Washington University School of Medicine Digital Commons@Becker

Open Access Publications

2017

Effects of two years of calorie restriction on aerobic capacity and muscle strength

Susan B. Racette Washington University School of Medicine in St. Louis

James Rochon Washington University School of Medicine in St. Louis

Mary L. Uhrich Washington University School of Medicine in St. Louis

Dennis T. Villareal Washington University School of Medicine in St. Louis

Sai Krupa Das Washington University School of Medicine in St. Louis

See next page for additional authors

Follow this and additional works at: https://digitalcommons.wustl.edu/open_access_pubs

Recommended Citation

Racette, Susan B.; Rochon, James; Uhrich, Mary L.; Villareal, Dennis T.; Das, Sai Krupa; Fontana, Luigi; Bhapkar, Manjushri; Martin, Corby K.; Redman, Leanne M.; Fuss, Paul J.; Roberts, Susan B.; and Kraus, William E., "Effects of two years of calorie restriction on aerobic capacity and muscle strength." Medicine & Science in Sports & Exercise.49,11.2240-2249. (2017). https://digitalcommons.wustl.edu/open_access_pubs/6686

This Open Access Publication is brought to you for free and open access by Digital Commons@Becker. It has been accepted for inclusion in Open Access Publications by an authorized administrator of Digital Commons@Becker. For more information, please contact engeszer@wustl.edu.

Authors

Susan B. Racette, James Rochon, Mary L. Uhrich, Dennis T. Villareal, Sai Krupa Das, Luigi Fontana, Manjushri Bhapkar, Corby K. Martin, Leanne M. Redman, Paul J. Fuss, Susan B. Roberts, and William E. Kraus

Effects of Two Years of Calorie Restriction on Aerobic Capacity and Muscle Strength

SUSAN B. RACETTE^{1,2}, JAMES ROCHON^{3,4}, MARY L. UHRICH², DENNIS T. VILLAREAL^{2,5}, SAI KRUPA DAS⁶, LUIGI FONTANA^{2,7,8}, MANJUSHRI BHAPKAR³, CORBY K. MARTIN⁹, LEANNE M. REDMAN⁹, PAUL J. FUSS⁶, SUSAN B. ROBERTS⁶, and WILLIAM E. KRAUS¹⁰

¹Program in Physical Therapy, Washington University School of Medicine, St. Louis, MO; ²Department of Medicine, Washington University School of Medicine, St. Louis, MO; ³Duke Clinical Research Institute, Durham, NC; ⁴Rho Federal Systems, Chapel Hill, NC; ⁵Michael E DeBakey VA Medical Center, Baylor College of Medicine, Houston, TX; ⁶Jean Mayer USDA Human Nutrition Research Center on Aging, Tufts University, Boston, MA; ⁷Department of Clinical and Experimental Sciences, Brescia University Medical School, Brescia, Italy; ⁸Healthy Aging Center, CEINGE Biotecnologie Avanzate, Napoli, Italy; ⁹Pennington Biomedical Research Center, Baton Rouge, LA; and ¹⁰Department of Medicine and Duke Molecular Physiology Institute, Duke University School of Medicine, Durham, NC

ABSTRACT

RACETTE, S. B., J. ROCHON, M. L. UHRICH, D. T. VILLAREAL, S. K. DAS, L. FONTANA, M. BHAPKAR, C. K. MARTIN, L. M. REDMAN, P. J. FUSS, S. B. ROBERTS, and W. E. KRAUS. Effects of Two Years of Calorie Restriction on Aerobic Capacity and Muscle Strength. Med. Sci. Sports Exerc., Vol. 49, No. 11, pp. 2240-2249, 2017. Purpose: Calorie restriction (CR) improves health span and delays age-related diseases in many species. The multicenter Comprehensive Assessment of Long-term Effects of Reducing Intake of Energy (CALERIE) study was the first randomized controlled trial of CR in nonobese humans. The aim of this investigation was to determine the effects of CR on VO_{2max} and muscle strength in the CALERIE trial. Methods: Healthy, normal-weight, and mildly overweight women and men (n = 218, mean \pm SE age = 37.9 \pm 0.5 yr) were randomized to 25% CR or an *ad libitum* (AL) control condition in a 2:1 allocation (143 CR, 75 AL). VO_{2max} was determined with an incremental treadmill test; the strength of the knee flexors and extensors was assessed by dynamometry at baseline, 1 yr, and 2 yr. Results: The CR group achieved an average $11.9\% \pm 0.7\%$ CR during the 2-yr intervention. Body weight decreased in CR (-7.7 ± 0.4 kg), but not AL ($+0.2 \pm 0.5$ kg). Absolute \dot{VO}_{2max} (Lmin⁻¹) decreased at 1 and 2 yr with CR, whereas \dot{VO}_{2max} expressed relative to body mass increased at both time points (1 yr: +2.2 ± 0.4; 2 yr: +1.9 \pm 0.4; 0.5 mL·kg⁻¹·min⁻¹) and relative to AL. The CR group increased their treadmill test time and workload at 1 and 2 yr. Strength results in CR were similar, with decreases in absolute flexor and extensor strength, but increases when expressed relative to body mass. No changes were observed for VO_{2max} expressed relative to lean body mass or leg lean mass. Conclusions: Two years of modest CR without a structured exercise component did not appear to compromise aerobic capacity in healthy nonobese adults. The clinical implications of the observed changes in VO_{2max} and muscle strength will be important to explore in future studies. Key Words: VO_{2max}, CR, ENDUR-ANCE, CARDIORESPIRATORY FITNESS

alorie restriction (CR), characterized by a reduction in energy intake without nutrient deficiency, has been shown to extend health span and life span in a variety of organisms, including nonhuman primates (19).

Address for correspondence: Susan B. Racette, Ph.D., Washington University School of Medicine, Campus Box 8502, 4444 Forest Park Ave., St. Louis, MO 63108; E-mail: racettes@wustl.edu. Submitted for publication December 2016. Accepted for publication June 2017.

0195-9131/17/4911-2240/0

MEDICINE & SCIENCE IN SPORTS & EXERCISE_ ${\scriptstyle \circledast}$

Copyright © 2017 The Author(s). Published by Wolters Kluwer Health, Inc. on behalf of the American College of Sports Medicine. This is an open-access article distributed under the terms of the Creative Commons Attribution-Non Commercial-No Derivatives License 4.0 (CCBY-NC-ND), where it is permissible to download and share the work provided it is properly cited. The work cannot be changed in any way or used commercially without permission from the journal.

DOI: 10.1249/MSS.000000000001353

CR is the only known intervention that affects both primary aging (i.e., the inevitable age-related deterioration in tissue structure and function) and secondary aging (i.e., deterioration in tissue structure and function due to adverse lifestyle and environmental factors) (13). A series of observational human studies of adults who have been following a CR diet voluntarily for several years provides intriguing evidence that CR has many health benefits in nonobese humans (7) that are similar to those observed in experimental animal and rodent models. CR is a promising intervention that may enhance cardiovascular health in humans (3).

However, in addition to promoting fat oxidation that causes reductions in adipose tissue with related metabolic health benefits, CR generally induces loss of lean body mass (21,23) because of the catabolism of skeletal muscle and myocardial tissue. There is also some evidence that prolonged CR in humans results in lower physical activity (18), whereas CR in animal models results in preservation of or even increases in physical activity (32). Collectively, these effects of CR on muscle mass and muscle activity in humans may cause adverse changes in cardiorespiratory fitness and muscle strength. On the basis of the well-established association of higher cardiorespiratory fitness and reduced mortality (27), an intervention that adversely affects cardiorespiratory fitness may be cause for concern.

The Comprehensive Assessment of Long-term Effects of Reducing Intake of Energy (CALERIE) study was the first randomized controlled trial of prolonged CR in humans (25). In the CALERIE phase 1 study, we observed that 1 yr of CR in a middle-age (50-60 yr) cohort of nonobese men and women caused reductions in lean body mass, thigh muscle volume, absolute muscle strength, and absolute maximal oxygen uptake ($\dot{V}O_{2max}$) (30). However, when $\dot{V}O_{2max}$ and muscle strength were expressed relative to whole body mass, lean body mass, or muscle volume, the reductions from baseline were no longer significant. Whether the absolute decrements in aerobic capacity and strength in response to CR are clinically important is unclear. Furthermore, it is unknown whether nonobese younger adults would demonstrate similar decrements in muscle mass, strength, and aerobic capacity during long-term CR.

As part of the CALERIE 2 multicenter study, which explored the effects of 2 yr of CR on various metabolic, physiological, and psychological outcomes in healthy young and middle-age adults (22,25), we sought to determine whether long-term CR adversely affects aerobic capacity and muscle strength in normal weight and mildly overweight individuals.

METHODS

Overview. The CALERIE 2 study was a multicenter, randomized controlled trial of 2 yr of CR versus an *ad libitum* (AL) control condition. The design and primary outcomes of the study have been described in detail elsewhere (22,25). The focus of the current analysis is on aerobic capacity and strength measures, which were assessed at baseline, 1 yr, and 2 yr.

Subjects. Participants were enrolled at three clinical CALERIE sites: Pennington Biomedical Research Center, Baton Rouge, LA; Tufts University, Boston, MA; and Washington University School of Medicine, St. Louis, MO. Eligibility criteria included body mass index (BMI) of 22.0 to <28.0 kg·m⁻², age 21-47 yr for women (to avoid menopause) and 21-50 yr for men, nonsmoking, absence of chronic diseases, low risk for eating disorders, and willingness to be randomized to an intervention or control group for 2 yr. An exclusion criterion was engagement in a regular exercise program of vigorous physical activity for ≥ 30 min five or more times per week during the past year. After a series of screening assessments and interviews, eligible participants were randomly assigned to the CR intervention or to an AL control group in a 2:1 ratio, with stratification for site, sex, and BMI (i.e., BMI $<25.0 \text{ vs} \ge 25.0 \text{ kg} \cdot \text{m}^{-2}$). The CR intervention and all study procedures were uniform across the three study sites, and the protocol was approved by the Institutional Review Board at each site. Safety was monitored throughout the trial by a data and safety monitoring board. All participants provided written informed consent.

CR intervention. The CR prescription was set at 25%; the goal was to reduce daily energy intake by 25% relative to daily energy expenditure at baseline and to maintain that intake level for 2 yr. Baseline energy expenditure was determined using the doubly labeled water method during two consecutive 2-wk periods (i.e., 4 wk) of weight stability (22). Thus, the specific calorie prescription was individualized for each participant. The composition of the CR diet was flexible to accommodate individual preferences. During the first 27 d of the 2-yr intervention, participants received three diets (9 d each) of differing macronutrient composition, prepared in the metabolic kitchen at each study site. This strategy promoted early adherence and familiarity with 25% CR using different dietary approaches. Extensive education, counseling, encouragement, and support were provided by registered dietitians, psychologists, and licensed behavior counselors through a structured intervention delivered through a combination of individual and group sessions to facilitate adherence (24). Individual sessions were held weekly (during month 1), twice per month (months 2-12), and monthly (months 13-24); group sessions occurred twice monthly (months 1-6) and monthly (months 7-24). There was no exercise intervention, and no efforts were made to alter participants' physical activity levels (PAL). However, all CALERIE participants (CR and AL) were informed about the Centers for Disease Control and Prevention's physical activity guidelines, which at that time recommended at least 30 min of moderate intensity physical activity on at least 5 d·wk⁻¹.

AL control group. Participants in the AL group were instructed to continue their habitual dietary patterns; they did not receive education or counseling from research personnel beyond instructions needed to complete food diaries and physical activity questionnaires. AL group participants completed the same assessments as participants in the CR group.

Whole body mass, lean body mass, and leg lean mass. Whole body mass was measured to the nearest 0.1 kg on a calibrated scale (Scale Tronix 5200, White Plains, NY) in the research centers in the morning in the fasted state, with the participant wearing a preweighed hospital gown. Lean body mass and leg lean mass were assessed by dual-energy x-ray absorptiometry (DXA; Hologic Inc., Bedford, MA) using a standardized protocol to ensure consistent subject positioning and scan mode over time and across the three clinical sites. Analyses of all DXA scans were performed using Hologic software (Apex version 3.3) at the CALERIE DXA reading center (University of California, San Francisco). The average lean mass of the right and left legs was used in the muscle strength analyses.

Physical activity energy expenditure. Energy expended in physical activity was estimated from total daily energy expenditure (TDEE), resting metabolic rate (RMR), and thermic effect of food (TEF) using the following formula: activity energy expenditure (AEE) = TDEE – RMR – TEF. TDEE was assessed during 4 wk at baseline and for 2 wk at the 1- and 2-yr time points using the doubly labeled water method. RMR was assessed by indirect calorimetry with a calibrated Vista-MX metabolic cart (Vacumed, Ventura, CA). TEF was estimated to be 10% of daily energy intake. At baseline, energy intake was assumed to equal TDEE because participants were weight stable; at the 1- and 2-yr time points, energy intake was computed from TDEE and changes in body energy stores based on DXA (20). PAL was computed as TDEE/RMR and used as a relative estimate of physical activity (16) as subjects lost weight.

Dietary protein intake. Daily protein intake was quantified from 6-d food diaries that were analyzed using the University of Minnesota Nutrition Data System for Research dietary analysis program at the CALERIE nutrition reading center (University of Cincinnati). Protein intake was expressed in absolute $(g \cdot d^{-1})$ and relative (g·kg body mass⁻¹·d⁻¹) quantities.

Aerobic capacity and physiologic responses to maximal treadmill exercise. Maximal oxygen uptake (VO_{2max}) was measured using the Cornell incremental treadmill test (28), with the speed and/or grade of the treadmill changing every 2 min (stage 1, 1.7 mph/5% grade; stage 2, 1.7 mph/10%; stage 3, 2.0 mph/10%; stage 4, 2.5 mph/12%; stage 5, 3.0 mph/13%; stage 6, 3.4 mph/14%; stage 7, 3.8 mph/ 16%; stage 8, 4.2 mph/16%; stage 9, 4.8 mph/17%; stage 10, 5.0 mph/17%; stage 11, 5.0 mph/18%; stage 12, 5.5 mph/ 18%). Expired gases were collected continuously, and ventilatory variables were calculated at 15-s intervals using a calibrated metabolic cart (TrueOne® 2400 or TrueMax 2400; Parvo Medics, Sandy, UT). The flowmeter was calibrated with a 3-L syringe, and gas calibration was performed with precision-analyzed gas mixtures before and after each test. Participants were instructed not to take medications the morning of the assessment and not to eat or drink beverages other than water for at least 90 min before their treadmill test. Before the baseline test, participants practiced walking on the treadmill without holding onto the handrails. Before each test, participants were familiarized with the testing protocol, the Borg RPE scale (4), and hand signals. HR was monitored continuously throughout the test with 12-lead ECG. Blood pressure (BP) was measured by auscultation at every other workload and at the end of the test. RPE was recorded during the last 30 s of each workload. Research personnel provided verbal encouragement and support to assist participants in reaching a maximal effort.

The two highest consecutive 15-s $\dot{V}O_2$ values during the last 2 min of exercise were averaged to determine $\dot{V}O_{2max}$ if the following three criteria were met: a plateau in $\dot{V}O_2$, defined as an increase of ≤ 150 mL between the final two workloads (i.e., stages) of the test, RER ≥ 1.10 , and HR_{max} within 10 bpm of age-predicted HR_{max}. Age-predicted HR_{max} was computed as 208 - (0.7 × age) (29). Oxygen pulse at maximal exercise was calculated as absolute $\dot{V}O_{2max}$ divided by HR_{max}. Rate pressure product at maximal exercise was computed as the product of HR_{max} and maximal systolic BP.

Muscle strength and muscle endurance. Muscle strength and endurance were assessed for knee extensors and

knee flexors using Biodex System 3 dynamometers (Biodex Medical Systems, Shirley, NY) at all three sites; a Cybex II Isokinetic Dynamometer (Cybex Division of Lumex, Inc., Ronkonkoma, NY) was used with some participants at the Pennington Biomedical Research Center. The test sequence and procedures were standardized and programmed into the dynamometers. Personnel at all sites followed the same script for explaining the assessments to participants. The \dot{VO}_{2max} and strength tests were performed 1 wk apart at approximately the same time of day at all assessment time points.

The testing series began with a 10-min warm-up on a cycle ergometer (50 W, 70 rpm). The participant was then positioned properly in the dynamometer chair, and the right leg was weighed at 30°. All results were adjusted (by the dynamometer computer) for gravity effect by subtracting the lower leg mass from flexion torque values and adding lower leg mass to extension torque values. Subjects were familiarized with the testing procedures by performing three submaximal repetitions (~75% effort) at each angular velocity. The isokinetic strength tests included five repetitions of concentric knee extension and concentric knee flexion at $60^{\circ} \cdot s^{-1}$, then at $180^{\circ} \cdot s^{-1}$, with a 30-s rest between sets. After a 5-min rest period, muscular endurance was assessed using 30 repetitions of concentric knee extension and flexion at $180^{\circ} \cdot s^{-1}$. Participants were instructed to contract maximally with each repetition rather than to pace themselves. After another 5-min rest period, three 5-s maximal isometric contractions, with a 30-s rest between, were performed at 45° for knee extension and flexion. Within a set, all repetitions were performed continuously, without rest between repetitions. The entire sequence was repeated with the left leg.

Muscular strength was interpreted as peak torque from the isokinetic and isometric tests. Results of the right and left legs were averaged and expressed in absolute terms, relative to whole body mass, and relative to leg lean mass. Muscular endurance was determined from muscle fatigue index, which was computed for both knee extension and knee flexion as the percent decrease in peak torque during the 30-repetition isokinetic sets at 180° ·s⁻¹ using the following equation:

muscle fatigue index = $100 - \frac{\text{mean of last 5 repetitions}}{\text{mean of highest consecutive 5 repetitions}} \times 100$

Results from the right and left legs were averaged. A composite muscle fatigue index was computed for knee extension and flexion by averaging those results.

Statistical analyses. Standard intention-to-treat analyses were performed by including all available observations for each time point. Baseline characteristics were compared using the Wilcoxon–Mann–Whitney test for continuous and ordinal variables. Typical for outcomes dependent on maximal exertion, an initial examination of outcome variables revealed numerous outliers. Because of concerns that they could exert undue influence on the results, robust regression procedures (15) were used to evaluate responses to CR. Specifically, *M*-estimation (14) using the bisquare weight function

as implemented in PROC ROBUSTREG in SAS was applied. The advantage is that this method is nearly fully efficient when the data are normally distributed yet provides resistant and stable inference in the presence of outliers. The dependent variable was the change from baseline to 1 yr or 2 yr, with treatment assignment as the between-group factor. Design variables (i.e., site, sex, and BMI stratum) and the baseline value centered about the mean of the outcome were included as covariates. The predicted mean and SE are the adjusted values from this model. Within- and between-group changes from baseline were tested by defining contrasts among the regression parameters. Type I error was controlled at $\alpha = 0.05$ using the sequentially rejective Bonferroni procedure (12); P values were adjusted using the method of Wright (31). Consistency between the two sexes was evaluated by adding and testing the treatment-sex interaction term. Spearman correlations were used to determine univariate associations. Analyses were performed using SAS Software version 9.2 and higher (SAS Institute, Cary, NC). Data are expressed as means \pm SE.

RESULTS

Subjects. As reported in detail previously (22), 218 subjects began the study (143 CR, 75 AL); 69.7% were women. The racial distribution was 77.1% White, 11.9% Black, and 11.0% other; 3.2% were Hispanic. Mean age was 37.9 ± 0.5 yr. Attrition during the 2-yr study resulted in an analysis sample of 201 subjects (130 CR, 71 AL) at 1 yr and 188 subjects (117 CR, 71 AL) at 2 yr. The CR group achieved an average 11.9% \pm 0.7% CR during the 2-yr intervention, which was significantly

lower than the prescribed 25% CR. No sex differences were observed in the average %CR achieved during year 1 (females, $15.1\% \pm 0.8\%$; males, $15.3\% \pm 1.2\%$; P = 0.74) or throughout the 2-yr intervention (females, $11.9\% \pm 0.8\%$; males, $11.7\% \pm 1.2\%$; P = 0.85).

Whole body mass, lean body mass, and leg lean mass. At baseline, 48% of subjects were categorized as normal weight (i.e., BMI, 22.0 to <25.0 kg·m⁻²) and 52% were overweight (i.e., BMI, 25.0 to <28.0 kg·m⁻²). As shown in Table 1, whole body mass decreased significantly in the CR group, but not in the AL group, at 1 and 2 yr, with a significant difference between groups at both time points. Likewise, lean body mass and leg lean mass decreased in response to the CR intervention. There was a significant correlation between change in whole body mass and change in lean body mass in the CR group at 2 yr (r = 0.649, P < 0.0001). Comprehensive analyses of body composition changes by treatment and sex are provided by Das et al. (6).

Physical AEE. Absolute AEE (kcal·d⁻¹) decreased significantly in the CR group at 1 and 2 yr (Table 1). In the CR group, this tended to differ by sex (P = 0.055), with decreases observed in females (-129 ± 23 and -142 ± 26 kcal·d⁻¹ at 1 and 2 yr, respectively, both P < 0.0001), but not in males (P = 0.701). When physical activity was expressed in relative terms as PAL, no significant changes were observed in the CR or AL group.

Dietary protein intake. Absolute protein intake $(g \cdot d^{-1})$ did not differ between groups (Table 1). Relative protein intake exceeded the recommended dietary allowance of 0.8 g·kg body mass⁻¹·d⁻¹ (16) throughout the intervention. The change in lean body mass in the CR group from baseline to 2 yr was

TABLE 1. Body mass, physical activity, and dietary protein in the CR and AL groups.

	CR			AL	
	$\textbf{Mean} \pm \textbf{SE}$	Within-Group P Value	$\text{Mean} \pm \text{SE}$	Within-Group P Value	Between-Group P Value
Body mass (kg)					
Baseline	72.0 ± 0.8		71.5 ± 1.1		0.978
Change at 1 yr	-8.5 ± 0.3	<0.0001	-0.4 ± 0.4	0.529	<0.0001
Change at 2 yr	-7.7 ± 0.4	<0.0001	$+0.2\pm0.5$	0.688	<0.0001
Lean body mass (kg)					
Baseline	46.0 ± 0.8		45.2 ± 1.0		0.479
Change at 1 yr	-2.2 ± 0.1	<0.0001	-0.2 ± 0.2	0.439	<0.0001
Change at 2 yr	-2.3 ± 0.2	<0.0001	-0.1 ± 0.2	0.533	<0.0001
Leg lean mass (kg)					
Baseline	7.7 ± 0.1		7.5 ± 0.2		0.302
Change at 1 yr	-0.36 ± 0.03	<0.0001	+0.01 \pm 0.03	0.813	<0.0001
Change at 2 yr	-0.41 ± 0.03	<0.0001	-0.01 ± 0.04	0.813	<0.0001
Physical AEE (kcal·d ⁻¹)					
Baseline	808 ± 20		758 ± 27		0.183
Change at 1 yr	-64 ± 21	0.010	-8 ± 27	0.876	0.244
Change at 2 yr	-84 ± 23	0.002	-5 ± 29	0.876	0.105
PAL					
Baseline	1.75 ± 0.02		1.72 ± 0.02		0.363
Change at 1 yr	-0.03 ± 0.02	0.686	-0.01 ± 0.03	0.686	0.686
Change at 2 yr	-0.04 ± 0.02	0.349	-0.01 ± 0.03	0.686	0.686
Protein intake, absolute (g·d ⁻¹)					
Baseline	87 ± 2		87 ± 3		0.973
Change at 1 yr	-1 ± 2	0.916	-1 ± 2	0.916	0.916
Change at 2 yr	-4 ± 2	0.073	-7 ± 2	0.017	0.916
Protein intake, relative (g·kg body mass ⁻¹ ·d ⁻¹)					
Baseline	1.20 ± 0.02		1.21 ± 0.04		0.951
Change at 1 yr	$+0.12\pm0.03$	0.0001	-0.01 ± 0.04	0.856	0.012
Change at 2 yr	$+0.06 \pm 0.03$	0.062	-0.12 ± 0.04	0.004	<0.001

Sample sizes at baseline, 1 yr, and 2 yr were 143, 130, and 120 for CR and 75, 71, and 71 for AL, respectively.

not related to absolute protein intake during the intervention (r = -0.062, P = 0.502).

Aerobic capacity and physiologic responses to maximal treadmill exercise. Table 2 presents aerobic capacity results for participants who achieved $\dot{V}O_{2max}$: n =195 at baseline (130 CR, 65 AL); *n* = 158 at 1 yr (101 CR, 57 AL); n = 141 at 2 yr (92 Cr, 49 AL). Treadmill exercise time increased significantly in CR at 1 and 2 yr (baseline 13.0 ± 0.2 ; 1 yr 14.3 ± 0.3 ; 2 yr 14.6 ± 0.3 min); the increases were significant relative to AL at both time points (P < 0.001). Absolute \dot{VO}_{2max} (L·min⁻¹), however, decreased significantly in the CR group at 1 and 2 yr; a small nonsignificant decrement was observed in AL at 2 yr. By contrast, VO_{2max} expressed relative to body mass increased significantly in CR subjects at both time points relative to baseline and relative to AL. When VO_{2max} was expressed relative to lean body mass or lean leg mass, no changes were observed from baseline to 1 yr or 2 yr in the CR or AL group.

When the analyses were performed using absolute and relative $\dot{V}O_{2peak}$ (i.e., the single highest 15-s $\dot{V}O_2$ value)

instead of $\dot{V}O_{2max}$ and included all participants who completed a treadmill test, the overall results and conclusions were not different. Furthermore, changes in absolute and relative $\dot{V}O_{2max}$ in response to 2 yr of CR did not differ between participants who were normal weight or overweight at baseline. Data on oxygen pulse, rate pressure product, HR, BP, RER, and RPE at maximal exercise are presented in Table 2.

Sex differences in body mass and aerobic capacity in the CR group are shown in Table 3. Men had higher body mass, lean body mass, absolute $\dot{V}O_{2max}$, and relative $\dot{V}O_{2max}$ values than women at all time points (as expected). Absolute $\dot{V}O_{2max}$ and $\dot{V}O_{2max}$ expressed relative to lean body mass decreased significantly only among females in response to CR. $\dot{V}O_{2max}$ expressed relative to whole body mass tended to increase more among CR males than females. Treadmill test time increased significantly among females and males at both time points; a statistically greater increase was observerd among males after 2 yr of CR. Changes in other exercise test variables in response to CR, such as oxygen pulse at maximal exercise, rate pressure product, HR_{max},

TABLE 2. Aerobic capacity and physiologic responses to maximal treadmill exercise in the CR and AL groups.

		CR		AL		
	$\textbf{Mean} \pm \textbf{SE}$	Within-Group P Value	$\textbf{Mean} \pm \textbf{SE}$	Within-Group P Value	Between-Group P Valu	
VO _{2max} , absolute (L·min ^{−1})						
Baseline	2.53 ± 0.06		2.51 ± 0.09		0.517	
Change at 1 yr	-0.14 ± 0.03	<0.0001	0.00 ± 0.03	0.895	<0.001	
Change at 2 yr	-0.15 ± 0.03	<0.0001	-0.08 ± 0.04	0.129	0.134	
VO _{2max} , relative to body mass (mL·kg b	body mass ^{-1} ·min ^{-1})					
Baseline	34.8 ± 0.6		34.8 ± 1.0		0.996	
Change at 1 yr	$+2.2 \pm 0.4$	<0.0001	$+0.2 \pm 0.5$	0.648	0.004	
Change at 2 yr	+1.9 ± 0.5	<0.001	-1.1 ± 0.7	0.191	<0.001	
VO _{2max} , relative to lean body mass (mL	\cdot kg body mass ⁻¹ ·min ⁻¹)					
Baseline	54.4 ± 0.6		54.8 ± 1.1		0.738	
Change at 1 yr	-0.78 ± 0.54	0.453	$+0.47 \pm 0.70$	0.657	0.453	
Change at 2 yr	-1.05 ± 0.59	0.381	-1.47 ± 0.81	0.381	0.657	
Oxygen pulse at maximal exercise (mL		0.001		0.001	0.001	
Baseline	14.1 ± 0.3		13.9 ± 0.5		0.537	
Change at 1 yr	-0.65 ± 0.15	0.0001	$+0.04 \pm 0.19$	0.848	0.006	
Change at 2 yr	-0.90 ± 0.16	<0.0001	-0.32 ± 0.21	0.241	0.045	
Rate pressure product	0.50 ± 0.10	<0.0001	0.52 ± 0.21	0.241	0.040	
Baseline	31.026 ± 404		30.710 ± 532		0.75	
Change at 1 yr	-2308 ± 530	<0.0001	-323 ± 655	0.621	0.003	
Change at 2 yr	-3850 ± 1241	0.008	-3376 ± 1356	0.038	0.621	
Treadmill time (min)	3030 ± 1241	0.000	5570 ± 1550	0.030	0.021	
Baseline	13.0 ± 0.2		13.2 ± 0.3		0.60	
Change at 1 yr	+0.7 ± 0.1	<0.0001	-0.1 ± 0.2	0.739	<0.001	
Change at 2 yr	+0.7 ± 0.1 +2.9 ± 0.2	<0.0001	-0.1 ± 0.2 +1.8 ± 0.2	<0.0001	<0.001	
HR at maximal exercise (bpm)	+2.9 ± 0.2	<0.0001	$+1.0 \pm 0.2$	<0.0001	<0.0001	
	185 ± 1		185 ± 1		0.94	
Baseline	-2 ± 1	0.003	185 ± 1 -1 ± 1	0.630		
Change at 1 yr	-2 ± 1 -2 ± 1				0.630	
Change at 2 yr		0.113	-0 ± 1	0.642	0.630	
Systolic blood pressure at maximal exe			107 . 0		0.00	
Baseline	168 ± 2	0.001	167 ± 3	0.710	0.80	
Change at 1 yr	-10 ± 3	< 0.001	-1 ± 3	0.710	0.005	
Change at 2 yr	-21 ± 6	0.001	-17 ± 6	0.017	0.569	
Diastolic blood pressure at maximal ex						
Baseline	77 ± 1		77 ± 2		0.52	
Change at 1 yr	-1 ± 1	0.643	+1 ± 2	0.643	0.643	
Change at 2 yr	-6 ± 3	0.361	-8 ± 4	0.140	0.643	
RER at maximal exercise						
Baseline	1.23 ± 0.01		1.23 ± 0.01		0.52	
Change at 1 yr	-0.01 ± 0.01	0.499	-0.02 ± 0.01	0.366	0.975	
Change at 2 yr	-0.00 ± 0.01	0.975	+0.00 \pm 0.01	0.975	0.975	
Borg RPE at maximal exercise						
Baseline	18.0 ± 0.2		18.0 ± 0.2		0.93	
Change at 1 yr	+0.5 \pm 0.2	0.014	+0.3 \pm 0.2	0.357	0.585	
Change at 2 yr	+0.6 ± 0.2	<0.001	$+0.2 \pm 0.2$	0.504	0.444	

VO_{2max}, maximal oxygen uptake. Table includes subjects who achieved VO_{2max}. Sample sizes at baseline, 1 yr, and 2 yr were 130, 101, and 92 for CR and 65, 57, and 49 for AL, respectively.

		and the second	and the second	and the states	and the last of	and the state	. 00
TABLE 3. Body	/ mass,	aerodic ca	pacity, and	muscle strer	ngth by	sex in th	e CK group.

	CR Females		C		
	$\textbf{Mean} \pm \textbf{SE}$	Within-Sex P Value	$\textbf{Mean} \pm \textbf{SE}$	Within-Sex P Value	Between-Sex P Value
Body mass (kg)					
Baseline	67.7 ± 0.6		81.6 ± 1.2		< 0.0001
Change at 2 yr	-7.1 ± 0.4	< 0.0001	-8.5 ± 0.7	<0.0001	0.136
Lean body mass (kg)					
Baseline	40.9 ± 0.4		57.5 ± 0.9		< 0.0001
Change at 2 yr	-1.6 ± 0.2	< 0.0001	-3.2 ± 0.4	< 0.0001	0.003
\dot{VO}_{2max} , absolute (L·min ⁻¹)			012 - 011		01000
Baseline	2.19 ± 0.04		3.22 ± 0.07		<0.0001
Change at 2 yr	-0.20 ± 0.04	<0.0001	-0.09 ± 0.06	0.180	0.180
\dot{VO}_{2max} , relative to body mass (mL·kg body mass		<0.0001	0.05 ± 0.00	0.100	0.100
Baseline	32.5 ± 0.6		39.4 ± 0.8		<0.0001
Change at 2 yr	+0.9 ± 0.6	0.146	$+3.3 \pm 0.9$	0.001	0.073
\dot{VO}_{2max} , relative to lean body mass (mL·kg lean b		0.140	$+0.0 \pm 0.3$	0.001	0.075
					0.004
Baseline	53.7 ± 0.8	0.001	55.9 ± 1.0	0.055	0.024
Change at 2 yr	-2.8 ± 0.7	<0.001	$+0.9\pm1.0$	0.355	0.009
Treadmill time (min)					
Baseline	12.2 ± 0.2		14.5 ± 0.3		<0.001
Change at 2 yr	$+2.3\pm0.2$	<0.0001	$+3.5\pm0.3$	<0.0001	0.005
Isokinetic measures, absolute					
Knee extension at 60°·s ^{−1} (N·m)					
Baseline	122 ± 3		190 ± 5		< 0.0001
Change at 2 yr	-19.5 ± 2.5	< 0.0001	$+0.1 \pm 4.7$	0.976	0.003
Knee extension at 180°⋅s ⁻¹ (N·m)					
Baseline	79 ± 2		128 ± 4		< 0.0001
Change at 2 yr	-10.8 ± 1.6	<0.0001	$+4.9 \pm 2.9$	0.191	0.0001
Knee flexion at $60^{\circ} \cdot \text{s}^{-1}$ (N·m)					
Baseline	64 ± 2		101 ± 3		<0.0001
Change at 2 yr	-5.9 ± 1.3	<0.0001	-1.0 ± 2.4	0.673	0.307
Knee flexion at $180^{\circ} \text{s}^{-1}$ (N·m)	0.0 ± 1.0	-0.0001	1.0 ± 2.4	0.070	0.007
Baseline	51 ± 1		79 ± 3		<0.0001
Change at 2 yr	-6.6 ± 1.1	<0.0001	+0.6 ± 2.0	0.772	0.015
Isokinetic measures, relative to body mass	0.0 ± 1.1	<0.0001	$\pm 0.0 \pm 2.0$	0.772	0.015
Knee extension at 60° ·s ⁻¹ (N·m·kg body mass	-1,				
			0.00 . 0.00		0.0001
Baseline	1.81 ± 0.04	0.070	2.33 ± 0.06	0.0005	< 0.0001
Change at 2 yr	-0.08 ± 0.04	0.079	+0.24 \pm 0.06	0.0005	0.0001
Knee extension at 180°·s ⁻¹ (N·m·kg body mas	,				
Baseline	1.18 ± 0.03		1.57 ± 0.04		<0.0001
Change at 2 yr	-0.00 ± 0.02	0.864	$+0.19 \pm 0.04$	< 0.0001	< 0.001
Knee flexion at 60°·s ⁻¹ (N·m·kg body mass ⁻¹)				
Baseline	0.94 ± 0.02		1.24 ± 0.04		< 0.0001
Change at 2 yr	$+0.02 \pm 0.02$	0.311	$\textbf{+0.12}\pm0.03$	0.002	0.065
Knee flexion at 180°.s ⁻¹ (N·m·kg body mass ⁻	¹)				
Baseline	0.76 ± 0.02		0.97 ± 0.03		< 0.0001
Change at 2 yr	-0.01 ± 0.02	0.696	$+0.10 \pm 0.03$	0.0009	0.005

Sample sizes for VO_{2max} at baseline and 2 yr were 97 and 60 for CR females and 43 and 32 for CR males, respectively.

maximal blood pressure, and peak RER, did not differ by sex (data not shown).

Figures 1 and 2 show changes in \dot{VO}_{2max} (absolute and relative, respectively) versus changes in body mass at 2 yr for each CR and AL subject, by sex. The change in absolute \dot{VO}_{2max} in the CR group from baseline to 2 yr did not correlate with change in whole body mass (P = 0.569), whereas there was a trend for a weak correlation with change in lean body mass (r = 0.204, P = 0.055).

Muscle strength and muscle endurance. As shown in Table 4, absolute knee extensor and knee flexor strength declined significantly in response to CR; at 2 yr, however, these changes were not statistically different than that in the AL group. When expressed relative to body mass, knee extension and knee flexion torque generally increased in CR compared with AL. No group differences were observed when peak torque was expressed relative to leg lean mass (data not shown). Muscle fatigue index increased comparably in CR and AL at both time points. Changes in absolute and relative strength measures in response to 2 yr of CR did not differ between participants who were normal weight or overweight at baseline. Knee extensor and flexor strength measures by sex are shown in Table 3. All strength values were greater for CR males than females, whether expressed in absolute terms or relative to body mass or leg lean mass. Additional sex differences were observed in response to 2 yr of CR. Females had significant decreases in absolute strength and strength expressed relative to leg lean mass, whereas males did not. By contrast, males demonstrated strength increases when expressed relative to body mass. Isometric results were consistent with those of the isokinetic tests (data not shown).

DISCUSSION

This is the first randomized controlled trial to investigate the effects of long-term (i.e., >1 yr) CR on aerobic capacity and muscle strength in nonobese humans. In our population

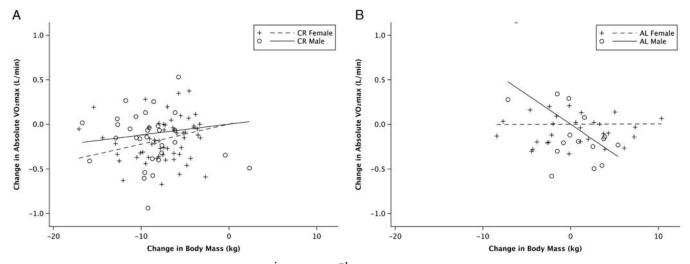


FIGURE 1—Change in absolute maximal oxygen uptake (\dot{VO}_{2max} , $L \min^{-1}$) in relation to change in whole body mass from baseline to 2 yr. CR group (A); AL control group (B). Each symbol represents a single subject. Regression lines are shown for females (dashed lines) and males (solid lines). CR females: $R^2 = 0.3350$, P < 0.0001; CR males: $R^2 = 0.1172$, P = 0.040; AL females: $R^2 = 0.0001$, P = 0.957; AL males: $R^2 = 0.257$, P = 0.007.

of healthy adults, 2 yr of a modest CR diet resulted in decrements in absolute \dot{VO}_{2max} and strength of the knee extensor and flexor muscles, whereas increases in these parameters were observed when expressed relative to body mass. Importantly, CR participants exercised longer on the treadmill and achieved a higher work rate after 2 yr of ~12% CR. This demonstrates that despite significant reductions in lean body mass, leg strength, and oxygen uptake at maximal exercise, cardiorespiratory fitness was preserved.

 \dot{VO}_{2max} is affected primarily by age, sex, and training status in healthy individuals. The reduction in \dot{VO}_{2max} that occurs with normal aging generally averages 10% per decade, as observed in both cross-sectional and longitudinal studies (9), but is highly variable between individuals. The observed decrease in absolute \dot{VO}_{2max} (L·min⁻¹) at 2 yr in our study exceeded the expected age-related decline of approximately 1% per year and was greater in CR than in AL.

Vigorous endurance exercise training has been shown to attenuate the age-related decline in $\dot{V}O_{2max}$ (8,26) but was not part of the CR intervention.

Relative \dot{VO}_{2max} (mL·kg body mass⁻¹·min⁻¹), in contrast to absolute \dot{VO}_{2max} , increased in response to CR. Males in our study tended to increase relative \dot{VO}_{2max} to a greater extent than females, which may be explained by the observation that physical AEE was maintained in males, whereas it decreased in females in response to CR. It is noteworthy that CR participants achieved a higher work rate during the graded treadmill test after 2 yr of CR, reaching stage 8 (4.2 mph, 16% grade) at 2 yr, compared with stage 7 (3.8 mph, 16% grade) at baseline. The AL group reached stage 7 at baseline and 2 yr. Therefore, the increase in the CR group does not appear to be a learning effect of repeated testing. Importantly, the American College of Sports Medicine cardiorespiratory fitness categories (1) are based on relative \dot{VO}_{2max} . There is no evidence of which

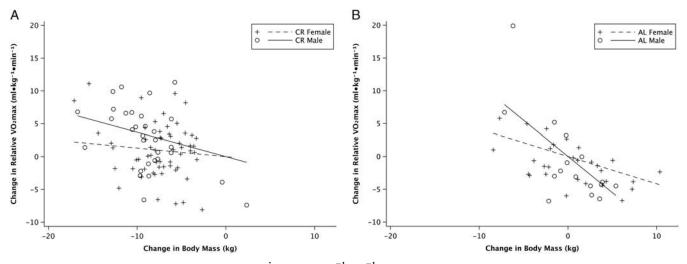


FIGURE 2—Change in relative maximal oxygen uptake (\dot{VO}_{2max} , mL·kg⁻¹·min⁻¹) in relation to change in whole body mass from baseline to 2 yr. CR group (A); AL control group (B). Each symbol represents a single subject. Regression lines are shown for females (dashed lines) and males (solid lines). CR females: $R^2 = 0.052$, P = 0.049; CR males: $R^2 = 0.364$, P < 0.0001; AL females: $R^2 = 0.299$, P < 0.001; AL males: $R^2 = 0.374$, P < 0.0001.

		CR		AL	
	$\textbf{Mean} \pm \textbf{SE}$	Within-Group P Value	$\textbf{Mean} \pm \textbf{SE}$	Within-Group P Value	Between-Group P Value
Isokinetic measures, absolute					
Knee extension at 60°⋅s ⁻¹ (N·m)					
Baseline	143 ± 4		138 ± 5		0.527
Change at 1 yr	-10.9 ± 1.7	<0.0001	-1.3 ± 2.1	0.527	0.0004
Change at 2 yr	-10.3 ± 2.3	<0.0001	-7.2 ± 2.7	0.023	0.527
Knee extension at 180°·s ⁻¹ (N·m)					
Baseline	94 ± 3		92 ± 3		0.487
Change at 1 yr	-6.2 ± 1.1	<0.0001	-0.3 ± 1.4	0.841	0.001
Change at 2 yr	-3.2 ± 1.4	0.084	-2.2 ± 1.7	0.583	0.841
Knee flexion at 60°⋅s ⁻¹ (N⋅m)					
Baseline	75 ± 2		75 ± 3		0.963
Change at 1 yr	-4.4 ± 1.1	0.0003	-1.6 ± 1.4	0.333	0.318
Change at 2 yr	-3.0 ± 1.2	0.044	-1.4 ± 1.4	0.333	0.333
Knee flexion at 180°·s ⁻¹ (N·m)					
Baseline	60 ± 2		59 ± 2		0.751
Change at 1 yr	-4.0 ± 0.8	<0.0001	-1.1 ± 1.1	0.602	0.088
Change at 2 yr	-3.2 ± 1.0	0.044	-0.4 ± 1.2	0.744	0.123
Isokinetic measures, relative to body mass					
Knee extension at 60°·s ⁻¹ (N·m·kg body r	mass ⁻¹)				
Baseline	, 1.97 ± 0.04		1.93 ± 0.05		0.441
Change at 1 yr	$+0.08 \pm 0.03$	0.008	-0.00 ± 0.03	0.908	0.061
Change at 2 yr	$+0.07 \pm 0.03$	0.061	-0.10 ± 0.04	0.061	0.004
Knee extension at 180°.s ⁻¹ (N·m·kg body	$mass^{-1}$)				
Baseline	1.30 ± 0.03		1.27 ± 0.03		0.543
Change at 1 yr	$+0.07 \pm 0.02$	<0.0001	$+0.00 \pm 0.02$	0.926	0.015
Change at 2 yr	$+0.09 \pm 0.02$	<0.0001	-0.03 ± 0.03	0.396	0.0002
Knee flexion at 60°·s ⁻¹ (N·m·kg body mas	ss ⁻¹)				
Baseline	1.03 ± 0.02		1.04 ± 0.03		0.996
Change at 1 yr	$+0.06 \pm 0.02$	0.0003	-0.01 ± 0.02	0.544	0.005
Change at 2 yr	$+0.07 \pm 0.02$	0.0002	-0.02 ± 0.02	0.544	0.001
Knee flexion at 180°·s ⁻¹ (N·m·kg body ma	ass^{-1})				
Baseline	0.82 ± 0.02		0.82 ± 0.02		0.676
Change at 1 yr	$+0.04 \pm 0.01$	0.005	-0.01 ± 0.02	0.983	0.053
Change at 2 yr	$+0.05 \pm 0.01$	0.005	-0.00 ± 0.02	0.983	0.071
Isometric measures, absolute					
Isometric knee extension at 45° (N·m)					
Baseline	124 ± 3		122 ± 4		0.830
Change at 1 yr	-5.0 ± 1.8	0.0193	$+2.7 \pm 2.3$	0.274	0.019
Change at 2 yr	-6.1 ± 1.9	0.0104	-2.6 ± 2.4	0.274	0.274
Isometric knee flexion at 45° (N·m)					
Baseline	97 ± 2		96 ± 3		0.695
Change at 1 yr	-5.6 ± 1.3	<0.0001	-3.1 ± 1.6	0.173	0.381
Change at 2 yr	-6.3 ± 1.6	0.0005	-4.9 ± 2.0	0.048	0.552
Isometric measures, relative to body mass					
Isometric knee extension at 45° (N·m·kg b	body mass $^{-1}$)				
Baseline	1.70 ± 0.03		1.70 ± 0.05		0.666
Change at 1 yr	$+0.13 \pm 0.03$	< 0.0001	$\textbf{+0.05} \pm 0.03$	0.295	0.112
Change at 2 yr	+0.09 \pm 0.03	0.0102	-0.04 ± 0.04	0.295	0.014
Isometric knee flexion at 45° (N·m·kg bod	y mass ⁻¹)				
Baseline	1.33 ± 0.03		1.33 ± 0.03		0.949
Change at 1 yr	$+0.09 \pm 0.02$	< 0.0001	-0.03 ± 0.02	0.182	< 0.0001
Change at 2 yr	$+0.07 \pm 0.02$	0.014	-0.06 ± 0.03	0.087	0.001
Muscle endurance					
Work fatigue index (%), average of knee f	lexion and extension at 1	80°·s ⁻¹			
Baseline	28.9 ± 1.2		29.7 ± 1.5		0.69
Change at 1 yr	+3.1 ± 0.7	< 0.0001	+4.2 ± 0.9	< 0.0001	0.482
Change at 2 yr	$+3.8 \pm 0.8$	< 0.0001	$+2.9 \pm 1.1$	0.019	0.482

Values for knee extension and flexion represent peak torque, averaged across the right and left legs. Sample sizes at baseline, 1 yr, and 2 yr were 141, 128, and 114 for CR and 72, 67, and 65 for AL, respectively.

we are aware that a decrease in absolute $\dot{V}O_{2max}$ without a decrease in relative $\dot{V}O_{2max}$ is detrimental or has clinical implications with respect to cardiovascular health or mortality risk. This is an important area of investigation for future CR studies.

The smaller body mass resulting from long-term CR may enable greater economy of movement that likely offsets the lower absolute $\dot{V}O_2$. Long-term CR studies in rhesus monkeys support this view (32). Another intriguing possibility is that skeletal muscle mitochondrial adaptations may occur in response to CR, as reviewed by Hepple (10). Specifically, mitochondrial biogenesis is induced in response to CR in rodents (17) and humans (5); this may be explained by peroxisome proliferator-activated receptor γ coactivator 1 α gene expression in skeletal and cardiac muscle with CR, as demonstrated in rodents (11). Peroxisome proliferator-activated receptor γ coactivator 1 α is an inducible transcriptional coregulator of mitochondrial function that is activated in response to exercise, cold exposure, and CR (2).

Consistent with the results of our CALERIE phase 1 study (30), absolute muscle strength declined in response to CR in

the current study, whereas muscle strength expressed relative to body weight increased. The current multicenter CALERIE 2 results build upon the findings of the earlier, single-center report in which CR was compared with an exercise-induced energy deficit (30). Distinct characteristics of the CALERIE 2 study design include the longer duration of the CR intervention (2 vs 1 yr), greater %CR prescribed (25% vs 16%-20%), younger age range of participants (21-50 vs 50-60 yr), larger study sample size (218 vs 34), and a larger sample of CR participants (143 vs 18). The clinical significance of the observed decrements in absolute strength in these studies is unclear. In the absence of sex- and age-specific muscle strength reference values and their associations with long-term health outcomes, it is difficult to predict whether such changes are deleterious with respect to functional capacity and disease risk. A decline in absolute strength is to be expected after 2 yr of a CR intervention that does not include resistance training; future CR studies should include a structured resistance training component to evaluate its potential in preserving lean body mass and absolute strength.

Interestingly, the changes in strength in response to 2 yr of CR differed significantly by sex, with only the female CR participants demonstrating a decline in absolute strength and only the male participants demonstrating an increase in strength expressed relative to body weight. The explanation for this finding is unclear, particularly in light of the greater loss of lean body mass among male participants. There were no significant relationships observed between changes in absolute leg strength and changes in lean body mass or whole body mass at 2 yr in the overall sample. Likewise, dietary protein did not appear to influence the changes in lean body mass or muscle strength at 2 yr.

A notable limitation of this study is that the average level of CR achieved was substantially less than the 25% prescribed, despite frequent contact and encouragement by research

REFERENCES

- 1. American College of Sports Medicine. *ACSM's Guidelines for Exercise Testing and Prescription*. 9th ed. Philadelphia, PA: Lippincott Williams & Wilkins; 2014.
- Anderson RM, Barger JL, Edwards MG, et al. Dynamic regulation of PGC-1alpha localization and turnover implicates mitochondrial adaptation in calorie restriction and the stress response. *Aging Cell*. 2008;7(1):101–11.
- Bales CR, Kraus WE. Caloric restriction: implications for human cardiometabolic health. J Cardiopulm Rehabil Prev. 2013;33(4):201–8.
- Borg G. Perceived exertion as an indicator of somatic stress. Scand J Rehabil Med. 1970;2(2):92–8.
- Civitarese AE, Carling S, Heilbronn LK, et al. Calorie restriction increases muscle mitochondrial biogenesis in healthy humans. *PLoS Med.* 2007;4(3):e76.
- Das SK, Roberts SB, Bhapkar MV, et al. Body-composition changes in the Comprehensive Assessment of Long-term Effects of Reducing Intake of Energy (CALERIE)-2 study: a 2-y randomized controlled trial of calorie restriction in nonobese humans. *Am J Clin Nutr.* 2017;105(4):913–27.
- Fontana L, Hu FB. Optimal body weight for health and longevity: bridging basic, clinical, and population research. *Aging Cell*. 2014; 13(3):391–400.

personnel. Therefore, we are unable to determine whether 25% CR would result in deleterious changes in \dot{VO}_{2max} or muscle strength that might impair cardiorespiratory or functional capacity. Additional factors that may have influenced our results include differences in participant demographics at the three clinical sites and variability in diet composition and diet quality among participants. Furthermore, some subjects were unable to complete the \dot{VO}_{2max} or muscle strength tests for orthopedic reasons, some did not meet the criteria for achieving \dot{VO}_{2max} , and others dropped out of the study. Strengths of the CALERIE 2 trial include a relatively large and diverse sample of individuals from three distinct regions of the United States, a long-term CR intervention, common protocols for the CR intervention and all assessments, and a high degree of quality control.

In summary, we observed that 2 yr of CR in nonobese, healthy, young, and middle-age adults resulted in higher relative aerobic capacity, despite significant reductions in absolute $\dot{V}O_{2max}$, absolute strength, whole body mass, lean body mass, and leg lean mass. In addition, treadmill exercise time and work rate increased after 2 yr of CR, suggesting that long-term, modest CR with adequate nutrient intake does not adversely affect cardiorespiratory function in healthy adults.

This project was supported by grants U01AG020478, U01AG020480, U01AG020487, U01AG022132, and U24AG047121 from the National Institute on Aging; grants P30DK056341 and P30DK072476 from the National Institute of Diabetes and Digestive and Kidney Disease. The authors thank the study participants at each site for their dedication to this project and all the research personnel that contributed their expertise and skills. None of the authors or anyone involved with this research had a conflict of interest, and the results of the present study do not constitute endorsement by the American College of Sports Medicine.

On behalf of the CALERIE Study Group, the authors would like to thank the study participants at each site for their dedication to this project and all the research personnel who contributed their expertise and skills.

- Grimsmo J, Arnesen H, Maehlum S. Changes in cardiorespiratory function in different groups of former and still active male crosscountry skiers: a 28–30-year follow-up study. *Scand J Med Sci Sports.* 2010;20(1):e151–61.
- Hawkins S, Wiswell R. Rate and mechanism of maximal oxygen consumption decline with aging: implications for exercise training. *Sports Med.* 2003;33(12):877–88.
- Hepple RT. Why eating less keeps mitochondria working in aged skeletal muscle. *Exerc Sport Sci Rev.* 2009;37(1):23–8.
- Hepple RT, Baker DJ, McConkey M, Murynka T, Norris R. Caloric restriction protects mitochondrial function with aging in skeletal and cardiac muscles. *Rejuvenation Res.* 2006;9(2): 219–22.
- Hochberg Y. A sharper Bonferroni procedure for multiple tests of significance. *Biometrika*. 1988;75(4):800–2.
- Holloszy JO, Fontana L. Caloric restriction in humans. *Exp Gerontol.* 2007;42(8):709–12.
- Huber PJ. 1972 Wald Memorial Lectures—Robust Regression— Asymptotics, Conjectures and Monte-Carlo. Ann Stat. 1973;1(5): 799–821.
- 15. Huber PJ, Ronchetti EM. *Robust Statistics*. 2nd ed. Hoboken (NJ): Wiley; 2009.

- Institute of Medicine, and Food and Nutrition Board. Dietary Reference Intakes for Energy, Carbohydrate, Fiber, Fat, Fatty Acids, Cholesterol, Protein, and Amino Acids. Washington (DC): The National Academies Press, 2005.
- Lopez-Lluch G, Hunt N, Jones B, et al. Calorie restriction induces mitochondrial biogenesis and bioenergetic efficiency. *Proc Natl Acad Sci U S A*. 2006;103(6):1768–73.
- Martin CK, Das SK, Lindblad L, et al. Effect of calorie restriction on the free-living physical activity levels of nonobese humans: results of three randomized trials. J Appl Physiol (1985). 2011;110(4):956–63.
- Mattison JA, Colman RJ, Beasley TM, et al. Caloric restriction improves health and survival of rhesus monkeys. *Nat Commun.* 2017;8:14063.
- Racette SB, Das SK, Bhapkar M, et al. Approaches for quantifying energy intake and %calorie restriction during calorie restriction interventions in humans: the multicenter CALERIE study. *Am J Physiol Endocrinol Metab.* 2012;302(4):E441–8.
- Racette SB, Weiss EP, Villareal DT, et al. One year of caloric restriction in humans: feasibility and effects on body composition and abdominal adipose tissue. J Gerontol A Biol Sci Med Sci. 2006;61(9):943–50.
- 22. Ravussin E, Redman LM, Rochon J, et al. A 2-year randomized controlled trial of human caloric restriction: feasibility and effects on predictors of health span and longevity. *J Gerontol A Biol Sci Med Sci.* 2015;70(9):1097–104.
- Redman LM, Heilbronn LK, Martin CK, et al. Metabolic and behavioral compensations in response to caloric restriction: implications for the maintenance of weight loss. *PLoS One*. 2009;4(2):e4377.
- Rickman AD, Williamson DA, Martin CK, et al. The CALERIE study: design and methods of an innovative 25% caloric restriction intervention. *Contemp Clin Trials*. 2011;32(6):874–81.

- Rochon J, Bales CW, Ravussin E, et al. Design and conduct of the CALERIE study: comprehensive assessment of the long-term effects of reducing intake of energy. *J Gerontol A Biol Sci Med Sci*. 2011;66(1):97–108.
- Rogers MA, Hagberg JM, Martin WH 3rd, Ehsani AA, Holloszy JO. Decline in VO_{2max} with aging in master athletes and sedentary men. *J Appl Physiol (1985)*. 1990;68(5):2195–9.
- 27. Ross R, Blair SN, Arena R, et al.; American Heart Association Physical Activity Committee of the Council on Lifestyle and Cardiometabolic Health; Council on Clinical Cardiology; Council on Epidemiology and Prevention; Council on Cardiovascular and Stroke Nursing; Council on Functional Genomics and Translational Biology; Stroke Council. Importance of assessing cardiorespiratory fitness in clinical practice: a case for fitness as a clinical vital sign: a scientific statement from the American Heart Association. *Circulation.* 2016;134(24):e653–99.
- Tamesis B, Stelken A, Byers S, et al. Comparison of the Asymptomatic Cardiac Ischemia Pilot and modified Asymptomatic Cardiac Ischemia Pilot versus Bruce and Cornell exercise protocols. *Am J Cardiol.* 1993;72(9):715–20.
- 29. Tanaka H, Monahan KD, Seals DR. Age-predicted maximal heart rate revisited. *J Am Coll Cardiol*. 2001;37(1):153–6.
- Weiss EP, Racette SB, Villareal DT, et al. Lower extremity muscle size and strength and aerobic capacity decrease with caloric restriction but not with exercise-induced weight loss. *J Appl Physiol* (1985). 2007;102(2):634–40.
- Wright SP. Adjusted P-values for simultaneous inference. *Biometrics*. 1992;48:1005–13.
- 32. Yamada Y, Colman RJ, Kemnitz JW, et al. Long-term calorie restriction decreases metabolic cost of movement and prevents decrease of physical activity during aging in rhesus monkeys. *Exp Gerontol.* 2013;48(11):1226–35.