# **Nutrition & Metabolism**



Research Open Access

# Minimal nutrition intervention with high-protein/low-carbohydrate and low-fat, nutrient-dense food supplement improves body composition and exercise benefits in overweight adults: A randomized controlled trial

Christopher M Lockwood, Jordan R Moon, Sarah E Tobkin, Ashley A Walter, Abbie E Smith, Vincent J Dalbo, Joel T Cramer and Jeffrey R Stout\*

Address: Metabolic and Body Composition Research Laboratory, Department of Health and Exercise Science, University Of Oklahoma, Norman, OK 73019, USA

Email: Christopher M Lockwood - chrislockwood@ou.edu; Jordan R Moon - jordanmoon@ou.edu; Sarah E Tobkin - sarahtobkin@ou.edu; Ashley A Walter - ashannwalter@ou.edu; Abbie E Smith - abbiesmith@ou.edu; Vincent J Dalbo - vinnyjames@ou.edu; Joel T Cramer - jcramer@ou.edu; Jeffrey R Stout\* - jrstout@ou.edu

\* Corresponding author

Published: 21 April 2008

Nutrition & Metabolism 2008. 5:11 doi:10.1186/1743-7075-5-11

Received: 16 November 2007 Accepted: 21 April 2008

This article is available from: http://www.nutritionandmetabolism.com/content/5/1/11

© 2008 Lockwood et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<a href="http://creativecommons.org/licenses/by/2.0">http://creativecommons.org/licenses/by/2.0</a>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

### **Abstract**

**Background:** Exercise and high-protein/reduced-carbohydrate and -fat diets have each been shown separately, or in combination with an energy-restricted diet to improve body composition and health in sedentary, overweight (BMI > 25) adults. The current study, instead, examined the physiological response to 10 weeks of combined aerobic and resistance exercise (EX) versus exercise + minimal nutrition intervention designed to alter the macronutrient profile, in the absence of energy restriction, using a commercially available high-protein/low-carbohydrate and low-fat, nutrient-dense food supplement (EXFS); versus control (CON).

**Methods:** Thirty-eight previously sedentary, overweight subjects (female = 19; male = 19) were randomly assigned to either CON (n = 10), EX (n = 14) or EXFS (n = 14). EX and EXFS participated in supervised resistance and endurance training (2× and 3×/wk, respectively); EXFS consumed 1 shake/d (weeks 1 and 2) and 2 shakes/d (weeks 3–10).

**Results:** EXFS significantly decreased total energy, carbohydrate and fat intake (-14.4%, -27.2% and -26.7%, respectively; p < 0.017), and increased protein and fiber intake (+52.1% and +21.2%, respectively; p < 0.017). EX and EXFS significantly decreased fat mass (-4.6% and -9.3%, respectively; p < 0.017), with a greater (p < 0.05) decrease in EXFS than EX and CON. Muscle mass increase only reached significance in EXFS (+2.3%; p < 0.017), which was greater (p < 0.05) than CON but not EX (+1.1%). Relative VO<sub>2</sub>max improved in both exercise groups (EX = +5.0% and EXFS = +7.9%; p < 0.017); however, only EXFS significantly improved absolute VO<sub>2</sub>max (+6.2%; p = 0.001). Time-to-exhaustion during treadmill testing increased in EX (+9.8%) but was significantly less (p < 0.05) than in EXFS (+21.2%). Total cholesterol and LDL decreased only in the EXFS (-12.0% and -13.3%, respectively; p < 0.017). Total cholesterol-to-HDL ratio, however, decreased significantly (p < 0.017) in both exercise groups.

**Conclusion:** Absent energy restriction or other dietary controls, provision of a high-protein/low-carbohydrate and -fat, nutrient-dense food supplement significantly, 1) modified ad libitum macronutrient and energy intake (behavior effect), 2) improved physiological adaptations to exercise (metabolic advantage), and 3) reduced the variability of individual responses for fat mass, muscle mass and time-to-exhaustion – all three variables improving in 100% of EXFS subjects.

## **Background**

The American College of Sports Medicine (ACSM) Position Stand, "Appropriate intervention strategies for weight loss and prevention of weight regain for adults", states there is little evidence to suggest that exercise-alone is as effective as energy restriction for promoting weight loss [1]. Energy restriction-alone is not, however, a sustainable long-term solution for continued improvements in body composition and health, and instead it has been reported that greater than 50% of subjects that lose weight as a result of dietary restriction-alone eventually regain the weight,[2]. Chronic adjustments to exercise volume and intensity, on the other hand, are virtually limitless. In fact, exercise, in the absence of energy restriction, improves cardiovascular fitness and body composition in a dosedependent fashion; specifically, decreasing body fat and increasing, or at least preventing significant loss of metabolically active lean body tissue [3].

Another strategy that is gaining support for treating overweight and obesity, either when combined with energy restriction or its absence, is the manipulation of macronutrient composition; more precisely, reducing carbohydrate (CHO) and increasing protein (PRO) intake to improve body composition and blood lipids [4-6]. Krieger et al. [4], for example, concluded that, independent of energy intake, low-carbohydrate/high-protein diets elicit a metabolic advantage; significantly reducing body mass (BM), fat mass, percent body fat and retaining significantly more fat-free mass than diets consisting of greater than 42% of energy from CHO and  $\leq 1.05$  g/kg/d PRO, respectively. Similarly, Krauss et al. [7], which was further explored by Feinman and Volek [8], showed that in the absence of energy restriction, a reduction in CHO and concomitant increase in PRO and dietary fat (FAT) resulted in significant improvements in BM, total cholesterol, triglycerides, and total cholesterol-to-high-density lipoprotein ratio compared to a diet consisting of 54% CHO, 16% PRO and 30% FAT. Increased satiety and thermogenesis are also commonly reported in response to a high-protein versus normative-protein (~15% of total energy from PRO) diet [9].

Furthermore, it has been reported that the use of 1–2 macronutrient- and micronutrient-containing meal

replacements, per day, reduces BM and improves a variety of disease risk factors in overweight and obese populations [10]. Such results, however, have been found to be significantly and positively related solely to the hypocaloric state characteristic of said meal replacement interventions [10]. Nevertheless, a major advantage to the use of a portion-controlled, nutrient-dense food supplement appears to be behavioral in nature – their use minimizes drastic behavior modification and simplifies decisionmaking [11], resulting in greater long-term compliance [10]. Thus, the purpose to our minimal nutrition intervention, controlled design was to assess if provision of a high-protein/low-carbohydrate and low-fat (~52% PRO, 33% CHO, 15% FAT) nutrient-dense food supplement (Full Strength®, Phillips Performance Nutrition, LLC, Golden, CO) would, in the absence of energy restriction, alter ad libitum macronutrient intake (behavioral effect) and impose any added metabolic advantage [or detriment | to the physiological effects of 10 weeks of combined aerobic and resistance training in previously sedentary, overweight men and women.

### **Methods**

### Design, subjects and screening

This study involved a minimal nutritional intervention, controlled design to simulate "real world" use of a high-protein/low-carbohydrate and low-fat, nutrient-dense food supplement while participating in a supervised exercise program for 10 weeks. After screening, participants were randomized into one of three groups: exercise (EX), exercise plus food supplement (EXFS), or control (CON). This study was approved by the University of Oklahoma Institutional Review Board for Human Subjects, and written informed consent was obtained from each participant prior to testing.

Sixty sedentary (< 30 min. physical activity per week), overweight (BMI  $\ge$  25) men and women volunteered to participate in this study. Six subjects were lost from each of the exercise groups and ten from the CON. Reasons for attrition included lack of time, unwillingness to consume the food supplement (FS) and conflicts with work. Therefore, 38 adult men and women (Table 1) completed the study. Each participant was assessed by routine medical screening for inclusion. None of the participants reported

Table I: Baseline (PRE) descriptive data of the groups ( $\bar{x} \pm SE$ ).

| Variable         | <b>CON</b> (n = 10; 5F and 5M) | <b>EX</b> (n = 14; 7F and 7M) | <b>EXFS</b> (n = 14; 7F and 7M) |
|------------------|--------------------------------|-------------------------------|---------------------------------|
| Age (yr)         | 30.0 ± 1.6                     | 34.8 ± 1.3                    | 32.6 ± 1.6                      |
| Height (cm)      | 171.7 ± 3.9                    | 175.4 ± 2.3                   | 170.2 ± 2.4                     |
| Body Weight (kg) | 78.3 ± 5.0                     | 82.3 ± 4.1                    | 84.7 ± 5.5                      |
| BMI (kg/m2)      | 26.5 ± 1.4                     | 26.7 ± 1.2                    | 29.2 ± 1.5                      |

CON = control; EX = exercise only; EXFS = exercise + food supplement. No significant differences were observed between groups for height, weight or BMI (p > 0.05).

or exhibited: (a) A history of medical or surgical events that may significantly affect the study outcome, including cardiovascular disease, metabolic, renal, hepatic or musculoskeletal disorders; (b) None of the female participants were currently pregnant or breast feeding; (c) Use of any medicine that may significantly affect the study outcome; (d) Use of nutritional supplements, other than a multivitamin/mineral, in the four weeks prior to the start of the study; and, (d) Participation in another clinical trial or ingestion of another investigational product within 30 days prior to screening.

Prior to the start of the 10-week program (PRE), participants visited the laboratory on two occasions to complete all body composition, cardiorespiratory, strength and blood lipid tests. The same measures were performed during week 12 (POST), after 10 weeks of training. In addition, all participants completed 3-day food logs during PRE and each week of training for a total of 11 weeks. Each food log included two non-consecutive weekdays and one weekend day and was used to represent subjects' average weekly diets. Food logs were analyzed by the same investigator for total energy (kcal), macronutrients and fiber (grams), using Food Processor for Windows, Version 8.6 (ESHA Research, Salem, Oregon). Micronutrient intake was not assessed as part of this investigation.

## Measurements

Height (HT) was measured to the nearest 0.5 cm using a calibrated stadiometer; body mass (BM) was measured using a calibrated clinical scale to nearest 0.01 kg with participants wearing only Spandex shorts or tight-fitting bathing suit. Serum blood samples were drawn at the University of Oklahoma Goddard Health Center. Samples were separated by centrifugation and shipped to Laboratory Corporation of America (Oklahoma City, OK) for analysis. All samples were analyzed using established enzymatic assays for total cholesterol (TC), triglycerides (TRI) and high-density lipoprotein cholesterol (HDL). Low-density and very low-density lipoprotein cholesterol (LDL and VLDL, respectively) were calculated using Friedwald's equations [LDL = TC-TRI/2.2; VLDL = TC-(LDL+HDL)].

# Body composition

All body composition assessments were performed on the same day following a 12-hour fast (*ad libitum* water intake was allowed up to one hour prior to testing). Participants were instructed to avoid exercise for at least 24 hours prior to testing. Fat mass (FM), percent body fat (%FAT) and fat-free mass (FFM) were estimated using the five-compartment (5-C) model described by Wang et al. [12]:

• FM (kg) = 2.748(BV) - 0.715(TBW) + 1.129(Mo) + 1.222(Ms) - 2.051(BM)

- %FAT = (FM/BM) × 100
- FFM = BM FM

Where BV is total body volume, TBW is total body water, Mo is total body bone mineral, Ms is total body soft tissue mineral, and BM is body mass.

The test-retest reliability for the 5-C equation, as measured 24 to 48 hours apart in 11 men and women, resulted in an intraclass correlation (ICC) of 0.99 and a standard error of measurement (SEM) of 0.48%, 0.36 kg and 0.52 kg for %FAT, FM and FFM, respectively [13]. In addition, there were no significant differences (p > 0.05) from trial 1 to trial 2 for %FAT (mean  $\pm$  SE; 22.0  $\pm$  2.5% to 21.1  $\pm$  2.6%), FM (15.6  $\pm$  1.8 kg to 14.9  $\pm$  1.9 kg) and FFM (55.9  $\pm$  3.4 kg to 56.7  $\pm$  3.5 kg).

Dual-energy X-ray absorptiometry (DXA) (software version 10.50.086, Lunar Prodigy Advance, Madison, WI) was used to estimate total body bone mineral content and total body muscle mass (MM). Bone mineral content (BMC) was converted to Mo using the following equation: Mo = total body BMC × 1.0436 [12]. In addition, the sum of lean soft tissue for both arms and legs (ALST), as measured by DXA, was used to estimate MM from the validated equation of Kim et al. [14]: MM = (1.13 × ALST) - (0.02 × age) + [0.61 × sex (m = 0, f = 1)] + 0.97. Test-retest reliability for MM, as measured 24 to 48 hours apart in 11 male and female subjects resulted in an ICC and SEM of 0.99 and 0.04 kg, respectively [13].

Air displacement plethysmography (BOD POD®, Life Measurement, Inc., Concord, CA.) was used to estimate BV. Prior to each test, the BOD POD® was calibrated according to the manufacturer's instructions with the chamber empty and using a cylinder of known volume (49.558 L). The participant, wearing only minimal clothing (as described earlier) and swimming cap, entered and sat in the fiberglass chamber. The BOD POD® was sealed, and the participant breathed normally for 20 seconds while BV was measured. The participant was then connected to a breathing tube internal to the system to measure thoracic gas volume, which was used to correct the BV measurement.

Bioimpedance spectroscopy (BIS) was used to estimate TBW following the procedures recommended by the manufacturer (ImpediMed SFB7, Queensland, Australia). TBW estimates were taken while the participant lay supine on a table with arms  $\geq$  30 degrees from their torso and legs separated. Electrodes were placed at the distal ends of the participants' right hand and foot. Prior to electrode placement, excess body hair was removed, and the skin at

each site was cleaned with alcohol. Prior to analysis, each participant's HT, BW, age and sex were entered into the BIS device. Internal to the device, the BIS utilized 256 frequencies to estimate TBW. The average of two trials within  $\pm$  0.05 liters was used to represent each participant's TBW. The TBW estimate was then used to estimate Ms using the equation from Wang et al. [12]:  $Ms = TBW \times 0.0129$ . The BIS device used in the current study was recently examined in our laboratory as compared to deuterium oxide for estimating TBW in a heterogeneous sample of men and women (n = 30; 23.8 ± 4 yrs; 174.47 ± 7.34 cm; 73.4 ± 18.45 kg; 23.10  $\pm$  5.77 %FAT;  $\bar{x} \pm SD$ ). The results demonstrated a non-significant constant error (CE = -0.56L, p > 0.05) and high correlation (r = 0.97), which is similar to results obtained in other laboratories [15-17]. Therefore, we feel confident that our BIS device accurately estimated TBW.

#### Cardiorespiratory measurements

During a graded exercise test (GXT) on a Quinton® Q65 Series 90 Treadmill (Quinton Instrument Co., Seattle, WA), respiratory gases were monitored and continuously analyzed with open-circuit spirometry to calculate minute ventilation (VE), oxygen consumption rate (VO<sub>2</sub>), carbon dioxide expiration rate (VCO<sub>2</sub>) and respiratory exchange ratio (RER) using a metabolic cart and manufacturer's software (True One 2400®, Parvo-Medics, Inc., Provo, UT). The data were averaged over 30-second intervals. Prior to each test, the metabolic cart was calibrated using room air for the flow rate calibration and gases of known volume and concentration for the calibration of the O<sub>2</sub> and CO<sub>2</sub> analyzers. The highest 30-second VO2 value during the GXT was recorded as the maximal oxygen uptake (VO<sub>2</sub>max) if it coincided with at least two of the following criteria: (a) Plateau in heart rate (HR) or HR values within 10% of the age-predicted HR<sub>max</sub>; (b) Plateau in VO<sub>2</sub> (defined as an increase of not more than 150 ml/min); and/or, (c) RER value greater than 1.15. Test-retest reliability for VO<sub>2</sub>max, as measured using 10 male and female subjects, resulted in an ICC and SEM of 0.98 and 1.17 ml/ kg/min, respectively.

## Strength measurements

Each participant completed a 5-repetition maximum (5RM) for upper- and lower-body strength (bench press and squat, respectively), using a Cybex® Plate Loaded Smith Press (Cybex International, Medway, MA). Each participant completed a familiarization session prior to testing. Testing began with a warm-up consisting of 8 to 10 repetitions at approximately 50% of the tester-estimated 5RM load. Following adequate rest of 2 to 3 minutes, weight was added and participants attempted five repetitions through the full range of motion. After each

successful set of five repetitions, the weight was increased until subjects could no longer complete five repetitions. Participants rested for 2 to 3 minutes between sets. Testretest, for bench press and squat, as measured using 10 male and female subjects, resulted in an ICC and SEM of 0.99, and 0.83 kg and 1.4 kg, respectively.

### Training protocol

The exercise program was designed using the American College of Sports Medicine (ACSM) recommended guidelines for apparently healthy adults; all participants were supervised and trained by an ACSM or National Strength and Conditioning Association certified trainer. Endurance training was performed three days per week. Participants were allowed to select the mode of exercise, provided it utilized large muscle groups and was rhythmic in nature. The progressive endurance training program used is summarized in Table 2. Resistance training was performed two days per week, providing at least 24 hours recovery between exercise sessions. Participants completed 10 exercises that incorporated all major muscle groups. Each exercise was performed once per session, and participants completed 8 to 12 repetitions per exercise until volitional exhaustion. Weight was increased when participants performed 12 repetitions at the same resistance during two consecutive lifting sessions. All lifts were performed on Nautilus Nitro® (Nautilus, Inc., Vancouver, WA) selectorized resistance machines.

## Nutritional protocol

Participants in the EXFS group were instructed to consume one FS per day (Full Strength\*, Phillips Performance Nutrition, LLC, Golden, CO) (Figure 1) for the first two weeks, and two servings per day for the remaining eight weeks. Subjects were provided a two-week supply of FS at the end of testing during PRE and at biweekly weigh-ins throughout the duration of the study. Time of day for consuming the FS, as well as how to incorporate the FS into each subject's diet (i.e. consuming as an additional meal or in place of another meal), was left to the subject's choosing. Participants were instructed to mix the FS in

Table 2: Endurance training protocol

| Week | Duration (min) | % Heart Rate Reserve |
|------|----------------|----------------------|
| I    | 15 – 20        | 40 – 50              |
| 2    | 20 – 25        | 40 – 50              |
| 3    | 25 - 30        | 50 — 60              |
| 4    | 25 - 30        | 50 — 60              |
| 5    | 25 - 30        | 60 – 70              |
| 6    | 25 - 30        | 60 – 70              |
| 7    | 25 - 30        | 60 – 70              |
| 8    | 30 – 35        | 60 – 70              |
| 9    | 30 – 35        | 60 – 70              |
| 10   | 30 – 35        | 60 – 70              |

| Nutriti   | on F  | acte   |  | Nutritio  | n E  | acte   |  |
|---|---|--|--|---|--|--|--|
| Serving Size  | 1 Packet  | t (82g)  |  | Serving Size 1  | Packet   | (85a)  |  |
| Servings Per  | Contain   | er 14  |  | Servings Per (  | Contain  | er 14  |  |
|   |   |  | 12.00  | TO A CONTRACT OF THE PARTY OF   |  | THE REAL PROPERTY AND ADDRESS.   | NAME OF TAXABLE PARTY.   |
| Amount Per Se   | erving  |  |  | Amount Per Sen  |  |  |  |
| Calories 300  |   | Calories from  | NAME AND ADDRESS OF TAXABLE PARTY.   | Calories 300  |  | Calories from  |  |
| Total Fat   | 5g  | % Da   | ily Value*   | Total Fat   | 5g   | % Da   | aily Value*  |
| Saturated Fat   |   |  | 5%   | Saturated Fat   | 1g   |  | 5%   |
| Trans Fat   | 0g  |  |  | Trans Fat   | 0g   |  | 070  |
| Cholesterol   | 15n   | ng   | 5%   | Cholesterol   | 15n  | na   | 5%   |
| Sodium  | 340   | mg   | 14%  | Sodium  | 350  |  | 15%  |
| Potassium   | 920   | mg   | 26%  | Potassium   | 157  | 0mg  | 45%  |
| Total Carbohyo  |   |  | 8%   | Total Carbohydra  | ate 25g  |  | 8%   |
| Dietary Fiber   | 6g  |  | 24%  | Dietary Fiber   | 6g   |  | 24%  |
| Sugars  | 1g  |  | -  | Sugars  | 3g   |  | 1500   |
| Protein   | 40g   |  | 0.1000000  | Protein   | 40g  |  |  |
| Vitamin A   | 100%  | Vitamin C  | 830%   | Vitamin A   | 100%   | Vitamin C  | 830%   |
| Calcium   | 50%   | Iron   | 10%  | Calcium   | 50%  | Iron   | 10%  |
| Vitamin D   | 50%   | Vitamin E  | 100%   | Vitamin D   | 50%  | Vitamin E  | 100%   |
| Vitamin K   | 50%   | Thiamin  | 50%  | Vitamin K   | 50%  | Thiamin  | 50%  |
| Riboflavin  | 50%   | Niacin   | 50%  | Riboflavin  | 50%  | Niacin   | 50%  |
| Vitamin B6  | 50%   | Folate   | 50%  | Vitamin B6  | 50%  | Folate   | 50%  |
| Vitamin B12 Pantothenic Ad  | 50%   | Biotin<br>Phosphorus   | 50%<br>35%   | Vitamin B12 Pantothenic Acid  | 50%  | Biotin   | 50%  |
| lodine  | 50%   | Magnesium  |  | lodine  | 50%  | Phosphorus<br>Magnesium  |  |
| Zinc  | 50%   |  | 100%   | Zinc  | 50%  |  | 100%   |
| Copper  | 25%   | Manganese  | 25%  | Copper  | 25%  | Manganese  | 25%  |
| Chromium  | 50%   | Molybdenum   |  | Chromium  | 50%  | Molybdenum   |  |
|   | nay be higher   | on a 2,000 calorie<br>or lower dependin  | g on   | *Percent Daily Values<br>Your daily values may<br>your calorie needs<br>INGREDIENTS: FS Tru-  | be higher  | or lower depending   | g on   |
| rotein Concentrats colate, L-Glutamin nzyme), FS Energ lour, Fibersol [Sol unflower Oil, FS V hiloride, Magnesiu odium Ascorbate, ocopheryl Acetate lelenium Amino Acid chelate, Molybdent lydrochloride, Ribin homium Picolina otassium lodide, latural & Artificial ellulose Gum, Sal lantarum, Lactoba lantarum, Lactoba | e, Taurine, T<br>y'm (Glucose<br>uble Fiber), I<br>itamin and N<br>im Oxide, Tri<br>Zinc Amino<br>, Niacinamid<br>cid Chelate, c<br>I Chelate, o<br>I Chelate, o<br>offavin, Thian<br>te, Vitamin E<br>Vitamin K, ar<br>Flavors, Carr<br>t, ctobacillus a | yrosine, Aminoge of Polymers, Hydrosey, Hig Onlydextrose), Hig Onlydextrose), Hig Aller Blend (Pot calcium Phospha Acid Chelate, d-Ae, Iron Amino Aci -Calcium Pantoth nganese Amino A cid Chelate, Pyridd Chelate, Pyridd Chelate, Pyridd Oil Gaylocobalami ageenan, Soy Lee ene, Sucralose, Eddobnius, Lacto (Edobnius, Lacto) | n<br>lyzed Oat<br>jh Oleic<br>asssium<br>te,<br>ipha<br>d Chelate,<br>lenate,<br>icid<br>oxine<br>et,<br>in),<br>ithin,<br>Sbacillus | Protein Concentrate, C<br>Isolate, L-Glutamier,<br>Enzyme), FS Energy <sup>M</sup><br>Flour, Fibersol (Solubl<br>Dutch-Processed Coc<br>Vitamin and Mineral B<br>Magnesium Oxide, Tri<br>Ascorbate, Zinc Amino<br>Acetate, Niacinamide,<br>Amino Acid Chelate, d<br>Amino Acid Chelate, d<br>Amino Acid Chelate, M<br>Nolybdenum Amino A<br>Hydrochloride, Ribofl<br>Chromium Picolinate,<br>Potassium Iodide, Vita<br>Natural & Artificial Fla<br>Soy Lecithin, Celluloss<br>Sucralose, FS Probiol<br>Lactobacillus plantaru | Faurine, Ty<br>(Glucose<br>e Fiber], Ioa, High O<br>Jend (Pot.<br>calcium Plo<br>Acid Che<br>Iron Amin<br>-Calcium I<br>Ianganese<br>Icid Chelat<br>avin, Thian<br>Vitamin D<br>amin K, an<br>vors, Hone<br>e Gum, Sal | rrosine, Aminogei Polymers, Hydro Polydextrose), leic Sunflower Oil assium Chloride, hosphate, Sodium late, d-Alpha Too o Acid Chelate, Spantothenate, Coo Amino Acid Chele, Pyridoxine min Hydrochlorid S, Folic Acid, Biot d Cyanocobalami py Powder, Carrag It, Beta Carotene, Lactobacillus acie | n lyzed Oal, FS n copheryl elenium pper ate, e, in, n), geenan, dophilus do |

Figure I
Nutrition Facts and Ingredients of the Food Supplement (Full Strength®, Phillips Performance Nutrition,
Golden, CO). Subjects in EXFS (exercise + food supplement) consumed IFS/d (Week I and 2) and 2FS/d (Week 3–10) in
combination with ad libitum diet. Participants were free to choose either chocolate- or vanilla-flavored FS. Two-week supplies
of FS were provided at bi-weekly weigh-ins.

accordance with the label directions. All participants in CON, EX and EXFS were advised to maintain current (*ad libitum*) diet.

### Statistical analyses

Separate two-way mixed factorial ANOVAs [group (CON vs. EX vs. EXFS) × time (PRE vs. POST)] were used to identify any group x time interactions. If a significant interaction was observed, the statistical model was decomposed by examining the simple main effects with one-way repeated measures ANOVAs for each group and one-way factorial ANOVAs for each time. In the event of significant simple main effects, Tukey post-hoc comparisons were performed among the groups; all pair-wise comparison dependent samples t-tests with Bonferroni corrections (p  $\leq$  0.017) were performed across time. If there was no interaction, main effects were analyzed by collapsing across the non-interacting variable as described above for simple main effects. The level of significance was set at  $p \le 0.05$ . Analyses were performed using SPSS 14.0 (SPSS Inc. Chicago, IL). SPSS-derived p-values of less than  $10^{-3}$  (i.e., "p = 0.000") are reported in the manuscript as the critical pvalue necessary to obtain significance (i.e., " $p \le 0.05$ " or "p≤ 0.017"). Statistical power calculations demonstrated power in this investigation ranged from 0.69 to 0.92.

# Results Nutritional profile

Baseline measures, including estimated dietary intake did not differ (p > 0.05) among subjects in each of the groups (Table 1 and 3, 4, 5, 6, 7). Estimated mean daily energy intake did not differ (p > 0.05) between groups, however,

a significant decrease did occur for total energy within EXFS from PRE to Week 3–10 (-14.4%; p = 0.002); no significant changes appeared within EX or CON. Individual analysis revealed 85.7% (n = 12) of subjects in EXFS reduced average energy intake by -118.62 to -761.12 kcal/ d, with only two subjects reporting increases from PRE to Week 3–10; 21.4% (n = 3) of EX subjects reduced energy intake by -557.87 to -2,184.50 kcal/d (Figure 2). A twoway interaction (p = 0.001; 1 -  $\beta = 0.97$ ) was identified for mean protein intake from PRE to Week 3-10 (Table 3). Post-hoc analysis revealed the change in protein intake for EXFS was significantly greater than EX (+52.1% versus -9.2%, respectively;  $p \le 0.05$ ). Individual analysis revealed 92.9% (n = 13) of subjects in EXFS increased protein intake by +31 to +112.13 g/d, whereas EX values ranged from -68 to +41 g/d (Figure 3). No difference (p > 0.05) was identified for mean total carbohydrate and fiber intake among groups; however, a significant decrease (-27.2%; p = 0.002) in mean daily carbohydrate and an increase (+21.2%;  $p \le 0.017$ ) in mean daily fiber intake did occur within EXFS from PRE to Week 3-10, while no significant changes appeared in EX or CON (Table 3). Individual analysis revealed that 78.6% (n = 11) of subjects in EXFS reduced mean carbohydrate intake by -52.25 to -173.62 g/d (Figure 4). There was no difference (p >0.05) for mean daily fat intake among groups; however, there was a significant decrease in mean daily fat intake within EXFS from PRE to Week 3–10 (-26.7%;  $p \le 0.017$ ).

## Body composition

A significant (p = 0.044; 1 -  $\beta = 0.61$ ) main effect for time was observed across all groups for BM and FFM; however,

Table 3: Average daily energy and macronutrient intake from PRE to Week I-2 and Week 3-10 (  $\bar{x} \pm SE$ ).

|                                  | PRE          | Week I-2               | Week 3-10               |
|----------------------------------|--------------|------------------------|-------------------------|
| Fotal Energy Intake (kcal/d)*    |              |                        |                         |
| CON                              | 2076 ± 168.0 | 1853 ± 167             | 1945 ± 66               |
| EX                               | 2039 ± 196.0 | 2021 ± 127             | 1951 ± 115              |
| EXFS                             | 2166 ± 160.0 | 1865 ± 99 <sup>a</sup> | 1854 ± 98 <sup>a</sup>  |
| Fotal Protein Intake (g/d)**     |              |                        |                         |
| CON                              | 84.1 ± 8.8   | 90.9 ± 8.8             | 90.6 ± 11.6             |
| EX                               | 82.4 ± 5.9   | 75.8 ± 5.2             | 74.8 ± 5.1              |
| EXFS                             | 86.3 ± 8.5   | 102.2 ± 3.3            | $131.3 \pm 4.0^{a,b,c}$ |
| Fotal Carbohydrate Intake (g/d)* |              |                        |                         |
| CON                              | 253.0 ± 23.1 | 205.1 ± 19.0           | 238.0 ± 16.6            |
| EX                               | 268.4 ± 23.1 | 263.1 ± 19.4           | 253.5 ± 20.3            |
| EXFS                             | 261.5 ± 20.9 | $205.0 \pm 19.4^{a}$   | 190.3 ± 10.5a           |
| Total Fat Intake (g/d)*          |              |                        |                         |
| CON                              | 80.9 ± 6.7   | 74.3 ± 7.0             | 70.1 ± 5.5              |
| EX                               | 70.6 ± 10.6  | 73.9 ± 5.7             | 70.7 ± 4.3              |
| EXFS                             | 86.1 ± 7.7   | $70.7 \pm 4.5^{a}$     | $63.1 \pm 4.3^{a,b}$    |
| Γotal Fiber Intake (g/d)**       |              |                        |                         |
| CON                              | 15.7 ± 1.3   | 15.3 ± 1.7             | 16.8 ± 2.1              |
| EX                               | 16.3 ± 1.8   | 16.5 ± 1.6             | 15.5 ± 1.8              |
| EXFS                             | 18.9 ± 2.1   | 19.5 ± 0.8             | $22.9 \pm 0.9$ a,b      |

Representative daily intake was calculated as the mean of values as determined from 3-day food logs, recorded weekly by each participant. CON = control; EX = exercise only; EXFS = exercise + food supplement. One shake per day, was consumed during Week I-2; two shakes, per day, were consumed during Week 3-10. \*Main effect for time ( $p \le 0.05$ ); \*\*Group-by-Time interaction ( $p \le 0.05$ ). \*Different from PRE ( $p \le 0.017$ ); \*Different from Week I-2 ( $p \le 0.05$ ); \*Different from EX ( $p \le 0.05$ ).

Table 4: Changes in body composition from PRE to POST ( $\overline{x} \pm$  SE).

|                     | PRE        | POST           | CHANGE                 |
|---------------------|------------|----------------|------------------------|
|                     |            |                |                        |
| Body Mass (kg)*     |            |                |                        |
| CON                 | 78.2 ± 5.0 | 77.9 ± 5.0     | $-0.3 \pm 0.5$         |
| EX                  | 82.3 ± 4.1 | $82.0 \pm 4.0$ | $-0.3 \pm 0.5$         |
| EXFS                | 84.7 ± 5.5 | 82.9 ± 5.2     | -1.8 ± 1.0             |
| Fat Mass (kg)**     |            |                |                        |
| CON                 | 22.6 ± 2.0 | 22.2 ± 2.0     | $-0.4 \pm 0.4$         |
| EX                  | 23.9 ± 2.2 | 22.8 ± 2.9     | $-1.1 \pm 0.4^{a}$     |
| EXFS                | 28.9 ± 2.3 | 26.2 ± 2.3     | $-2.7 \pm 0.4^{a,b,c}$ |
| % Body Fat**        |            |                |                        |
| CÓN                 | 28.9 ± 2.4 | 28.5 ± 2.6     | $-0.4 \pm 0.4$         |
| EX                  | 29.0 ± 2.2 | $27.8 \pm 2.3$ | $-1.2 \pm 0.4^{a}$     |
| EXFS                | 34.1 ± 1.3 | 31.6 ± 1.4     | $-2.5 \pm 0.4^{a,b,c}$ |
| Fat-Free Mass (kg)* |            |                |                        |
| CON                 | 55.6 ± 4.3 | 55.7 ± 4.4     | $0.10 \pm 0.3$         |
| EX                  | 58.4 ± 3.5 | 59.2 ± 3.7     | $0.80 \pm 0.6$         |
| EXFS                | 55.8 ± 3.5 | 56.7 ± 3.4     | $0.90 \pm 0.5$         |
| Muscle Mass (kg)**  |            |                |                        |
| CON                 | 26.8 ± 2.5 | 26.8 ± 2.6     | 0.0 ± 0.1              |
| EX                  | 28.1 ± 2.2 | 28.4 ± 2.3     | 0.3 ± 0.2              |
| EXFS                | 26.5 ± 2.1 | 27.1 ± 2.1     | $0.6 \pm 0.1^{a,b}$    |

CON = control; EX = exercise only; EXFS = exercise + food supplement. \*Main effect for time ( $p \le 0.05$ ); \*\*Group-by-Time interaction ( $p \le 0.05$ ). \*Different from PRE ( $p \le 0.017$ ); \*Different from CON ( $p \le 0.05$ ); \*Different from EX ( $p \le 0.05$ ).

dependent t-tests showed no significant change for any single group (Table 4). A two-way interaction (p = 0.002;  $1 - \beta = 0.94$ ) was identified for %FAT and FM from PRE to POST; *post-hoc* analyses revealed significantly greater decreases in %FAT and FM for EXFS (-2.5% and -2.7 kg, respectively;  $p \le 0.05$ ) than in EX (%FAT = -1.2%; FM = -1.1 kg) and CON (%FAT = -0.4%; FM = -0.4 kg). In addition, there was a two-way interaction (p = 0.014;  $1 - \beta = 0.77$ ) for MM from PRE to POST. The increase in MM was

Table 5: Changes in upper- and lower-body strength from PRE to POST (  $\overline{x} \pm \text{SE}$  ).

|                    | PRE        | POST           | CHANGE              |
|--------------------|------------|----------------|---------------------|
| Bench Press (kg)** |            |                |                     |
| CON                | 49.9 ± 8.8 | 51.3 ± 9.0     | 1.4 ± 0.4           |
| EX                 | 47.7 ± 7.4 | 54.0 ± 5.7     | $6.3 \pm 0.8^{a,b}$ |
| EXFS               | 50.0 ± 6.4 | 58.5 ± 7.2     | $8.5 \pm 1.0^{a,b}$ |
| Squat (kg)**       |            |                |                     |
| CON                | 60.9 ± 8.4 | 61.9 ± 8.0     | $1.0 \pm 0.6$       |
| EX                 | 59.2 ± 7.2 | $70.8 \pm 7.3$ | 11.6 ± 1.4a,b       |
| EXFS               | 61.0 ± 6.9 | 74.9 ± 7.2     | 13.9 ± 1.8a,b       |

Upper- and lower-body strength, as assessed by five-repetition maximum (5RM) bench press (upper) and squat (lower). CON = control; EX = exercise only; EXFS = exercise + food supplement. \*Main effect for time ( $p \le 0.05$ ); \*\*Group-by-Time interaction ( $p \le 0.05$ ). \*Different from PRE ( $p \le 0.017$ ); \*Different from CON ( $p \le 0.05$ )

significantly (p = 0.05) greater in EXFS (+2.3%) than CON (0.0%) but did not differ significantly from EX (+1.1%); EX was not significantly different (p > 0.05) from CON (Table 4). Individual analysis revealed that 100% (n = 14) of subjects in EXFS reduced FM (range = -0.62 to -5.3 kg) and increased MM (range = +0.19 to +1.67 kg); 85.7% (n = 12) of subjects in EX reduced FM (range = -0.04 to -4.14 kg) and 78.6% increased MM (range = +0.05 to +1.36 kg) (Figures 5 and 6).

### Strength measurements

There was a two-way interaction (p = 0.001;  $1 - \beta = 0.99$ ) for 5RM strength for both the bench press and squat (Table 5). Upper- and lower-body strength significantly improved for EX (+13.2% and +19.6%;  $p \le 0.017$ ) and EXFS (+17.0% and +22.8%;  $p \le 0.017$ ), with no changes

Table 6: Changes in cardiorespiratory fitness from PRE to POST ( $\bar{x} \pm SE$ ).

|  | PRE          | POST           | CHANGE                   |
|--|--------------|----------------|--------------------------|
| VO₂max (ml·kg <sup>-1</sup> ·min <sup>-1</sup> )** |              |                |                          |
| CON  | 36.8 ± 2.1   | $36.9 \pm 2.0$ | $0.1 \pm 0.5$            |
| EX   | 35.7 ± 2.7   | 37.5 ± 2.4     | $1.8 \pm 0.6^{a}$        |
| EXFS   | 32.9 ± 1.9   | 35.5 ± 1.8     | $2.6 \pm 0.4^{a}$        |
| VO <sub>2</sub> (L·min <sup>-1</sup> )*            |              |                |                          |
| CON  | 2.87 ± 0.25  | 2.88 ± 0.25    | 0.01 ± 0.01              |
| EX   | 2.87 ± 0.23  | 3.00 ± 0.82    | $0.13 \pm 0.05$          |
| EXFS   | 2.76 ± 0.20  | 2.93 ± 0.21    | $0.17 \pm 0.04^{a}$      |
| Minute Ventilation (L·min-1)*                      |              |                |                          |
| CON  | 79.4 ± 7.6   | 79.0 ± 6.0     | $0.4 \pm 2.5$            |
| EX   | 83.3 ± 5.0   | 89.1 ± 5.5     | $5.8 \pm 2.4^{a}$        |
| EXFS   | 83.7 ± 4.7   | 88.0 ± 5.1     | $4.3 \pm 1.4^{a}$        |
| Maximum Heart Rate (bpm)*                          |              |                |                          |
| CON  | 192.1 ± 2.7  | 190.9 ± 2.0    | -1.2 ± 1.8               |
| EX   | 190.6 ± 2.6  | 188.6 ± 2.8    | -2.0 ± 1.3               |
| EXFS   | 187.0 ± 6.2  | 184.6 ± 6.0    | -2.4 ± 1.5               |
| Time-to-Exhaustion (sec)**                         |              |                |                          |
| CON  | 643.9 ± 44.8 | 653.0 ± 46.4   | 9.1 ± 13.4               |
| EX   | 681.8 ± 62.0 | 748.6 ± 65.4   | $66.8 \pm 13.8^{a,b}$    |
| EXFS   | 559.9 ± 46.8 | 678.6 ± 50.1   | $118.7 \pm 14.3^{a,b,c}$ |

Cardiorespiratory fitness, as assessed by graded treadmill test to exhaustion. CON = control; EX = exercise only; EXFS = exercise + food supplement. \*Main effect for time ( $p \le 0.05$ ); \*\*Group-by-Time interaction ( $p \le 0.05$ ). a Different from PRE ( $p \le 0.017$ ). b Different from CON ( $p \le 0.05$ ). c Different from EX ( $p \le 0.05$ ).

Table 7: Changes in fasting lipid concentrations from PRE to POST ( $\bar{x} \pm SE$ ).

|                            | PRE             | POST           | CHANGE             |
|----------------------------|-----------------|----------------|--------------------|
| Total Cholesterol (mg/dL)* |                 |                |                    |
| CON                        | 179.8 ± 11.4    | 173.2 ± 10.9   | -6.6 ± 6.6         |
| EX                         | $186.4 \pm 7.3$ | 178.2 ± 7.1    | -8.2 ± 3.4         |
| EXFS                       | 197.5 ± 10.1    | 173.9 ± 13.1   | -23.6 ± 6.7a       |
| HDL (mg/dL)*               |                 |                |                    |
| CON                        | 53.9 ± 3.4      | 51.3 ± 2.4     | -2.6 ± 1.8         |
| EX                         | 45.1 ± 2.2      | 41.9 ± 2.0     | -3.2 ± 1.5         |
| EXFS                       | 51.3 ± 4.1      | 50.9 ± 4.2     | -0.7 ± 1.6         |
| LDL (mg/dL)*               |                 |                |                    |
| CON                        | 107.3 ± 8.7     | 99.8 ± 8.7     | -7.5 ± 4.5         |
| EX                         | 119.7 ± 7.3     | 112.9 ± 5.8    | $-6.8 \pm 2.7$     |
| EXFS                       | 119.5 ± 8.4     | 103.6 ± 10.5   | -15.9 ± 5.2a       |
| TC:HDL*                    |                 |                |                    |
| CON                        | $3.6 \pm 0.3$   | $3.3 \pm 0.3$  | -0.3 ± 0.1         |
| EX                         | $4.6 \pm 1.3$   | 4.2 ± 1.2      | $-0.4 \pm 0.1^{a}$ |
| EXFS                       | 4.2 ± 1.3       | 3.8 ± 1.2      | $-0.4 \pm 0.1^{a}$ |
| VLDL (mg/dL)*              |                 |                |                    |
| CON                        | 21.2 ± 3.8      | 20.0 ± 3.1     | -1.2 ± 1.8         |
| EX                         | 24.7 ± 3.2      | $20.9 \pm 2.9$ | $-3.8 \pm 2.6$     |
| EXFS                       | 25.6 ± 3.9      | $22.3 \pm 3.4$ | -3.3 ± 2.2         |
| Triglycerides (mg/dL)*     |                 |                |                    |
| CON                        | 106.7 ± 18.6    | 98.1 ± 13.9    | -8.6 ± 8.5         |
| EX                         | 124.0 ± 16.1    | 104.1 ± 14.4   | -19.9 ± 12.9       |
| EXFS                       | 138.0 ± 19.0    | 111.2 ± 17.0   | -26.8 ± 10.0       |

CON = control; EX = exercise only; EXFS = exercise + food supplement. HDL = high-density lipoproteins; LDL = low-density lipoproteins; TC:HDL = total cholesterol-to-high-density lipoproteins ratio; VLDL = very low-density lipoproteins. \*Main effect for time ( $p \le 0.05$ ). aDifferent from PRE ( $p \le 0.017$ ).

(p > 0.05) in CON. No difference (p > 0.05) was evident between EX and EXFS.

# Cardiorespiratory measurements

 $VO_2$ max (ml/kg/min) improved ( $p \le 0.017$ ) for EX (+5.0%) and EXFS (+7.9%), however, no changes were

observed for CON (Table 6). When expressed in absolute terms (L/min), VO<sub>2</sub>max increased significantly in EXFS (+6.2%; p = 0.001), whereas EX demonstrated a non-significant (p > 0.017) increase of 4.5%. There was a two-way interaction (p = 0.001;  $1 - \beta = 0.99$ ) for time-to-exhaustion (TTE). *Post-hoc* analysis determined that increases in TTE

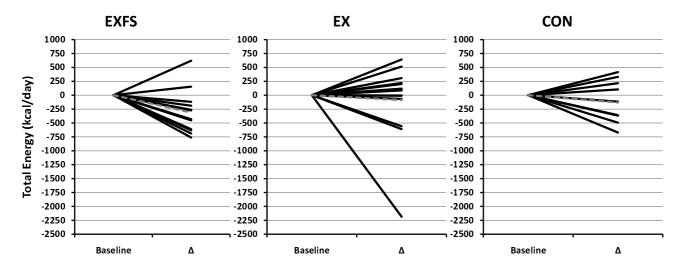


Figure 2 Change in Average Daily Energy Intake by Subject. Individual responses for average daily intake of Total Energy from Baseline to Week 3–10 ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. Total Energy was significantly reduced over time ( $p \le 0.017$ ) in EXFS.

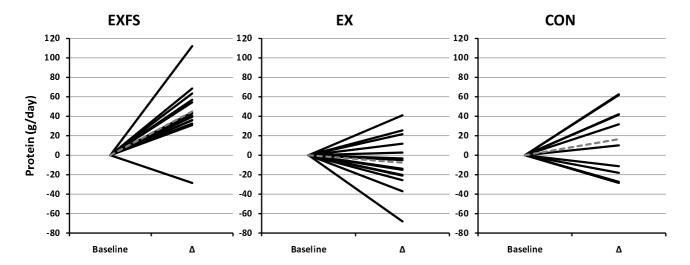


Figure 3 Change in Average Daily Protein Intake by Subject. Individual responses for average daily intake of Protein from Baseline to Week 3–10 ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. Protein intake was significantly increased over time ( $p \le 0.017$ ) in EXFS, and significantly greater than EX ( $p \le 0.05$ ).

observed in the EX (+9.8%) and EXFS (+21.2%) groups were significantly greater ( $p \le 0.05$ ) than changes in CON (+1.4%); the improvement in TTE was significantly greater ( $p \le 0.05$ ) in EXFS than EX. Individual analysis revealed 100% (n = 14) of EXFS improved TTE (+45 to +210 sec), versus 92.9% (n = 13) of EX (-46 to +147 sec) and 40% (n = 4) of CON (Figure 7).

### Blood lipids

Fasting lipid concentrations are presented in Table 7, and individual responses for total cholesterol (TC) and triglycerides (TRI) in Figures 8 and 9. TC and low-density lipoproteins (LDL) decreased significantly in EXFS (-12.0% and -13.3%, respectively;  $p \le 0.017$ ), whereas no significant changes in TC and LDL were observed in EX or CON. The TC-to-high-density lipoprotein ratio (TC:HDL)

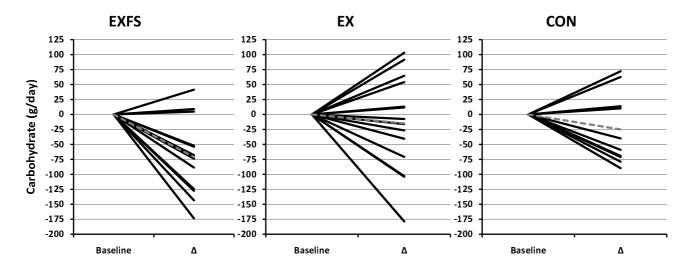


Figure 4 Change in Average Daily Carbohydrate Intake by Subject. Individual responses for average daily intake of Carbohydrate from Baseline to Week 3–10 ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. Carbohydrate intake was significantly reduced over time ( $p \le 0.017$ ) in EXFS.

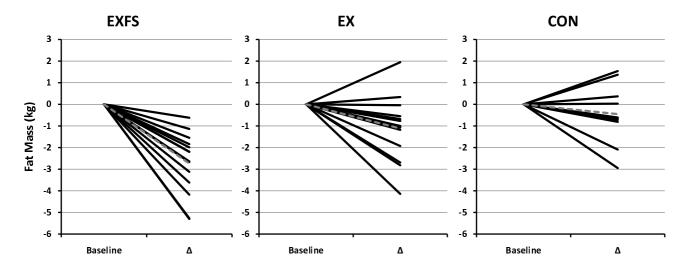


Figure 5 Change in Fat Mass by Subject. Individual responses for changes in Fat Mass from Baseline to Post Week 3 ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. Fat Mass was significantly reduced over time ( $p \le 0.017$ ) in EXFS, and reduced significantly more than EX ( $p \le 0.05$ ).

decreased significantly in both EX and EXFS (-8.7% and -9.5%, respectively;  $p \le 0.017$ ), with no changes in CON.

# **Discussion**

As hypothesized, consumption of the food supplement (FS) significantly increased protein (PRO) and fiber intake in EXFS; PRO rising from 15.9% (86.3 ± 8.5 g/d; ~1.02 g/kg/d) of total energy at baseline (PRE) to 28.3%

(131.3  $\pm$  4.0 g/d; ~1.58 g/kg/d) during weeks 3–10. Simultaneously, subjects in EXFS realized a significant decrease in carbohydrate (CHO), fat (FAT) and total energy. Specifically, CHO decreased from 48.3% (261.5  $\pm$  20.9 g/d) to 41.1% (190.3  $\pm$  10.5 g/d) of total energy; FAT, from 35.8% to 30.6%; and total energy, from 2166 ( $\pm$  160.0) to 1854 ( $\pm$  98) kcals/d [Table 3]. The reduction in total energy intake may be explained by the satiating effect of

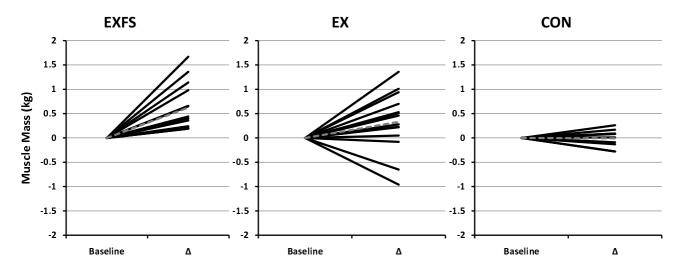


Figure 6 Change in Muscle Mass by Subject. Individual responses for changes in Muscle Mass from Baseline to Post Week ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. Muscle Mass increased significantly over time ( $p \le 0.017$ ) in EXFS, and was significantly greater than CON ( $p \le 0.05$ ).

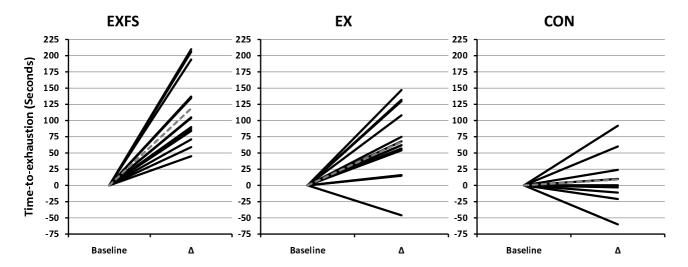


Figure 7 Change in Time-to-Exhaustion by Subject. Individual responses for Time-to-Exhaustion (TTE) from Baseline to Post ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. TTE increased significantly over time (p = 0.001) in EXFS; EXFS was significantly greater than EX and CON ( $p \le 0.05$ ).

PRO and/or fiber [18,19]; though it is our conclusion the modest increase (+4 g/d) in dietary fiber, albeit statistically significant, was not a major contributing factor. Instead, the observed non-significant increase in PRO, during weeks 1–2, and the significant decrease in CHO, FAT and energy intake over that same time period in EXFS would seem to support the separate hypotheses that CHO or FAT reduction spontaneously reduce energy consump-

tion [20-22]. Such a conclusion runs contrary to findings by Weigle et al. [23] that increasing PRO-alone elicits a spontaneous reduction in *ad libitum* energy intake, or that only PRO intake, within EXFS, was significantly different from both control (CON) and EX during weeks 3–10. Instead, it is proposed that both an increase in dietary PRO and a reduction in CHO are equally necessary [24]; however, in accordance with our study design, we hypoth-

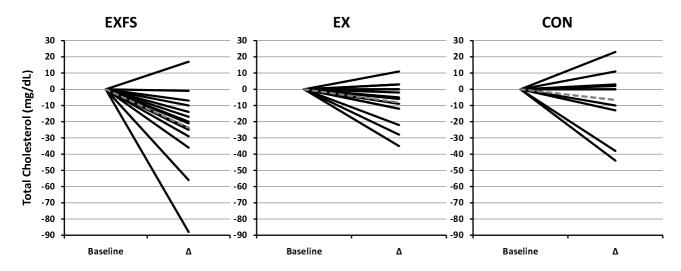


Figure 8 Change in Total Cholesterol by Subject. Individual responses for changes in Total Cholesterol (TC) from Baseline to Post ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. TC was significantly reduced over time ( $p \le 0.017$ ) in EXFS.

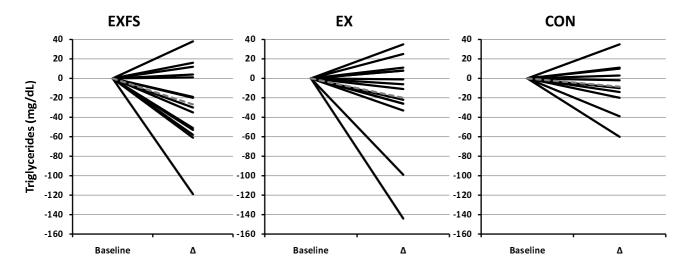


Figure 9 Change in Triglycerides by Subject. Individual responses for changes in Triglycerides (TRI) from Baseline to Post ( $\Delta$ ). Exercise + FS (EXFS; left); Exercise-alone (EX; center); Control (CON; right). Dashed line represents the mean. TRI did not change significantly across treatments ( $p \ge 0.05$ ).

esize that the decrease in CHO was predicated by the initial increase in PRO. Other possible explanations for the reduced energy intake are 1) the high viscosity of the nutritional shake [25], 2) added multivitamin and mineral supplementation [26], 3) reduced portion sizes [27], and/or 4) limiting variety and adding structure to the diet [28].

It has been well documented that, in the absence of dieting, exercise elicits only minor effects on total body mass (BM) despite significant improvements in cardiovascular fitness and strength [29,30]. King et al. ([31] propose that "inter-individual variability," or behavioral and metabolic compensatory events in response to exercise-induced increases in energy expenditure, may largely explain nonsignificant changes in BM from exercise-only interventions. In agreement, we found little individual variability in BM (92.9% of EXFS subjects' ABM occurred within +1.01 and -3.85 kg; 92.9% of EX subjects'  $\Delta BM$  occurred within +2.39 and -2.17 kg). Individual responses for ad libitum energy intake (Figure 2) did, however, provide evidence of what appears to be an apparent trend toward compensatory increase in energy intake in 57.1% (n = 8) of subjects in EX but only14.3% (n = 2) of subjects in EXFS. Consequently, average energy intake was significantly reduced within EXFS (Table 3); supportive of the FS provoking a satiating effect. Additionally, Lofgren et al. [32] state that even modest changes in BM (<5%), in response to reduced energy and CHO intake and increased physical activity, improves cardiovascular health as assessed by low-density lipoprotein (LDL) cholesterol. In agreement, reductions in LDL only reached significance in EXFS ( $\Delta$ BM = -2.13%; p > 0.05); the only group within the current study that realized a significant reduction in energy and CHO intake (Tables 3 and 7).

Expectedly, significant improvements in strength, cardiovascular fitness and blood lipids were observed in both EX and EXFS (Tables 5, 6, 7). Of special note, however, the change in time-to-exhaustion (TTE), in EXFS, was significantly greater than both CON and EX (EXFS = +118.7 sec > EX = +66.8 sec > CON = +9.1 sec; Table 6), with all subjects in EXFS resulting in a minimum improvement in fatigue threshold of +45 sec (Figure 7). Speculatively, the 100% improvement rate in TTE, within EXFS, may be attributable to the rise in muscle mass (MM; +2.3%) and greater reduction in fat mass (FM) [33], stable blood glucose, hormonal or other physiological adaptation [24], specific macro- and/or micro-ingredients of the food supplement or improved recovery nutrition between exercise bouts [34], improved hydration due to twice daily liquid supplementation, or possibly that the EXFS group was not blinded to the intervention. Further research controlling for such variables is warranted.

Interestingly, whereas neither EX or EXFS realized a significant reduction in plasma triglycerides (-16.1% and -19.4%, respectively), only EXFS experienced significant reductions in total cholesterol (-23.6 mg/dL) and LDL (-15.9 mg/dL) (Table 7 and Figures 8 and 9). The non-significant change in triglycerides, within EXFS, is noteworthy because the  $\%\Delta$  (-19.4%) is, in fact, consistent with

findings involving low-carbohydrate and/or energy-restricted diets [6,7,32]. One explanation for the non-significant change may simply be sample size-dependent, whereas it is also plausible that EXFS achieved neither a great enough absolute reduction in CHO and/or energy to elicit such a response [35]. Layman and Walker [36], on the other hand, posit that both CHO must be below 150 g/d and PRO greater than 1.5 g/kg/d to elicit effective treatment against obesity and metabolic syndrome; only the latter was, in fact, achieved in EXFS (CHO = 190.3  $\pm$  10.5 g/d; PRO  $\approx$  1.58 g/kg/d).

According to a recent meta-regression by Krieger et al. [4], the reduction in CHO to <41.4% of total energy and increase in PRO to >1.05 g/kg/d, observed within EXFS, can account for the 1.6 kg, 1.3%, 1.5 kg and 0.3 kg greater improvements in FM, percent body fat, BM and MM respectively, compared to EX [Note: The regression analysis by Krieger et al. stated an additional 0.60 kg of fat-free mass was associated with PRO intakes of >1.05 g/kg/d.]. Of particular value is that supplementation with FS reduced the variability in FM and MM responses to exercise, such that 100% of subjects in EXFS realized a significant improvement; a finding that would seem to support a metabolic advantage of low-carbohydrate/high-protein diet modification [37]. However, changes in CHO and PRO alone cannot, in the current study, be viewed in lieu of modifications in dietary FAT. According to a prediction equation developed by Astrup et al. [22], 1.17 kg of the 1.8 kg of FM lost by EXFS can be accounted for by the 26.71% reduction in dietary FAT. Thus, it seems prudent that future research incorporate isocaloric manipulations of varying macronutrient contributions such that contributing factors and covariates become more evident.

If, instead, we assume 0.45 kg of FM is equivalent to an ~3500 kcal deficit, the -2.7 kg change in FM, within EXFS, could almost completely be accounted for by the -312 kcal/d (-14.4%) reduction in energy intake:

$$\frac{3500kcals}{0.45kgFM} \times \frac{-2.7kgFM}{70days} = -300kcals / d$$

Such assumptions, simplified to "calories in versus calories out," would however fail to recognize the increased energy demands requisite for the +0.6 kg of MM observed within EXFS. Instead, a cumulative metabolic advantage, as postulated by Fine and Feinman [37] and reported by Scott and Devore [38], combined with the anabolic response to increased amino acid availability [24,39], and potentially sustained thyroid hormone levels and reduced insulin response [6] are more probable mechanisms to explain the significant mean changes in EXFS body composition measures. Layman et al. [39], for example, suggests that a hypocaloric diet with carbohydrate-to-protein

ratio (CHO:PRO, in g/d) of 1.5:1.0 or less would be more effective in altering body composition than the 3.5:1.0 ratio currently recommended [39]. Consequently, these authors [39] reported decreases in FM (-22%) and no loss in lean body mass after 16 weeks of an energy-restricted diet composed of a CHO:PRO ratio of ~1.5:1.0, during an exercise program similar to that of the present study. Meckling and Sherfey [40] postulated similar conclusions in response to energy restriction and a 1:1 versus a 3:1 CHO:PRO ratio, with or without exercise, in overweight and obese women. It was reported that both a 1.5:1.0 dietonly and 0.96:1.00 diet + exercise treatment was more effective than traditional high carbohydrate, energy restriction-alone or with exercise. Our findings support these hypotheses and suggest that addition of the FS to the EXFS group's ad libitum diet lowered the CHO:PRO ratio from 3.03:1.00 to 1.46:1.00 for the 10-week training period and, as described by Wood et al. [21], spontaneously reduced ad libitum energy intake via an as of yet fully understood behavioral effect. This change may have accounted for the greater improvements and reduced variability of individual responses for FM and MM in EXFS, when compared to the 3.40:1.00 CHO:PRO ratio of EX. Another plausible explanation, though not directly assessed in the current study, may be that subjects in EXFS consumed one of the FS shakes as a breakfast meal; the addition of the second shake enabling for more frequent, protein-rich meals throughout the day. Such would be supportive of the hypothesis raised by Laymen [24]; that, consuming a minimum of 30 g of PRO for breakfast is "the most critical meal" for supporting an anabolic environment as is consuming PRO every 5-6 hours.

## **Conclusion**

In summary, in the absence of energy restriction or other dietary controls, provision of a commercially available high-protein/low-carbohydrate and fat, nutrient-dense food supplement (Full Strength®; EXFS), consumed daily, during a 10-week combined aerobic and resistance training intervention: 1) elicited a behavioral effect in previously sedentary, overweight adults such that subjects' macronutrient profiles were significantly modified (protein increased; carbohydrate and fat decreased) and total energy intake decreased spontaneously; 2) physiological adaptations to exercise were improved; and lastly, 3) though the significant mean differences between the exercise-only and EXFS groups may not be impressive in the absolute, consumption of the food supplement reduced the variability of individual responses for fat mass, muscle mass and time-to-exhaustion - all three variables improving in 100% of subjects in EXFS. This is of particular interest because virtually all of the available information pertaining to the use of food supplement interventions in overweight and obese populations has involved both an energy-restricted diet and the methodological replacement of at least one traditional meal per day [10]. It is recommended that future research explore the use of the food supplement under various isocaloric and controlled macronutrient ratios, as well as assess the potential impact of the added micronutrients and other ingredients, and whether or not meal timing and frequency are covariates to the improvements noted in body composition, fitness and cardiovascular health.

# **Competing interests**

The authors declare that they have no competing interests.

## **Authors' contributions**

CML: obtained funding, study design, intervention protocols, analysis and interpretation of data, and preparation of manuscript. JRM: study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. SET: study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. AAW: study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. AES: study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. VJD: intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. JTC: study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. JRS: principal researcher, obtained funding, study design, intervention protocols, acquisition of data, analysis and interpretation of data, and preparation of manuscript. All authors read and approved the final manuscript.

# **Acknowledgements**

We would like to thank Shawn Phillips and Phillips Performance Nutrition, LLC (Golden, CO) for funding this study, and all of the men and women who participated.

### References

- Jakicic JM, Clark K, Coleman E, Donnelly JE, Foreyt J, Melanson E, Volek |, Volpe SL: Appropriate strategies for intervention weight loss and prevention of weight regain for adults. Medicine and Science in Sports and Exercise 2001, 33(12):2145-2156.
- Donnelly JE, Smith BK: Is exercise effective for weight loss with ad libitum diet? Energy balance, compensation, and gender differences. Exerc Sport Sci Rev 2005, 33(4):169-174.
- Slentz CA, Duscha BD, Johnson JL, Ketchum K, Aiken LB, Samsa GP, Houmard JA, Bales CW, Kraus WE: **Effects of the amount of** exercise on body weight, body composition, and measures of central obesity: STRRIDE--a randomized controlled study. Arch Intern Med 2004, 164(1):31-39.
- Krieger JW, Sitren HS, Daniels MJ, Langkamp-Henken B: Effects of variation in protein and carbohydrate intake on body mass and composition during energy restriction: a meta-regression 1. The American journal of clinical nutrition 2006, 83(2):260-274.
- Schoeller DA, Buchholz AC: Energetics of obesity and weight control: does diet composition matter? Journal of the American Dietetic Association 2005, 105(5 Suppl 1):S24-8.
- Layman DK, Boileau RA, Erickson DJ, Painter JE, Shiue H, Sather C, Christou DD: A reduced ratio of dietary carbohydrate to protein improves body composition and blood lipid profiles dur-

- ing weight loss in adult women. The Journal of nutrition 2003, 133(2):411-417.
- Krauss RM, Blanche PJ, Rawlings RS, Fernstrom HS, Williams PT: Separate effects of reduced carbohydrate intake and weight loss on atherogenic dyslipidemia. The American journal of clinical nutrition 2006, 83(5):1025-31; quiz 1205.
- Feinman RD, Volek |S: Low carbohydrate diets improve atherogenic dyslipidemia even in the absence of weight loss. Nutr Metab (Lond) 2006, 3:24.
- Halton TL, Hu FB: The effects of high protein diets on thermogenesis, satiety and weight loss: a critical review. Journal of the American College of Nutrition 2004, 23(5):373-385
- Heymsfield SB, van Mierlo CA, van der Knaap HC, Heo M, Frier HI: Weight management using a meal replacement strategy: meta and pooling analysis from six studies. Int J Obes Relat Metab Disord 2003, 27(5):537-549. Levy AS, Heaton AW: Weight Control Practices of United-
- States Adults Trying to Lose Weight. Annals of Internal Medicine 1993, **119(7):**661-666.
- Wang Z, Pi-Sunyer FX, Kotler DP, Wielopolski L, Withers RT, Pierson RN Jr., Heymsfield SB: Multicomponent methods: evaluation of new and traditional soft tissue mineral models by in vivo neutron activation analysis. The American journal of clinical nutrition 2002, 76(5):968-974.
- Weir JP: Quantifying test-retest reliability using the intraclass correlation coefficient and the SEM. J Strength Cond Res 2005,
- Kim J, Wang Z, Heymsfield SB, Baumgartner RN, Gallagher D: Totalbody skeletal muscle mass: estimation by a new dual-energy X-ray absorptiometry method. The American journal of clinical nutrition 2002, 76(2):378-383.
- Matthie J, Zarowitz B, De Lorenzo A, Andreoli A, Katzarski K, Pan G, Withers P: Analytic assessment of the various bioimpedance methods used to estimate body water. | Appl Physiol 1998, 84(5):1801-1816.
- Sardinha LB A.M. Silva, and P.J. Teixeira: Usefulness of ageadjusted equations to estimate body fat with air displacement Plethysmography in male adolescent athletes. Acta Diabetol 2003, 40(1):51-55.
- Van Loan MD, Withers P, Matthie J, Mayclin PL: Use of bioimpedance spectroscopy to determine extracellular fluid, intracellular fluid, total body water, and fat-free mass. Basic Life Sciences 1993, 60:67-70.
- Howarth NC, Saltzman E, Roberts SB: Dietary fiber and weight regulation. Nutrition reviews 2001, 59(5):129-139.
- Astrup A: The satiating power of protein--a key to obesity
- prevention? The American journal of clinical nutrition 2005, **82(1)**:1-2. Dumesnil JG, Turgeon J, Tremblay A, Poirier P, Gilbert M, Gagnon L, St-Pierre S, Garneau C, Lemieux I, Pascot A, Bergeron J, Despres JP: Effect of a low-glycaemic index--low-fat--high protein diet on the atherogenic metabolic risk profile of abdominally obese men. The British journal of nutrition 2001, 86(5):557-568
- Wood RJ, Volek JS, Davis SR, Dell'Ova C, Fernandez ML: Effects of a carbohydrate-restricted diet on emerging plasma markers for cardiovascular disease. Nutr Metab (Lond) 2006, 3:19.
- Astrup A, Ryan L, Grunwald GK, Storgaard M, Saris W, Melanson E, Hill JO: The role of dietary fat in body fatness: evidence from a preliminary meta-analysis of ad libitum low-fat dietary intervention studies. The British journal of nutrition 2000, 83 Suppl
- Weigle DS, Breen PA, Matthys CC, Callahan HS, Meeuws KE, Burden VR, Purnell JQ: A high-protein diet induces sustained reductions in appetite, ad libitum caloric intake, and body weight despite compensatory changes in diurnal plasma leptin and ghrelin concentrations. The American journal of clinical nutrition 2005, **82(1):**41-48.
- Layman DK: Protein quantity and quality at levels above the RDA improves adult weight loss. Journal of the American College of Nutrition 2004, 23(6 Suppl):631S-636S.
- Mattes RD, Rothacker D: Beverage viscosity is inversely related to postprandial hunger in humans. Physiol Behav 2001, 74(4-
- Major GC, Doucet E, Jacqmain M, St-Onge M, Bouchard C, Tremblay A: Multivitamin and dietary supplements, body weight and appetite: results from a cross-sectional and a randomised

- double-blind placebo-controlled study. The British journal of nutrition 2007:1-11.
- Egger G: Are meal replacements an effective clinical tool for weight loss? Med J Aust 2006, 184(2):52-53.
- Ditschuneit HH, Flechtner-Mors M: Value of structured meals for weight management: risk factors and long-term weight maintenance. Obes Res 2001, 9 Suppl 4:284S-289S.
- 29. Utter AC, Nieman DC, Shannonhouse EM, Butterworth DE, Nieman CN: Influence of diet and/or exercise on body composition and cardiorespiratory fitness in obese women. Int J Sport Nutr 1998, 8(3):213-222.
- Garrow JS, Summerbell CD: Meta-analysis: effect of exercise, with or without dieting, on the body composition of overweight subjects. Eur J Clin Nutr 1995, 49(1):1-10.
- King NA, Caudwell P, Hopkins M, Byrne NM, Colley R, Hills AP, Stubbs JR, Blundell JE: Metabolic and behavioral compensatory responses to exercise interventions: barriers to weight loss. Obesity (Silver Spring) 2007, 15(6):1373-1383.
- Lofgren I, Zern T, Herron K, West K, Sharman MJ, Volek JS, Shachter NS, Koo SI, Fernandez ML: Weight loss associated with reduced intake of carbohydrate reduces the atherogenicity of LDL in premenopausal women. Metabolism 2005, 54(9):1133-1141.
- Maughan RJ: The limits of human athletic performance. Ann Transplant 2005, 10(4):52-54.
- Levenhagen DK, Carr C, Carlson MG, Maron DJ, Borel MJ, Flakoll PJ: Postexercise protein intake enhances whole-body and leg protein accretion in humans. Med Sci Sports Exerc 2002, 34(5):828-837.
- Wal JS, McBurney MI, Moellering N, Marth J, Dhurandhar NV: Moderate-carbohydrate low-fat versus low-carbohydrate high-fat meal replacements for weight loss. Int J Food Sci Nutr 2007, 58(4):321-329.
- Layman DK, Walker DA: Potential importance of leucine in treatment of obesity and the metabolic syndrome. The Journal of nutrition 2006, 136(1 Suppl):319S-23S.
- Fine EJ, Feinman RD: Thermodynamics of weight loss diets. Nutr Metab (Lond) 2004, 1(1):15.
- Scott CB, Devore R: Diet-induced thermogenesis: variations among three isocaloric meal-replacement shakes. Nutrition 2005, 21(7-8):874-877.
- Layman DK, Évans E, Baum JI, Seyler J, Erickson DJ, Boileau RA: Dietary protein and exercise have additive effects on body composition during weight loss in adult women. The Journal of nutrition 2005, 135(8):1903-1910.
- Meckling KA, Sherfey R: A randomized trial of a hypocaloric high-protein diet, with and without exercise, on weight loss, fitness, and markers of the Metabolic Syndrome in overweight and obese women. Appl Physiol Nutr Metab 2007, 32(4):743-752.

Publish with **Bio Med Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours you keep the copyright

Submit your manuscript here: http://www.biomedcentral.com/info/publishing\_adv.asp

