FFA (r = 0.623; P = 0.017) (Figure 2). No correlation was found between waist circumference and any FFA in the ADF-LF group.

Relationship between changes in FFA and lipid concentrations. In the ADF-HF group, changes in LNA concentrations negatively correlated with total cholesterol (r = −0.601; P = 0.023) and HDL (r = −0.548; P = 0.042). In the ADF-HF group, DHA negatively correlated (r = −0.573; P = 0.032) with LDL-C. In the ADF-LF group, LA was positively correlated (r = 0.508; P = 0.044) with HDL-C.

*Table 1 | Participant characteristics before (week 3) and after (week 10) the weight loss period*

| Group | ADF-LF (n = 15) | ADF-HF (n = 14) |
|-------|----------------|----------------|
| **Time** | **Age (y)** | **Body weight (kg)** | **Body mass index (kg/m²)** | **% Fat** | **Fat mass (kg)** | **Waist circumference** | **Total cholesterol (mmol/L)** | **LDL cholesterol (mmol/L)** | **Triglycerides (mmol/L)** | **Glucose (mg/dL)** |
| **Week 3** | 43.2 ± 2.3 | 88.0 ± 3.0 | 34.4 ± 0.8 | 41.6 ± 1.1 | 36.9 ± 1.9 | 98.4 ± 1.9 | 5.0 ± 0.2 | 3.0 ± 0.2 | 1.5 ± 0.1 | 1.1 ± 0.1 | 107.4 ± 2.9 |
| **Week 10** | 43.2 ± 2.3 | 83.7 ± 2.7 | 32.7 ± 0.7 | 41.8 ± 1.1 | 35.4 ± 1.9 | 90.4 ± 2.3 | 4.2 ± 0.2 | 2.3 ± 0.2 | 1.5 ± 0.1 | 0.9 ± 0.1 | 101.5 ± 2.0 |
| **Week 3** | 42.4 ± 3.0 | 89.2 ± 2.8 | 34.6 ± 0.7 | 40.4 ± 0.8 | 36.1 ± 1.4 | 98.0 ± 1.9 | 5.0 ± 0.3 | 2.8 ± 0.2 | 1.6 ± 0.1 | 1.4 ± 0.2 | 108.4 ± 3.0 |
| **Week 10** | 42.4 ± 3.0 | 85.2 ± 2.7 | 32.8 ± 0.7 | 39.2 ± 1.0 | 33.2 ± 1.5 | 90.8 ± 1.9 | 4.3 ± 0.2 | 2.3 ± 0.2 | 1.6 ± 0.1 | 1.3 ± 0.2 | 105.7 ± 2.2 |

Values reported as mean ± SEM. Alternate day fasting high fat diet (ADF-HF); alternate day fasting low fat diet (ADF-LF).

*P < 0.05 within group difference (from week 3 to week 10).*

*Figure 1 | Total FFAs in ADF groups from week 3 to week 10. Mean total fatty acid concentrations in ADF-HF and ADF-LF groups for weeks 3 and 10 of the study. Values reported as mean ± SEM. Alternate day fasting high fat diet (ADF-HF); alternate day fasting low fat diet (ADF-LF).*
Table 2 | FFA changes during an ADF-LF and ADF-HF weight loss diet

| FFA (µmol/L) | ADF-LF | ADF-HF |
|-------------|---------|--------|
|              | Week 3  | Week 10| Week 3  | Week 10|
| LNA         | 11.08 ± 0.31 | 10.96 ± 0.22* | 12.07 ± 0.38 | 11.59 ± 0.29a |
| DHA         | 11.61 ± 0.21 | 11.36 ± 0.31 | 12.01 ± 0.25 | 11.59 ± 0.30 |
| MA          | 19.79 ± 0.51 | 19.15 ± 0.46a | 20.72 ± 0.46 | 20.14 ± 0.48 |
| AA          | 11.88 ± 0.38 | 11.34 ± 0.34a | 12.67 ± 0.42 | 12.30 ± 0.28 |
| LA          | 79.92 ± 2.98 | 75.02 ± 2.47a | 89.39 ± 5.42 | 85.10 ± 3.88a |
| PA          | 48.35 ± 5.21 | 39.99 ± 5.85a | 60.75 ± 6.41 | 60.00 ± 5.56 |
| OA          | 37.71 ± 3.95 | 31.47 ± 4.26a | 46.10 ± 4.10 | 45.14 ± 3.60 |
| Total FFA   | 260.89 ± 22.69 | 162.27 ± 24.35b | 264.14 ± 27.89 | 210.21 ± 24.16 |

Values reported as mean ± SEM. Alternate day fasting high fat diet (ADF-HF); alternate day fasting low fat diet (ADF-LF).

Discussion
Elevated total and individual plasma FFA concentrations observed in obesity have been linked with cardiovascular disease risk1,2.13,14. During weight loss, diet composition may affect FFA concentrations7,8. Our study examined the effect of weight loss with a high fat ADF diet on total and individual FFA concentrations compared to a low fat ADF diet. We found that after weight loss, both groups had decreases in total FFA concentrations. However, even though weight loss and regional body composition results were similar between the two groups, almost all FFAs in the ADF-LF group decreased whereas changes were only noted in LA and LNA in the ADF-HF group. Despite similar changes in waist circumference, decreases in WC were correlated with decreases in several individual FFAs in the ADF-HF group only.

In our study, we evaluated circulating concentrations of FFA. Elevated total and individual plasma FFA concentrations observed in obesity have been linked with cardiovascular disease risk and atherosclerosis1,2,14,15. FFAs has been observed to impair vasodilation in blood vessels through activating apoptosis in the endothelium10. Furthermore, FFA is the main energy source for the myocardium; high levels of FFA play a role in damaging the myocardium, exacerbating heart failure and leading to myocardial infarction11. Thus far, no ADF studies have analysed circulating FFA profiles in order to determine whether any cardio-protective changes occur at the FFA level. The results of our FFA analysis show that weight loss with an ADF diet decreases total FFA concentrations, and that both the ADF-LF and ADF-HF diets could be effective at reducing FFA concentrations. Though no significant group*time interactions were observed, the decrease in total FFA concentrations with an ADF-LF diet was almost 2 times that of the ADF-HF diet. Previously, effects of weight loss on FFA concentrations appear to be variable with some studies showing decreases in FFA11,13,14, while others show no change22–24 or even increases25–27 in FFA. Some of these studies achieved weight loss even increases25–27 in FFA. Some of these studies achieved weight loss using a very low calorie diet, surgery or hypocaloric diets with varying macronutrient composition. Therefore, the dietary regimen used in these studies during the weight loss period may be a source of variability in changes of FFA to weight loss.

Individual FFA concentrations have been shown to contribute to CVD risk differently. Thrombosis and arrhythmia have been associated with a number of saturated FFA including palmitic acid28,29. Palmitic, oleic, and myristic acid have been positively associated with lipid risk factors of CVD. Our study also showed that LA, DHA and LNA were associated with lipid risk factors of CVD. Patients post-myocardial infarction were also shown to have greater palmitic and oleic acid and lower linoleic acid compared to age-matched controls30. With an ADF-LF diet, we observed a significant reduction in more types of FFAs (LNA, MA, AA, LA, PA and OA) vs. in the ADF-HF group (LNA and LA). This suggests that the low fat ADF diet was more effective at decreasing FFA concentrations vs. the high fat ADF diet. To our knowledge this is the first study to examine changes in individual FFA composition in the context of a weight loss program using ADF with varying macronutrient composition. Normal weight participants who consumed a high fat diet also observed changes in the proportion of LA and LNA compared with a low fat diet31. These results suggest that LA and LNA are more sensitive to changes with the consumption of a high fat diet. Therefore, combining a high fat diet with weight loss via an ADF protocol would likely have a greater effect on these FFAs, as observed in this study.

There were strong correlations in the ADF-HF group where decreases in waist circumference were associated with decreases in several FFAs. In contrast, there were no such correlations between waist circumference and FFAs in the ADF-LF group. Waist circumference is considered a surrogate marker of abdominal adiposity and visceral fat mass32. We hypothesize that in weight loss, the levels of dietary fat intake and changes in abdominal fat have a collective effect on decreases in FFA concentrations. The relatively larger contribution of a diet low in fat to changes in FFA concentrations may mask the relationship between abdominal fat mass and FFA concentrations. In contrast, weight loss with a high fat diet may have a smaller effect in lowering FFA concentrations in comparison to the greater effect of decreases in abdominal fat. Regardless of the relative contribution of dietary fat content on FFA lowering, changes in abdominal fat mass during weight loss with an ADF diet resulted in cardioprotective changes in FFA concentrations.

A major strength of the study was that both the ADF-LF and ADF-HF group were matched for body composition. Additionally, body composition changed equally in the ADF-LF and ADF-HF groups. Thus, the effects of a HF and LF diet on FFA concentrations could be examined without the confounding effects of changes in body composition. Additionally, since all the meals were provided and the ratio of SFA:MUFA:PUFA was similar in the high and low fat diets, the effects of overall fat content on FFA concentrations could be isolated.

Due to a communication error during sample processing, hormonal regulators of lipolysis were not measured. While insulin inhibits FFA release, other hormones such as epinephrine stimulate FFA release over a wide range33,34. Thus, we are not able to determine the mechanisms that underlie the changes we saw in FFA composition as a result of a ADF-HF or ADF-LF diet. Another limitation of this study was that both intervention groups lost weight during the weight maintenance period. However, there were no between group differences in weight changes. It is conceivable that decreases in weight during the weight maintenance period diluted observable decreases in FFA concentrations during the weight loss that followed.
As we observed significant changes in FFA concentrations during the weight loss period, the effect of the initial weight lost during the weight maintenance period on FFA concentrations was likely to be minimal.

In summary, our findings are the first to demonstrate that a weight loss intervention with ADF-HF or ADF-LF can reduce total and individual plasma FFA concentrations though it appears that a LF diet may be more effective at lowering FFA. Though both diets equally reduced waist circumference, we found several strong correlations between WC and changes in individual FFAs in the ADF-HF group that were not present in the ADF-LF group. Since there were no differences in changes in FFA composition between groups, the correlations between WC and FFAs found only in the ADF-HF group imply that ADF-HF and ADF-LF diets may lower FFA concentrations through different mechanisms. Overall, the decreases in FFA concentrations after ADF weight loss may be cardio-protective. Future studies might include individual FFA profiles when evaluating macronutrient composition, as the effects of specific FFAs in this context are not well known.

Methods

Subjects. Participants were recruited from the Chicago area and were randomized by stratified random sampling based on BMI and age. Subjects were then randomized to a ADF-HF or ADF-LF group, as previously described. Briefly, potential participants were eligible if they were female, 25 to 65 years of age, with a body mass index (BMI) between 30 and 39.9 kg/m². Participants had to be weight stable for 3 months prior to the beginning of the study, sedentary or lightly active, non-smokers, non-diabetic, and not taking any medications. Postmenopausal women were included provided they maintained a stable dose of their current hormone replacement therapy. All volunteers provided written informed consent. All study protocols were carried out in accordance with guidelines approved by the Office for the Protection of Research Subjects at the University of Illinois at Chicago, Chicago, USA and the Human Research Ethics Committee at Concordia University, Montreal, Canada.

Diet. Food provided was prepared in the metabolic kitchen of the Human Nutrition Research Unit at the University of Illinois, Chicago. All subjects underwent a 2-week high fat or low fat weight maintenance period (week 1–2), followed by an 8-week ADF weight-loss diet (week 3–10). The ADF-HF group received 45% fat (14% saturated fat, 20% monounsaturated fat, 11% polyunsaturated fat) and the ADF-LF group received 25% fat (6% saturated fat, 13% monounsaturated, 6% polyunsaturated fat). A detailed description of the diet composition and energy requirements is provided elsewhere. Subjects were provided with 3 meals on each feed day and one meal on each fast day. All subjects were asked to consume their meals at the same time each day.

Weight loss and body composition. Body weight measurements were taken weekly following an overnight fast (to the nearest 0.5 kg) in light clothing and without shoes using a balance beam scale (HealthoMeter; Sunbeam Products, Boca Raton, FL). Height was measured using a wall-mounted stadiometer (to the nearest 0.1 cm). BMI was assessed as kg/m². Total fat mass and fat-free mass was determined by dual energy X-ray absorptiometry (DXA) (QDR 4500 W, Hologic Inc. Arlington, MA) at baseline, week 3 and week 10. Waist circumference was taken by a flexible measuring tape (to the nearest 0.1 cm), midway between the lower costal margin and super iliac crest during a period of expiration. Participants wore a pedometer (Digi-Walker SW-200, Yamax Corp., Japan) during the study to ensure maintenance of regular physical activity habits.
Plasma lipid and free fatty acid analysis. Blood samples were collected in the morning after an overnight fast at baseline, week 3 and week 10. Exercise, alcohol, and coffee were restricted for 24 h before each visit. Blood was centrifuged for 10 min at 320 G in 4°C and plasma was stored at −80°C until analyzed. Plasma lipid samples were analysed as previously described24. FFA were quantified via ultra performance liquid chromatography (UPLC) (Waters ACUITY UPLC, Milford, Massachusetts). n-3-linolenate (LNA), docosahexanoate (DHA), arachadonate (AA), and oleate (OA) standards were purchased from Sigma-Aldrich, Myristate (MA), palmitate (PA) and oleate (OA) standards were purchased from Nu-Chek prep. All standards were >99% pure. FFA standards were dissolved in 4% fatty acid free albumin phosphate buffer. The individual standard solutions were combined to form an external mixed FFA standard. D31 palmitate (Cambridge Isotope Laboratories, Inc., Tewksbury, Massachusetts) was used as an internal standard and added to plasma samples and external standards. Plasma samples and the mixed FFA standards were extracted and derivatized as previously described25. Samples were then transferred to a UPLC vial in 20:80 H2O/acetonitrile solution and 10 μL were injected onto a 55′ Cheated Acquity UPLC BEH C18 1.7 μm 2.1 × 150 mm column eluted with H2O/acetonitrile: 20:80 at a mL/min rate. Peaks were detected with a photodiode array (PDA) detector at 242 nm.

Statistical methods. All data was analyzed using SPSS software (version 20.0, SPSS Inc., Chicago, IL). Results are presented as mean ± SEM. Normality was assessed by Shapiro-Wilk tests. Independent samples t-tests were used to test baseline differences between groups. Repeated-measures ANOVA were used to assess for significance and time as the within-subject factor and diet as the between-subject factor. Post-hoc analyses were done using a Tukey’s test. Pearson correlations were performed to assess the relationships between body composition and FFA concentrations. Differences were considered significant at P < 0.05.

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Author contributions
K.A.V assisted with the design of the experiment and drafted/revised the manuscript. V.T.D. analyzed and interpreted the data, and drafted/revised the manuscript. M.C.K. conceived and designed the experiment, conducted the clinical trial, analyzed the data and was the study manager. M.H. and R.C. generated, collected, analyzed the data and revised the manuscript. C.M.K. assisted with the collection of study data. S.S. obtained funding for the analyses, assisted with the design of the experiment, analyzed the data, drafted/revised the manuscript. All authors read and approved the final manuscript.

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Corrigendum: Effects of weight loss via high fat vs. low fat alternate day fasting diets on free fatty acid profiles

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The Competing financial interests statement in this Article should read:

Krista Varady is an author of the book, “The Every Other Day Diet”, which is a guide to alternate day fasting for the general public.