

# Dietary omega-3 fatty acid supplementation increases the rate of muscle protein synthesis in older adults: a randomized controlled trial<sup>1–3</sup>

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## ABSTRACT

**Background:** Loss of muscle mass with aging is a major public health concern. Omega-3 (n-3) fatty acids stimulate protein anabolism in animals and might therefore be useful for the treatment of sarcopenia. However, the effect of omega-3 fatty acids on human protein metabolism is unknown.

**Objective:** The objective of this study was to evaluate the effect of omega-3 fatty acid supplementation on the rate of muscle protein synthesis in older adults.

**Design:** Sixteen healthy, older adults were randomly assigned to receive either omega-3 fatty acids or corn oil for 8 wk. The rate of muscle protein synthesis and the phosphorylation of key elements of the anabolic signaling pathway were evaluated before and after supplementation during basal, postabsorptive conditions and during a hyperaminoacidemic-hyperinsulinemic clamp.

**Results:** Corn oil supplementation had no effect on the muscle protein synthesis rate and the extent of anabolic signaling element phosphorylation in muscle. Omega-3 fatty acid supplementation had no effect on the basal rate of muscle protein synthesis (mean  $\pm$  SEM:  $0.051 \pm 0.005\%/h$  compared with  $0.053 \pm 0.008\%/h$  before and after supplementation, respectively;  $P = 0.80$ ) but augmented the hyperaminoacidemia-hyperinsulinemia-induced increase in the rate of muscle protein synthesis (from  $0.009 \pm 0.005\%/h$  above basal values to  $0.031 \pm 0.003\%/h$  above basal values;  $P < 0.01$ ), which was accompanied by greater increases in muscle mTOR<sup>Ser2448</sup> ( $P = 0.08$ ) and p70s6k<sup>Thr389</sup> ( $P < 0.01$ ) phosphorylation.

**Conclusion:** Omega-3 fatty acids stimulate muscle protein synthesis in older adults and may be useful for the prevention and treatment of sarcopenia. This trial was registered at clinicaltrials.gov as NCT00794079. *Am J Clin Nutr* 2011;93:402–12.

## INTRODUCTION

Loss of muscle mass with aging is a major public health concern because it impairs physical function, which reduces quality of life and may lead to frailty and premature death (1–4). A major cause for the loss of muscle mass with advanced age is the inability of aging muscle to adequately increase the rate of muscle protein synthesis in response to nutritional stimuli (eg, amino acids and insulin) (5–7). The blunted anabolic response to nutritional stimuli is at least in part due to defects in the anabolic signaling cascade in muscle (ie, decreased activation of the mTOR-p70s6k signaling pathway) (5, 6), which may be mediated by increased inflammatory activity (8–10). Interventions that can overcome this anabolic resistance are therefore likely to prevent or slow the progression of muscle loss with aging.

Some evidence suggests that fish-oil-derived omega-3 fatty acids (n-3 polyunsaturated fatty acids) might be a potentially useful therapeutic agent for the treatment and prevention of sarcopenia. It has been shown (11) that providing feed enriched in menhaden oil to growing steers increases the activation (phosphorylation) of anabolic signaling proteins in muscle during administration of insulin and amino acids and increases the nonoxidative whole-body disposal of amino acids—an index of increased whole-body protein synthesis (the effect on muscle protein synthesis was not determined). Furthermore, omega-3 fatty acid supplementation has been shown to prevent loss of muscle mass in burned guinea pigs (12). In addition, omega-3 fatty acids have antiinflammatory properties (13), which may also help alleviate the muscle anabolic resistance in older adults. However, the effect of omega-3 fatty acid intake on human muscle protein metabolism and the intracellular signaling pathways that might regulate it are not known.

The purpose of the present study therefore was to determine the effect of dietary omega-3 fatty acid supplementation on the rate of muscle protein synthesis and the anabolic signaling cascade in older adults. We hypothesized that dietary omega-3 fatty acid supplementation increases the stimulatory effect of hyperaminoacidemia-hyperinsulinemia on anabolic signaling and protein synthesis in muscle.

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## SUBJECTS AND METHODS

### Subjects and preliminary testing

Sixteen older adults (10 men and 6 women  $\geq 65$  y of age) participated in this randomized controlled trial (ClinicalTrials.gov number NCT00794079). Subjects were considered eligible for the trial if they 1) were in good health (ie, no evidence of significant cardiovascular disease or organ dysfunction, including hypertension, dyslipidemia, and diabetes mellitus) after completing a comprehensive medical evaluation, which included a history and physical examination and standard blood tests; 2) were not obese [body mass index  $< 30.0$  (in  $\text{kg}/\text{m}^2$ )]; 3) did not engage in regular physical activities (ie, they exercised  $\leq 1$  h/wk); 4) did not consume fish-oil supplements; 5) did not take medications that could affect muscle protein metabolism; 6) did not consume tobacco products; and 7) did not report excessive alcohol intake. Each subject's total body fat-free mass (FFM) and fat mass were measured by using dual-energy X-ray absorptiometry (DXA; Delphi-W densitometer; Hologic, Waltham, MA) as part of the screening evaluation to help us adjust the amino acid infusion to FFM during the muscle protein study. We decided a priori to not perform DXA analysis after treatment to measure changes in appendicular lean body mass in this study because we determined that it would require a much bigger sample size to reliably measure the changes that could be achieved during the 8-wk intervention period in our study, or, alternatively, require a much longer duration of the intervention. On completion of the screening procedure, subjects were randomly assigned to either the omega-3 fatty acid or corn oil (control) group. One individual randomly assigned to the control group withdrew from the study after completing the initial assessment. Therefore, data from only 7 subjects are presented in the control group.

Written informed consent was obtained from all subjects before their participation in the study, which was approved by the Human Research Protection Office and the Clinical Research Unit Advisory Committee at Washington University School of Medicine (St Louis, MO).

### Experimental protocol

Each subject completed 2 stable-isotope-labeled tracer infusion studies to determine the effect of omega-3 fatty acid or corn oil (control group) supplementation on the rate of muscle protein synthesis and anabolic signaling in muscle during basal, postabsorptive conditions and combined amino acid and insulin infusions. The first study was performed within 1–3 wk of screening (before the intervention); the second study took place after 8 wk of dietary supplementation with omega-3 fatty acids [4 g Lovaza/d containing 1.86 g eicosapentaenoic acid (EPA, 20:5n-3) and 1.50 g docosahexaenoic acid (DHA, 22:6n-3)—both as ethyl esters—or an equal amount of corn oil]. We chose this dose because it is the dose approved by the Food and Drug Administration for lowering plasma triglyceride concentrations in hypertriglyceridemic subjects and has therefore previously been shown to be physiologically relevant in human subjects. Furthermore, similar doses (relative to body weight) increased whole-body protein synthesis (no muscle data were reported) in burned rats (14) and doubled the insulin-stimulated nonoxidative whole-body disposal of amino acids, a marker of increased whole-body protein synthesis (again, no measures of muscle

protein synthesis were obtained), in growing steer (11), whereas much higher doses (and possibly even toxic amounts of fish oil) resulted in no effect on protein metabolism or even decreased muscle protein synthesis rates in rats (15–18). Both the omega-3 fatty acid and corn oil treatments were provided by Reliant Pharmaceuticals Inc, Liberty Corner, NJ (now GlaxoSmith-Kline, Research Triangle Park, NC) and packaged in identical-looking capsules, and subjects were blinded to the treatment. Compliance was evaluated by pill count. To minimize falsification of the pill count, subjects were given an unknown (to them) number of pills in excess of needs and were asked to return any remaining pills at the end of the study.

Before each muscle protein metabolism study, subjects were instructed to adhere to their usual diet and to refrain from vigorous physical activities for 3 d. On the evening before the study, subjects were admitted to the Clinical Research Unit at Washington University School of Medicine. At 2000, they consumed a standard meal providing 50.2 kJ/kg body wt (15% as protein, 55% as carbohydrates, and 30% as fat). Subjects then rested in bed and fasted (except for water) until completion of the study the next day. At  $\approx 0600$  h on the following morning, a cannula was inserted into an antecubital vein for the infusion of stable-isotope-labeled tracers (ie, a phenylalanine tracer to measure the rate of muscle protein synthesis and phenylalanine rate of appearance in the systemic circulation and a glucose tracer to measure glucose rate of appearance in the systemic circulation); another cannula was inserted into a vein of the contralateral hand, which was warmed to 55°C for blood sampling. At  $\approx 0800$  h, primed, constant infusions of [ring- $^2\text{H}_5$ ]phenylalanine (priming dose: 2.8  $\mu\text{mol}/\text{kg}$  FFM, infusion rate: 0.08  $\mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$ ) and [6,6- $^2\text{H}_2$ ]glucose (priming dose: 18  $\mu\text{mol}/\text{kg}$  body wt, infusion rate: 0.22  $\mu\text{mol} \cdot \text{kg body wt}^{-1} \cdot \text{min}^{-1}$ ), both purchased from Cambridge Isotope Laboratories Inc (Andover, MA), were started and maintained for 7 h. Four hours after the start of the tracer infusions, a hyperaminoacidemic-hyperinsulinemic clamp was started and maintained for 3 h. Human insulin (Novolin R; Novo Nordisk, Princeton, NJ) was infused at a rate of 20  $\text{mU} \cdot \text{m}^2$  body surface area (BSA) $^{-1} \cdot \text{min}^{-1}$  (initiated with priming doses of 80  $\text{mU} \cdot \text{m}^2$  BSA $^{-1} \cdot \text{min}^{-1}$  for 5 min and then 40  $\text{mU} \cdot \text{m}^2$  BSA $^{-1} \cdot \text{min}^{-1}$  for an additional 5 min), and plasma amino acid availability was increased by providing an intravenous amino acid mixture (Travasol 10%; Baxter, Deerfield, IL) at a rate of 105 mg amino acids  $\cdot \text{kg FFM}^{-1} \cdot \text{h}^{-1}$  (priming dose: 35 mg amino acids/kg FFM) to raise plasma insulin and amino acid concentrations to within the range normally seen after meal consumption (19, 20). Euglycemia (blood glucose concentration of  $\approx 5.5$  mmol/L) was maintained during the clamp procedure by variable rate infusion of 20% dextrose (Baxter) enriched to 2.5% with [6,6- $^2\text{H}_2$ ]glucose. To minimize changes in plasma phenylalanine and glucose isotopic enrichment during the clamp due to the increased amino acid rate of appearance in plasma and reduced hepatic glucose production, the [ring- $^2\text{H}_5$ ]phenylalanine infusion rate was increased to 0.12  $\mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$  and the [6,6- $^2\text{H}_2$ ]glucose infusion rate was decreased to 0.11  $\mu\text{mol} \cdot \text{kg body wt}^{-1} \cdot \text{min}^{-1}$  during the clamp.

Blood samples (4 mL) were obtained before the tracer infusions began and at 60, 90, 120, 150, 180, 210, 220, 230, 240, 270, 300, 330, 360, 390, 400, 410, and 420 min to determine the labeling of phenylalanine and glucose in plasma and plasma substrate, hormone, and cytokine concentrations; plasma was



separated and stored at  $-80^{\circ}\text{C}$  until the final analyses. Additional blood ( $\approx 1$  mL) was obtained every 10 min during the clamp to monitor plasma glucose concentrations. Muscle tissue ( $\approx 100$  mg) was obtained under local anesthesia (lidocaine, 2%) from the quadriceps femoris by using a Tilley-Henkel forceps at 60 and 240 min to determine the basal rate of muscle protein synthesis (labeled phenylalanine incorporation into muscle protein; *see* section entitled "Calculations") and the basal concentrations (240 min) of phosphorylated elements of intramuscular signal transduction proteins (Akt; mTOR; p70s6k) involved in the regulation of glucose uptake and muscle protein synthesis (21, 22). A third muscle biopsy sample was obtained at 420 min (ie, 3 h after the clamp procedure was started) to determine both the rate of muscle protein synthesis and the intracellular signaling responses to hyperaminoacidemia-hyperinsulinemia. The second and third biopsy samples were obtained from the same incision on the leg contralateral to that biopsied first; the forceps was directed in proximal and distal directions, respectively, so that the 2 biopsy samples were collected  $\approx 5$ –10 cm apart. Muscle samples were initially frozen in liquid nitrogen and stored at  $-80^{\circ}\text{C}$  until the final analysis.

### Sample processing and analyses

Plasma glucose concentrations were measured with an automated glucose analyzer (Yellow Spring Instruments, Yellow Springs, OH). Plasma insulin concentrations were measured by radioimmunoassay (Linco Research, St Louis, MO). Commercially available enzyme-linked immunosorbent assay kits (R&D Systems Inc, Minneapolis, MN) were used to measure plasma concentrations of C-reactive protein (CRP), tumor necrosis factor- $\alpha$  (TNF- $\alpha$ ), and interleukin-6 (IL-6).

Muscle phospholipid fatty acid composition was determined after the extraction of lipids from  $\approx 30$  mg muscle tissue with 2 mL chloroform/methanol (2:1, vol:vol) containing 0.01% butylated hydroxytoluene. Phospholipids were then isolated by using thin-layer chromatography (Whatman TLC LK 6D; Fischer Scientific, Pittsburgh, PA), fatty acids were converted to their methyl esters by reacting with 10% acetyl chloride in methanol, and their relative concentrations were measured by using gas chromatography–mass spectrometry (GC-MS; MSD 5973 System; Hewlett-Packard). The proportional contribution of each fatty acid to total fatty acid content was calculated by dividing the peak area of each fatty acid by the sum of all peak areas.

To determine the labeling of plasma glucose, plasma proteins were precipitated with ice-cold acetone, and hexane was used to extract plasma lipids. The aqueous phase, containing glucose, was dried by speed-vac centrifugation (Savant Instruments, Farmingdale, NY), glucose was derivatized with heptafluorobutyric acid, and the tracer-to-tracee ratio (TTR) was determined by using GC-MS as previously described (23).

To measure the plasma concentrations of phenylalanine and leucine (thought to be a major regulator of muscle protein synthesis; 24) and the labeling of phenylalanine in plasma, known amounts of [ $1$ - $^{13}\text{C}$ ]phenylalanine and nor-leucine were added to an aliquot of each plasma sample, plasma proteins were precipitated, and the supernatant fluid (containing free amino acids) was collected to prepare the *t*-butyldimethylsilyl derivative of phenylalanine and leucine for analysis by GC-MS (25, 26). To determine phenylalanine labeling in muscle pro-

teins and in tissue fluid, samples ( $\approx 20$  mg) were homogenized in 1 mL trichloroacetic acid solution (3% wt:vol); proteins were precipitated by centrifugation and then hydrolyzed, and the supernatant fluid, containing free amino acids, was collected. Amino acids in the protein hydrolysate and supernatant fluid samples were purified on cation-exchange columns (Dowex 50W-X8-200; Bio-Rad Laboratories, Richmond, CA), and the *t*-butyldimethylsilyl derivative of phenylalanine prepared to determine its TTR by GC-MS analysis (25, 26).

Western analysis was used to measure the phosphorylation of Akt, mTOR, and p70s6k at 240 min (basal) and 420 min (clamp) as previously described (25–28). Briefly, frozen muscle tissue ( $\approx 20$  mg) was rapidly homogenized in ice-cold buffer, proteins were extracted, and 50  $\mu\text{g}$  protein from each sample was loaded onto 12% XT-Bis Tris gels, separated by sodium dodecyl sulfate–polyacrylamide gel electrophoresis, and transferred on ice to methanol prewetted 0.2- $\mu\text{m}$  PVDF membranes. Blots were then incubated sequentially with 5% (wt:vol) nonfat milk for 1 h, primary antibodies (Akt<sup>Thr308</sup>, mTOR<sup>Ser2448</sup>, p70s6k<sup>Thr389</sup>, and pan-actin as a loading control; all purchased from New England Biolabs) overnight at  $4^{\circ}\text{C}$ , and then secondary antibody (anti-rabbit; New England Biolabs, Ipswich, MA) for 1 h. Data were expressed in relation to GAPDH.

### Calculations

The muscle protein fractional synthesis rate (FSR) was calculated from the rate of incorporation of [ring- $^2\text{H}_5$ ]phenylalanine into muscle protein, by using a standard precursor-product model as follows:

$$\text{FSR} = \Delta E_p / E_{ic} \times 1/t \times 100 \quad (1)$$

where  $\Delta E_p$  is the change between 2 consecutive biopsy samples in extent of labeling (TTR) of protein-bound phenylalanine.  $E_{ic}$  is the phenylalanine labeling in the precursor pool, and  $t$  is the time between biopsy sample collections. We used both the free phenylalanine labeling in muscle tissue fluid (at 240 and 420 min) and the average plasma phenylalanine labeling between 60 and 240 min (basal) and 270 and 420 min (clamp) as surrogates for the phenylalanine labeling in the precursor pool (ie, aminoacyl-*t*-RNA) (29).

Glucose and phenylalanine rates of appearance (Ra) in plasma were calculated by dividing the tracer infusion rate by the average TTR in plasma during the last 30 min of the basal period and the last 30 min of the clamp. Glucose and phenylalanine Ra during basal conditions represent endogenous glucose and phenylalanine production; during the clamp procedure, glucose and phenylalanine Ra represent the sum of endogenous glucose and phenylalanine production and the rate of infused glucose and phenylalanine, respectively. Endogenous Ra during the clamp was therefore calculated by subtracting the glucose (dextrose plus tracer) and phenylalanine (Travasol and tracer) infusion rate from total glucose and phenylalanine Ra, respectively.

### Statistical analysis

Statistical analysis was performed by using SPSS version 13 software (SPSS Inc, Chicago, IL). All data sets were tested for normality, and skewed data sets were log transformed for data analysis or evaluated by using appropriate nonparametric testing procedures. Preliminary analysis of variance (ANOVA) was





performed to examine whether differences between the control and omega-3 fatty acid groups existed in subject characteristics; plasma glucose, insulin, phenylalanine, and leucine concentrations; mixed muscle protein FSR; and the concentration of muscle intracellular signaling elements before the intervention. No differences were detected. For each group (corn oil compared with omega-3 fatty acid), repeated-measures ANOVA and Tukey's post hoc test were therefore used to evaluate before-to-after intervention changes (effect of time) in plasma glucose, insulin, phenylalanine, and leucine concentrations; mixed muscle protein FSR; and muscle intracellular signaling elements during basal and postabsorptive conditions and during the clamp procedure in the omega-3 fatty acid and corn oil supplementation groups. Differences between groups in the treatment-induced change, if they existed, were evaluated by using Student's *t* test for independent samples or the Mann-Whitney *U* test (for skewed data sets). Differences in measurements made only during the basal postabsorptive state (eg, systemic inflammatory markers, muscle phospholipid fatty acid composition) before and after omega-3 fatty acid or corn oil supplementation were evaluated by using ANOVA with group and time as the factors. A *P* value  $\leq 0.05$  was considered significant. Data are presented as means  $\pm$  SEMs or medians with 25th and 75th percentiles for skewed data sets.

## RESULTS

### Subject characteristics, metabolic variables, and body composition

Subjects randomly assigned to receive omega-3 fatty acids and corn oil were similar in age, sex, BMI, and body composition, and

their health and metabolic status (ie, blood pressure, plasma lipid panel, and plasma inflammatory markers) were similar both before and after corn oil and omega-3 fatty acid supplementation (Table 1). Only diastolic blood pressure was somewhat lower in the omega-3 fatty acid group ( $P < 0.05$ ), and LDL-cholesterol and TNF- $\alpha$  concentrations slightly increased in response to both the corn oil and omega-3 fatty acid supplementation (main effect of time:  $P < 0.05$ ), but no significant group  $\times$  time interaction was observed.

### Compliance with omega-3 fatty acid and corn oil supplementation and muscle phospholipid fatty acid composition

All subjects consumed  $\geq 160$  of the 224 pills assigned to them. Average compliance, as judged by the leftover pill count, was 100% and  $96 \pm 4\%$  for the corn oil and omega-3 fatty acid groups, respectively. This was confirmed by analysis of the muscle phospholipid fatty acid profile. Omega-3 fatty acid supplementation approximately doubled ( $P < 0.01$ ) the muscle phospholipid omega-3 fatty acid content, whereas corn oil supplementation had no effect on the muscle phospholipid fatty acid composition (Table 2).

### Plasma insulin, glucose, and amino acid concentrations and glucose and phenylalanine labeling in plasma and muscle

Plasma insulin concentrations were not significantly different between the corn oil and omega-3 fatty acid supplementation groups, either before or after supplementation during both basal, postabsorptive conditions (before:  $6.3 \pm 1.1$  compared with  $6.1 \pm$

**TABLE 1**

Subject characteristics and basic metabolic variables before and after 8 wk of corn oil or omega-3 fatty acid supplementation<sup>1</sup>

	Corn oil ( <i>n</i> = 7)		Omega-3 fatty acids ( <i>n</i> = 8)		<i>P</i> value <sup>2</sup>		
	Before	After	Before	After	Group	Time	Interaction
Sex (M/F)	5/2	—	5/3	—	—	—	—
Age (y)	71 $\pm$ 2 <sup>3</sup>	—	71 $\pm$ 1	—	0.89 <sup>d</sup>	—	—
BMI (kg/m <sup>2</sup> )	25.7 $\pm$ 1.6	—	25.6 $\pm$ 1.0	—	0.95 <sup>d</sup>	—	—
Body mass (kg) <sup>5</sup>	77.3 $\pm$ 6.2	—	74.2 $\pm$ 4.6	—	0.69 <sup>d</sup>	—	—
Fat-free mass (kg) <sup>5</sup>	54.3 $\pm$ 4.7	—	52.5 $\pm$ 4.4	—	0.78 <sup>d</sup>	—	—
Appendicular MM (kg) <sup>5</sup>	21.7 $\pm$ 2.2	—	21.3 $\pm$ 2.1	—	0.90 <sup>d</sup>	—	—
Systolic BP (mm Hg)	133 $\pm$ 7	133 $\pm$ 5	121 $\pm$ 6	125 $\pm$ 5	0.17	0.59	0.73
Diastolic BP (mm Hg)	79 $\pm$ 2 <sup>6</sup>	75 $\pm$ 3	70 $\pm$ 3	70 $\pm$ 3	0.03	0.44	0.47
HOMA-IR	1.54 $\pm$ 0.34	1.41 $\pm$ 0.40	1.36 $\pm$ 0.23	1.33 $\pm$ 0.16	0.75	0.38	0.59
Total plasma TG (mg/dL)	72 (61, 80) <sup>7</sup>	84 (67, 1047)	81 (65, 94)	84 (73, 87)	0.47	0.52	0.87
Total plasma C (mg/dL)	169 $\pm$ 12	169 $\pm$ 11	172 $\pm$ 7	183 $\pm$ 8	0.53	0.07	0.09
LDL-C (mg/dL)	95 $\pm$ 11	100 $\pm$ 11	103 $\pm$ 8	116 $\pm$ 8	0.39	0.04	0.32
HDL-C (mg/dL)	52 $\pm$ 5	52 $\pm$ 5	44 $\pm$ 4	49 $\pm$ 4	0.42	0.19	0.14
CRP (mg/L)	1.28 (0.90, 1.45)	1.21 (0.61, 2.74)	0.71 (0.38, 1.00)	0.67 (0.24, 3.44)	0.36	0.99	0.72
TNF- $\alpha$ (pg/mL)	0.67 $\pm$ 0.15	0.86 $\pm$ 0.21	0.50 $\pm$ 0.13	0.53 $\pm$ 0.13	0.26	0.03	0.09
IL-6 (pg/mL)	2.05 (0.88, 7.02)	2.43 (1.83, 7.02)	1.29 (1.02, 2.74)	1.82 (1.35, 2.34)	0.34	0.19	0.61

<sup>1</sup> HOMA-IR, homeostasis model assessment of insulin resistance; BP, blood pressure; TG, triglycerides; LDL-C, LDL cholesterol; HDL-C, HDL cholesterol; MM, muscle mass; TNF- $\alpha$ , tumor necrosis factor- $\alpha$ ; IL-6, interleukin-6; CRP, C-reactive protein; C, cholesterol.

<sup>2</sup> *P* values determined by ANOVA with group (corn oil compared with omega-3 fatty acids) and time (before compared with after intervention) as factors after initial comparison of "before" values only by using Student's *t* or the Mann-Whitney *U* tests as appropriate.

<sup>3</sup> Mean  $\pm$  SEM (all such values).

<sup>4</sup> Determined by using Student's *t* test for independent samples.

<sup>5</sup> Measured by dual-energy X-ray absorptiometry as described in Subjects and Methods.

<sup>6</sup> Significantly different from the corresponding "before" value in the omega-3 fatty acid group,  $P = 0.04$  (initial Student's *t* test).

<sup>7</sup> Median; quartile in parentheses (all such values).

1.0  $\mu\text{U/mL}$ , respectively; after:  $6.0 \pm 1.5$  compared with  $5.6 \pm 0.6$   $\mu\text{U/mL}$ , respectively) and during the hyperinsulinemic-hyperaminoacidemic clamp (before:  $26.4 \pm 1.9$  compared with  $31.7 \pm 3.1$   $\mu\text{U/mL}$ , respectively; after:  $28.1 \pm 2.1$  compared with  $29.5 \pm 2.2$   $\mu\text{U/mL}$ , respectively). ANOVA showed a significant effect of clamp ( $P < 0.001$ ) but no significant effect of time or group  $\times$  time interaction.

No differences between the corn oil and omega-3 fatty acid groups in baseline (before intervention) plasma concentrations of glucose and amino acids (all  $P \geq 0.17$ ; **Tables 3** and **4**) were observed. During the clamp, plasma glucose was successfully maintained at  $\approx 5.5$  mmol/L both before and after corn oil and omega-3 fatty acid supplementation; plasma phenylalanine and leucine concentrations rose by  $\approx 80\%$  and  $45\%$  above basal values, respectively (all  $P < 0.001$ ). Omega-3 fatty acid supplementation led to a very small ( $\approx 6\%$ ) but significant ( $P < 0.05$ ) increase in basal plasma glucose concentration (Tables 3 and 4). Plasma phenylalanine and leucine concentrations were not affected by either corn oil or omega-3 fatty acid supplementation during basal, postabsorptive conditions or during the hyperinsulinemic-hyperaminoacidemic clamp (Tables 3 and 4).

Plasma phenylalanine enrichment between 60 and 420 min and plasma glucose enrichment during the last 30 min of the basal state and the clamp before and after corn oil and omega-3 fatty acid supplementation are shown in Tables 3 and 4. The muscle free phenylalanine enrichments (TTR) before and after corn oil supplementation were  $0.0612 \pm 0.0076$  and  $0.0619 \pm 0.0054$ , respectively, at the end of the basal, postabsorptive period and  $0.0748 \pm 0.0037$  and  $0.0739 \pm 0.0031$ , respectively, at the end

of the hyperinsulinemic-hyperaminoacidemic clamp; in the omega-3 fatty acid supplementation group, the respective values were  $0.0679 \pm 0.0030$  and  $0.0652 \pm 0.0021$  (at the end of the basal, postabsorptive period) and  $0.0797 \pm 0.0039$  and  $0.0762 \pm 0.0029$  (at the end of the hyperinsulinemic-hyperaminoacidemic clamp).

#### Muscle protein synthesis rate at baseline (before intervention)

The muscle protein synthesis rate at baseline was not different between the omega-3 fatty acid and corn oil groups, either during basal conditions ( $0.051 \pm 0.005\%$  compared with  $0.051 \pm 0.008\%/h$ , respectively with muscle free phenylalanine as the precursor enrichment and  $0.036 \pm 0.003\%$  compared with  $0.029 \pm 0.005\%/h$ , respectively, with plasma phenylalanine as the precursor enrichment) or during hyperinsulinemia-hyperaminoacidemia ( $0.060 \pm 0.007\%$  compared with  $0.063 \pm 0.009\%/h$ , respectively, with muscle free phenylalanine as the precursor enrichment and  $0.051 \pm 0.006\%$  compared with  $0.047 \pm 0.007\%/h$ , respectively with plasma phenylalanine as the precursor enrichment). ANOVA showed a significant effect of clamp ( $P < 0.001$ ) but no significant effect of group ( $P \geq 0.41$ ) and no interaction between the 2 ( $P \geq 0.60$ ).

To confirm the existence of anabolic resistance in our older adults, we compared these baseline data with values we recently published (27) for healthy, nonobese, young and middle-aged adults who completed the same muscle protein metabolism study and experienced similar increases in plasma insulin and amino

**TABLE 2**

Muscle phospholipid fatty acid profile before and after 8 wk of corn oil or omega-3 fatty acid supplementation<sup>1</sup>

	Corn oil (n = 7)		Omega-3 fatty acids (n = 8)	
	Before	After	Before	After
Saturated fatty acids (% of fatty acids)				
14:0	0.38 $\pm$ 0.02	0.35 $\pm$ 0.04	0.40 $\pm$ 0.09	0.32 $\pm$ 0.03
16:0	17.94 $\pm$ 2.14	17.16 $\pm$ 1.74	19.80 $\pm$ 1.72	18.53 $\pm$ 1.33
18:0	19.97 $\pm$ 0.82	19.14 $\pm$ 0.63	24.07 $\pm$ 1.79	21.28 $\pm$ 0.46
Total	38.24 $\pm$ 2.79	36.61 $\pm$ 2.11	44.17 $\pm$ 3.05	40.05 $\pm$ 1.49
Monounsaturated fatty acids (% of fatty acids)				
16:1n-7	0.35 $\pm$ 0.05	0.37 $\pm$ 0.10	0.34 $\pm$ 0.09	0.50 $\pm$ 0.09
18:1n-9	8.88 $\pm$ 0.55	10.71 $\pm$ 1.21	9.43 $\pm$ 0.54	8.77 $\pm$ 0.47
Total	9.23 $\pm$ 0.59	11.08 $\pm$ 1.24	9.77 $\pm$ 0.56	9.27 $\pm$ 0.50
Omega-6 polyunsaturated fatty acids (% of fatty acids)				
18:2n-6	30.72 $\pm$ 2.45	30.84 $\pm$ 2.50	25.56 $\pm$ 1.45	25.92 $\pm$ 0.49
20:3n-6	1.18 $\pm$ 0.12	1.31 $\pm$ 0.12	1.23 $\pm$ 0.12	1.30 $\pm$ 0.09
20:4n-6	16.11 $\pm$ 1.32	16.02 $\pm$ 0.98	14.24 $\pm$ 1.14	14.44 $\pm$ 0.85
Total	48.02 $\pm$ 3.12 <sup>2</sup>	48.17 $\pm$ 2.70 <sup>2</sup>	41.03 $\pm$ 2.41	41.65 $\pm$ 0.91
Omega-3 polyunsaturated fatty acids (% of fatty acids)				
20:5n-3	0.81 $\pm$ 0.10	0.84 $\pm$ 0.10	0.77 $\pm$ 0.15	2.41 $\pm$ 0.34
22:5n-3	1.62 $\pm$ 0.15	1.50 $\pm$ 0.13	2.23 $\pm$ 0.29	2.51 $\pm$ 0.27
22:6n-3	2.08 $\pm$ 0.25	1.79 $\pm$ 0.20	2.03 $\pm$ 0.28	4.11 $\pm$ 0.41
Total <sup>3</sup>	4.52 $\pm$ 0.37	4.14 $\pm$ 0.31	5.04 $\pm$ 0.45	9.03 $\pm$ 0.95 <sup>4,5</sup>

<sup>1</sup> All values are means  $\pm$  SEMs. The percentage of fatty acids was determined by dividing the peak area of each fatty acid by the sum of all peak areas after gas chromatographic analyses. There were no significant between-group differences in the "before" values,  $P \geq 0.10$  (Student's *t* tests).

<sup>2</sup> Significant group (corn oil compared with omega-3 fatty acids) effect,  $P = 0.05$  (ANOVA).

<sup>3</sup> Significant group  $\times$  time interaction ( $P < 0.05$ , ANOVA), which was followed by Tukey's post hoc analysis.

<sup>4</sup> Significantly different from corresponding value before intervention,  $P < 0.01$  (ANOVA).

<sup>5</sup> Significantly different from the corresponding value in the corn oil group,  $P < 0.01$  (ANOVA).

**TABLE 3**  
Plasma phenylalanine, leucine, and glucose concentrations and phenylalanine and glucose tracer-to-tracee ratios (TTR) during basal, postabsorptive conditions and during the hyperinsulinemic-hyperaminoacidemic clamp before and after corn oil supplementation ( $n = 7$ )<sup>1</sup>

Time	Concentration						Enrichment (TTR)					
	Phenylalanine		Leucine		Glucose		Phenylalanine		Glucose		Glucose	
	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After
Basal conditions	<i>nmol/L</i>											
60 min	65 ± 6	68 ± 9	120 ± 7	120 ± 9	5.31 ± 0.17	5.26 ± 0.16	0.0928 ± 0.0086	0.0928 ± 0.0079	—	—	—	—
90 min	65 ± 8	58 ± 5	117 ± 9	110 ± 8	—	—	0.1019 ± 0.0097	0.0962 ± 0.0091	—	—	—	—
120 min	58 ± 8	65 ± 10	99 ± 6	122 ± 16	5.35 ± 0.20	5.18 ± 0.12	0.1128 ± 0.0120	0.0987 ± 0.0081	—	—	—	—
150 min	64 ± 8	62 ± 10	114 ± 10	111 ± 14	—	—	0.1131 ± 0.0087	0.1085 ± 0.0097	—	—	—	—
180 min	68 ± 9	69 ± 8	119 ± 8	131 ± 7	5.30 ± 0.19	5.09 ± 0.13	0.1132 ± 0.0069	0.1083 ± 0.0098	—	—	—	—
210 min	69 ± 11	70 ± 4	120 ± 14	122 ± 5	5.30 ± 0.23	5.08 ± 0.14	0.1166 ± 0.0104	0.1023 ± 0.0073	0.0220 ± 0.0014	0.0222 ± 0.0014	0.0222 ± 0.0015	0.0222 ± 0.0015
220 min	—	—	—	—	5.28 ± 0.22	5.08 ± 0.14	—	—	0.0218 ± 0.0015	0.0222 ± 0.0015	0.0222 ± 0.0015	0.0222 ± 0.0015
230 min	—	—	—	—	5.26 ± 0.23	5.07 ± 0.13	—	—	0.0218 ± 0.0017	0.0225 ± 0.0016	0.0225 ± 0.0016	0.0225 ± 0.0016
240 min	74 ± 11	75 ± 12	132 ± 13	137 ± 11	5.28 ± 0.23	5.10 ± 0.14	0.1192 ± 0.0102	0.1090 ± 0.0089	0.0224 ± 0.0019	0.0218 ± 0.0015	0.0218 ± 0.0015	0.0218 ± 0.0015
Mean ± SEM	66 ± 8	66 ± 8	117 ± 8	121 ± 7	5.30 ± 0.21	5.12 ± 0.13	0.1099 ± 0.0091	0.1030 ± 0.0087	0.0220 ± 0.0016	0.0222 ± 0.0015	0.0222 ± 0.0015	0.0222 ± 0.0015
Hyperinsulinemic-hyperaminoacidemic clamp	<i>nmol/L</i>											
270 min	101 ± 13	99 ± 18	184 ± 16	168 ± 11	5.42 ± 0.20	5.47 ± 0.07	0.1206 ± 0.0059	0.1132 ± 0.0110	—	—	—	—
300 min	104 ± 13	118 ± 16	164 ± 11	197 ± 12	5.46 ± 0.19	5.52 ± 0.07	0.1193 ± 0.0075	0.0996 ± 0.0055	—	—	—	—
330 min	113 ± 18	127 ± 20	166 ± 25	196 ± 25	5.64 ± 0.06	5.58 ± 0.07	0.1014 ± 0.0087	0.0996 ± 0.0049	—	—	—	—
360 min	130 ± 17	126 ± 16	181 ± 15	177 ± 13	5.62 ± 0.06	5.60 ± 0.07	0.1019 ± 0.0062	0.0979 ± 0.0055	—	—	—	—
390 min	117 ± 20	111 ± 11	151 ± 18	159 ± 19	5.57 ± 0.07	5.60 ± 0.06	0.0985 ± 0.0087	0.0983 ± 0.0054	0.0249 ± 0.0018	0.0254 ± 0.0012	0.0255 ± 0.0012	0.0255 ± 0.0012
400 min	—	—	—	—	5.55 ± 0.06	5.59 ± 0.05	—	—	0.0247 ± 0.0016	0.0255 ± 0.0017	0.0255 ± 0.0017	0.0255 ± 0.0017
410 min	—	—	—	—	5.55 ± 0.06	5.59 ± 0.08	—	—	0.0255 ± 0.0017	0.0260 ± 0.0011	0.0260 ± 0.0011	0.0260 ± 0.0011
420 min	130 ± 20	118 ± 15	164 ± 18	155 ± 17	5.51 ± 0.06	5.60 ± 0.05	0.1015 ± 0.0057	0.0988 ± 0.0057	0.0253 ± 0.0016	0.0260 ± 0.0011	0.0260 ± 0.0011	0.0260 ± 0.0011
Mean ± SEM	116 ± 14 <sup>2</sup>	117 ± 14 <sup>2</sup>	168 ± 12 <sup>2</sup>	176 ± 13 <sup>2</sup>	5.54 ± 0.08 <sup>2</sup>	5.57 ± 0.04 <sup>2</sup>	0.1054 ± 0.005	0.1007 ± 0.0059	0.0251 ± 0.0016 <sup>2</sup>	0.0256 ± 0.0016 <sup>2</sup>	0.0256 ± 0.0016 <sup>2</sup>	0.0256 ± 0.0016 <sup>2</sup>

<sup>1</sup> All values are means ± SEMs. ANOVA on the "before" values from Tables 3 and 4 with group (corn oil compared with omega-3 fatty acid) and study stage (basal compared with clamp) as the factors showed no significant between-group differences (all  $P \geq 0.10$ ) or group × study stage interactions (all  $P \geq 0.36$ ).

<sup>2</sup> Significantly different from corresponding "basal" value,  $P < 0.001$  [ANOVA on values from the corn oil group presented here with study stage (basal compared with clamp) and time (before compared with after intervention) as factors].

**TABLE 4**  
Plasma phenylalanine, leucine, and glucose concentrations and phenylalanine and glucose tracer-to-trace ratios (TTR) during basal, postabsorptive conditions and during the hyperinsulinemic-hyperaminoacidemic clamp before and after omega-3 fatty acid supplementation ( $n = 8$ )<sup>1</sup>

Time	Concentration <sup>2</sup>										Enrichment (TTR)			
	Phenylalanine		Leucine		Glucose		Phenylalanine		Glucose		Before	After	Before	After
	Before	After	Before	After	Before	After	Before	After	Before	After				
Basal conditions														
60 min	58 ± 10	61 ± 14	112 ± 11	117 ± 7	5.03 ± 0.10	5.29 ± 0.07	0.0853 ± 0.0040	0.0914 ± 0.0093	—	—	—	—	—	—
90 min	63 ± 9	57 ± 9	119 ± 8	111 ± 6	—	—	0.0854 ± 0.0078	0.0968 ± 0.0073	—	—	—	—	—	—
120 min	56 ± 9	58 ± 10	107 ± 8	109 ± 7	5.06 ± 0.10	5.31 ± 0.08	0.0950 ± 0.0065	0.0987 ± 0.0071	—	—	—	—	—	—
150 min	58 ± 9	57 ± 12	112 ± 10	103 ± 9	—	—	0.0962 ± 0.0057	0.0991 ± 0.0084	—	—	—	—	—	—
180 min	54 ± 9	49 ± 5	104 ± 8	107 ± 6	5.06 ± 0.09	5.41 ± 0.10	0.0999 ± 0.0050	0.1086 ± 0.0062	—	—	—	—	—	—
210 min	59 ± 8	52 ± 8	116 ± 8	106 ± 5	4.99 ± 0.10	5.30 ± 0.11	0.1014 ± 0.0047	0.1023 ± 0.0062	—	—	0.0220 ± 0.0008	0.0218 ± 0.0009	0.0220 ± 0.0008	0.0218 ± 0.0009
220 min	—	—	—	—	4.99 ± 0.09	5.31 ± 0.10	—	—	—	—	0.0024 ± 0.0009	0.0218 ± 0.0009	0.0024 ± 0.0009	0.0218 ± 0.0009
230 min	—	—	—	—	4.97 ± 0.09	5.28 ± 0.09	—	—	—	—	0.0225 ± 0.0009	0.0217 ± 0.0009	0.0225 ± 0.0009	0.0217 ± 0.0009
240 min	55 ± 7	50 ± 7	110 ± 9	98 ± 7	4.97 ± 0.10	5.29 ± 0.11	0.1038 ± 0.0069	0.1044 ± 0.0055	—	—	0.0225 ± 0.0009	0.0217 ± 0.0009	0.0225 ± 0.0009	0.0217 ± 0.0009
Mean ± SEM	57 ± 8	56 ± 10	111 ± 7	108 ± 5	5.01 ± 0.09	5.31 ± 0.09 <sup>3</sup>	0.0955 ± 0.0052	0.0994 ± 0.0069	—	—	0.0223 ± 0.0090	0.0217 ± 0.0090	0.0223 ± 0.0090	0.0217 ± 0.0090
Hyperinsulinemic-hyperaminoacidemic clamp														
270 min	89 ± 9	91 ± 12	185 ± 10	183 ± 13	5.09 ± 0.18	5.37 ± 0.14	0.1015 ± 0.0030	0.0988 ± 0.0043	—	—	—	—	—	—
300 min	91 ± 9	95 ± 11	165 ± 9	172 ± 14	5.30 ± 0.17	5.46 ± 0.11	0.0939 ± 0.0024	0.0937 ± 0.0031	—	—	—	—	—	—
330 min	98 ± 13	99 ± 13	159 ± 12	158 ± 12	5.55 ± 0.06	5.48 ± 0.04	0.0917 ± 0.0032	0.0910 ± 0.0049	—	—	—	—	—	—
360 min	113 ± 14	109 ± 20	178 ± 18	160 ± 19	5.62 ± 0.09	5.39 ± 0.10	0.0914 ± 0.0036	0.0881 ± 0.0041	—	—	—	—	—	—
390 min	116 ± 16	108 ± 16	171 ± 10	146 ± 6	5.41 ± 0.07	5.39 ± 0.06	0.0866 ± 0.0076	0.0865 ± 0.0074	—	—	0.0260 ± 0.0005	0.0259 ± 0.0007	0.0260 ± 0.0005	0.0259 ± 0.0007
400 min	—	—	—	—	5.46 ± 0.04	5.47 ± 0.07	—	—	—	—	0.0260 ± 0.0005	0.0264 ± 0.0007	0.0260 ± 0.0005	0.0264 ± 0.0007
410 min	—	—	—	—	5.49 ± 0.06	5.48 ± 0.05	—	—	—	—	0.0264 ± 0.0005	0.0268 ± 0.0008	0.0264 ± 0.0005	0.0268 ± 0.0008
420 min	103 ± 15	101 ± 11	152 ± 13	150 ± 10	5.43 ± 0.04	5.41 ± 0.06	0.0894 ± 0.0052	0.0905 ± 0.0040	—	—	0.0264 ± 0.0005	0.0266 ± 0.0007	0.0264 ± 0.0005	0.0266 ± 0.0007
Mean ± SEM	102 ± 12 <sup>4</sup>	101 ± 13 <sup>4</sup>	168 ± 9 <sup>4</sup>	161 ± 10 <sup>4</sup>	5.42 ± 0.05 <sup>4</sup>	5.41 ± 0.04 <sup>4</sup>	0.0924 ± 0.0037	0.0914 ± 0.0039	—	—	0.0262 ± 0.0050 <sup>4</sup>	0.0264 ± 0.0070 <sup>4</sup>	0.0262 ± 0.0050 <sup>4</sup>	0.0264 ± 0.0070 <sup>4</sup>

<sup>1</sup> All values are means ± SEMs. ANOVA on the "before" values from Tables 3 and 4 with group (corn oil compared with omega-3 fatty acid) and study stage (basal compared with clamp) as the factors showed no significant between-group differences (all  $P \geq 0.10$ ) or group × study stage interactions (all  $P \geq 0.36$ ).

<sup>2</sup> With the exception of the change in basal plasma glucose concentration in the omega-3 fatty acid group ( $P < 0.05$  compared with the before-after change in basal plasma glucose concentration in the corn oil group), there were no significant differences in the before-after changes between the corn oil and the omega-3 fatty acid groups.

<sup>3</sup> Significantly different from corresponding "before" value,  $P < 0.05$  [ANOVA on values from the omega-3 fatty acid group presented here with study stage (basal compared with clamp) and time (before compared with after intervention) as factors and Tukey's post hoc analysis].

<sup>4</sup> Significantly different from corresponding basal value,  $P < 0.001$  [ANOVA on values from the omega-3 fatty acid group presented here with study stage (basal compared with clamp) and time (before compared with after intervention) as factors].

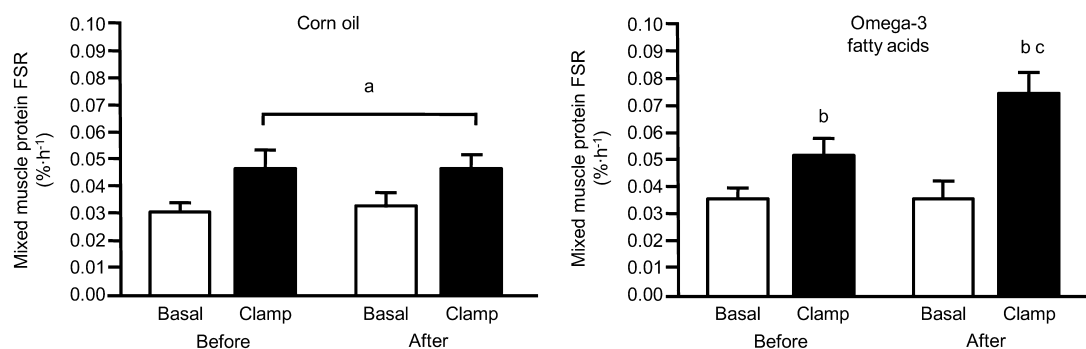
acid concentrations during the clamp as in our older subjects in the present study. Indeed, the hyperaminoacidemia-hyperinsulinemia induced increase in the rate of muscle protein synthesis above basal values was twice as high ( $P < 0.05$ ) in the young ( $0.022 \pm 0.004\%/h$ ) than in the older ( $0.011 \pm 0.003\%/h$ ) subjects (with muscle free phenylalanine as the precursor enrichment).

#### Effect of intervention on the muscle protein synthesis rate (plasma phenylalanine labeling as precursor pool enrichment)

Corn oil supplementation had no effect on either the basal muscle protein synthesis rate or the increase in the rate of muscle protein synthesis in response to amino acid and insulin infusion (change from basal values:  $0.019 \pm 0.004\%/h$  before and  $0.016 \pm 0.005\%/h$  after supplementation;  $P = 0.78$ ; **Figure 1**). In contrast, omega-3 fatty acid supplementation had no effect on the basal rate of muscle protein synthesis but approximately doubled the anabolic response to amino acid and insulin infusion (change from basal values:  $0.015 \pm 0.004\%$  compared with  $0.036 \pm 0.005\%/h$ ;  $P < 0.001$ ), and the before-after change in the anabolic response in the omega-3 fatty acid group was significantly greater than in the corn oil group ( $P = 0.01$ ).

#### Effect of intervention on the muscle protein synthesis rate (muscle free phenylalanine labeling as precursor pool enrichment)

Corn oil supplementation had no effect on either the basal muscle protein synthesis rate or the increase in the rate of muscle protein synthesis in response to amino acid and insulin infusion (change from basal values:  $0.013 \pm 0.005\%/h$  before and  $0.013 \pm 0.007\%/h$  after supplementation;  $P = 0.94$ ). In contrast, omega-3 fatty acid supplementation had no effect on the basal rate of muscle protein synthesis but significantly increased the anabolic response to amino acid and insulin infusion (change from basal values:  $0.031 \pm 0.003\%$  compared with  $0.009 \pm 0.005\%/h$ ;  $P = 0.004$ ).



**FIGURE 1.** Mean ( $\pm$ SEM) mixed skeletal muscle protein fractional synthesis rate (FSR), calculated by using the average plasma free phenylalanine enrichment as the precursor pool enrichment, during basal, postabsorptive conditions and during the hyperaminoacidemic-hyperinsulinemic clamp before and after 8 wk of supplementation with either corn oil ( $n = 7$ ) or omega-3 fatty acids ( $n = 8$ ). There was no difference in the muscle protein FSR between the omega-3 fatty acid and corn oil groups before the intervention [ANOVA showed a significant effect of clamp ( $P < 0.001$ ), no significant effect of group ( $P = 0.47$ ), and no interaction ( $P = 0.60$ )]. <sup>a</sup>In the corn oil group, ANOVA showed a significant main effect of clamp ( $P < 0.01$ ). In the omega-3 fatty acid group, ANOVA showed a significant effect of clamp ( $P < 0.01$ ) and an interaction ( $P < 0.001$ ), which was followed by Tukey's post hoc analysis. <sup>b</sup>Significantly different from the corresponding basal value,  $P < 0.01$ . <sup>c</sup>Significantly different from the corresponding value before omega-3 fatty acid supplementation,  $P < 0.01$ . Furthermore, the before-after intervention change in the anabolic response (increase in the muscle protein FSR from basal values) was significantly greater in the omega-3 fatty acid group than in the corn oil group ( $P = 0.01$ , Student's  $t$  test for independent samples).

#### Phosphorylation of signaling transduction proteins in muscle

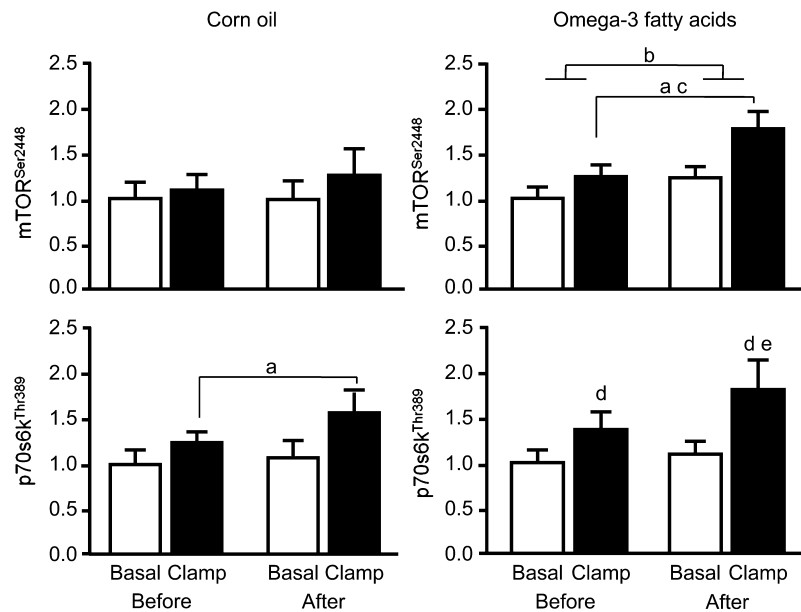
At baseline (before supplementation), no differences between the omega-3 fatty acid and corn oil groups were observed in the basal, postabsorptive concentrations of Akt<sup>Thr308</sup>, mTOR<sup>Ser2448</sup>, and p70s6k<sup>Thr389</sup> in muscle and the increase in Akt<sup>Thr308</sup>, mTOR<sup>Ser2448</sup>, and p70s6k<sup>Thr389</sup> phosphorylation in response to hyperaminoacidemia-hyperinsulinemia. For each of these, ANOVA showed a significant effect of clamp ( $P \leq 0.01$ ), no significant effect of group ( $P \geq 0.45$ ), and no interaction between the 2 ( $P \geq 0.17$ ).

Neither corn oil nor omega-3 fatty acid supplementation affected the concentration of Akt<sup>Thr308</sup> in muscle ( $P \geq 0.28$ ). The concentrations of Akt<sup>Thr308</sup> during basal conditions and the clamp were as follows: 0.96 (0.51, 1.32) and 1.37 (1.27, 1.90) before and 0.94 (0.48, 1.74) and 1.41 (0.64, 1.78) after corn oil supplementation and 0.94 (0.62, 1.42) and 1.31 (1.09, 1.57) before and 0.96 (0.61, 1.27) and 1.46 (1.19, 2.07) after omega-3 fatty acid supplementation. Corn oil supplementation also had no effect on the concentrations of mTOR<sup>Ser2448</sup> and p70s6k<sup>Thr389</sup> in muscle (**Figure 2**). In contrast, the concentrations of mTOR<sup>Ser2448</sup> and p70s6k<sup>Thr389</sup> during the clamp were greater after than before omega-3 fatty acid supplementation ( $P < 0.05$  for p70s6k<sup>Thr389</sup>;  $P = 0.08$  for mTOR<sup>Ser2448</sup>) (Figure 2), and the before-after changes in the insulin/amino acid-mediated increase in p70s6k and mTOR phosphorylation above basal values were greater in the omega-3 fatty acid group than in the corn oil group ( $P < 0.05$  and  $P = 0.07$ , respectively).

#### Whole-body glucose kinetics and phenylalanine rate of appearance

Whole-body glucose kinetics were not significantly different between the corn oil and omega-3 fatty acid groups at baseline and were not affected by either omega-3 fatty acid or corn oil supplementation (all  $P \geq 0.3$ ). Basal glucose Ra was  $14.8 \pm 1.0 \mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$  before and  $15.0 \pm 1.1 \mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$  after omega-3 fatty acid supplementation and  $14.4 \pm 1.2 \mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$  before and  $14.5 \pm$





**FIGURE 2.** Mean ( $\pm$ SEM) concentrations (arbitrary units) of mTOR<sup>Ser2448</sup> and p70s6k<sup>Thr389</sup> during basal, postabsorptive conditions and during the hyperaminoacidemic-hyperinsulinemic clamp before and after 8 wk of supplementation with either corn oil ( $n = 7$ ) or omega-3 fatty acids ( $n = 8$ ). <sup>a</sup>ANOVA showed a significant main effect of clamp ( $P < 0.01$ ). <sup>b</sup>ANOVA showed a significant main effect of time ( $P < 0.05$ ). <sup>c</sup>There was a trend for a greater clamp-induced increase in mTOR<sup>Ser2448</sup> after omega-3 fatty acid supplementation than before supplementation (interaction:  $P = 0.08$ ). <sup>d,e</sup>ANOVA showed a significant interaction ( $P < 0.05$ ), which was followed by Tukey's post hoc analysis. <sup>d</sup>Significantly different from corresponding basal value,  $P < 0.05$ . <sup>e</sup>Significantly different from corresponding value before omega-3 fatty acid supplementation,  $P < 0.05$ . Furthermore, the before-after intervention changes in the insulin/amino acid-mediated increase in p70s6k and mTOR phosphorylation above basal values were greater in the omega-3 fatty acid group than in the corn oil group ( $P < 0.05$  and  $P = 0.07$ , respectively; Mann-Whitney  $U$  test).

$1.3 \mu\text{mol} \cdot \text{kg FFM}^{-1} \cdot \text{min}^{-1}$  after corn oil supplementation. During the clamp, endogenous glucose Ra decreased by  $77 \pm 5\%$  before and  $75 \pm 5\%$  after omega-3 fatty acid supplementation, and by  $71 \pm 6\%$  before and  $68 \pm 4\%$  after corn oil supplementation [no significant difference between groups ( $P = 0.34$ ) and no significant effect of time ( $P = 0.44$ )]. Glucose Rd increased by  $137 \pm 24\%$  before and  $113 \pm 13\%$  after omega-3 fatty acid supplementation and by  $113 \pm 24\%$  before and  $109 \pm 28\%$  after corn oil supplementation [no significant difference between groups ( $P = 0.61$ ) and no significant effect of time ( $P = 0.43$ )].

Whole-body phenylalanine Ra was also not different between the corn oil and omega-3 fatty acid groups at baseline and was not affected by either omega-3 fatty acid or corn oil supplementation (all  $P > 0.3$ ). In the corn oil group, endogenous phenylalanine Ra was  $41.3 \pm 3.6 \mu\text{mol}/\text{min}$  before and  $43.8 \pm 4.0 \mu\text{mol}/\text{min}$  after supplementation during basal postabsorptive conditions and  $24.6 \pm 3.6 \mu\text{mol}/\text{min}$  before and  $26.3 \pm 4.3 \mu\text{mol}/\text{min}$  after supplementation during the clamp. ANOVA showed a main effect of clamp ( $P < 0.001$ ), no effect of time ( $P = 0.31$ ), and no clamp  $\times$  time interaction ( $P = 0.57$ ). In the omega-3 fatty acid group, the respective values were  $42.1 \pm 3.7$  and  $41.2 \pm 2.9 \mu\text{mol}/\text{min}$  in the basal postabsorptive state and  $30.8 \pm 4.9$  and  $31.3 \pm 4.7 \mu\text{mol}/\text{min}$  during the clamp. ANOVA showed a main effect of clamp ( $P = 0.005$ ), no effect of time ( $P = 0.84$ ), and no clamp  $\times$  time interaction ( $P = 0.50$ ).

## DISCUSSION

The normal age-related loss of muscle mass is considered to be largely due to anabolic resistance (ie, an inadequate response to anabolic stimuli such as increased amino acid availability) (5–7).

In the present study, we provide novel evidence that dietary omega-3 fatty acid supplementation augments the hyperaminoacidemia-hyperinsulinemia induced increase in the rate of muscle protein synthesis in older adults. Omega-3 fatty acids therefore probably attenuate the anabolic resistance and may potentially be useful as a therapeutic agent to treat sarcopenia.

The exact mechanisms by which omega-3 fatty acids act on the muscle protein synthesis process are not entirely clear, but our results indicate that it was at least partially mediated via increased activation of the mTOR-p70s6k signaling pathway. The mTOR-p70s6k signaling pathway is considered an integral control point for muscle cell growth (30–32), and Drummond et al recently showed that rapamycin, a potent inhibitor of mTOR, prevents the increase in the rate of muscle protein synthesis in response to feeding (33) and resistance exercise (34) in human subjects. Although we cannot determine the mechanism by which omega-3 fatty acids induced greater mTOR-p70s6k activation in response to increased amino acid and insulin supply, we can probably rule out Akt as the upstream signal because omega-3 fatty acid supplementation had no effect on Akt phosphorylation in our study.

It is thought that many of the well-known beneficial effects of omega-3 fatty acids (eg, reduction in plasma triglyceride concentration) are, at least partially, related to the antiinflammatory properties of omega-3 fatty acids (13, 35, 36). However, we were unable to discover significant differences in serum markers of inflammation and plasma triglyceride concentrations in response to omega-3 fatty acid supplementation, probably because we specifically selected healthy persons with low plasma concentrations of inflammatory markers and triglycerides for this study. Nevertheless, the fact that omega-3 fatty acids had anabolic effects in the absence of significant inflammatory activity or



changes thereof indicates that the antiinflammatory properties of omega-3 fatty acids are likely not responsible for the muscle anabolic action.

The improvement in muscle protein metabolism was not accompanied by significant changes in whole-body glucose metabolism, which is thought to be responsive to omega-3 fatty acid intake (37–40). This is not entirely surprising, however. Although results from studies in animals indicate consistently that omega-3 fatty acids stimulate muscle glucose uptake (37–40), the results from studies in human subjects concerning the insulin-sensitizing effect of omega-3 fatty acids on glucose metabolism are less clear-cut, with some showing a positive effect, some no effect, and some an adverse effect (35, 41–43). In our study we consider it possible that increased activation of mTOR by omega-3 fatty acids during hyperaminoacidemia-hyperinsulinemia may have prevented a potentially beneficial effect of omega-3 fatty acids on glucose uptake because there is some evidence for an amino acid induced decrease in insulin-mediated glucose transport via activation of the mTOR signaling pathway (44, 45).

In summary, we have shown that dietary omega-3 fatty acid, but not corn oil, supplementation increases muscle anabolic signaling activity and the insulin/amino acid-mediated increase in muscle protein synthesis above basal, postabsorptive values in older adults. Although the exact mechanisms by which omega-3 fatty acids stimulate muscle protein synthesis during hyperinsulinemia-hyperaminoacidemia remain to be resolved, our study provides compelling evidence of an interaction of omega-3 fatty acids and protein metabolism in human muscle and suggest that dietary omega-3 fatty acid supplementation could potentially provide a safe, simple, and low-cost intervention to combat sarcopenia.

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The authors' responsibilities were as follows—GIS: involved in conducting the study, processing the study samples, collecting the data, performing the final data analyses, and writing the manuscript; PA and MJR: involved in processing the study samples, collecting the data, and writing the manuscript; DNR and BSM: involved in conducting the study; DR: involved in sample processing and sample analyses; and BM: involved in designing and conducting the study, processing the study samples, collecting the data, performing the final data analyses, and writing the manuscript. None of the authors had any potential conflicts of interest. No funding entity had any role in the design, implementation, analysis, or interpretation of the data.

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