A high proportion of leucine is required for optimal stimulation of the rate of muscle protein synthesis by essential amino acids in the elderly

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Submitted 6 October 2005; accepted in final form 17 February 2006

Katsanos, Christos S., Hisamine Kobayashi, Melinda Sheffield-Moore, Asle Aarsland, and Robert R. Wolfe. A high proportion of leucine is required for optimal stimulation of the rate of muscle protein synthesis by essential amino acids in the elderly. Am J Physiol Endocrinol Metab 291:E381–E387, 2006—This study was designed to evaluate the effects of enriching an essential amino acid (EAA) mixture with leucine on muscle protein metabolism in elderly and young individuals. Four (2 elderly and 2 young) groups were studied before and after ingestion of 6.7 g of EAs. EAs were based on the composition of whey protein (26% leucine [26% Leu]) or were enriched in leucine (41% leucine [41% Leu]). A primed, continuous infusion of 1-[ring-2H5]phenylalanine was used together with vastus lateralis muscle biopsies and leg arteriovenous blood samples for the determinations of fractional synthetic rate (FSR) and balance of muscle protein. FSR increased following amino acid ingestion in both the 26% (basal: 0.048 ± 0.005%/h; post-EAA: 0.050 ± 0.007%/h) and the 41% (basal: 0.032 ± 0.003%/h, post-EAA: 0.051 ± 0.007%/h) Leu young groups (P < 0.05). In contrast, in the elderly, FSR did not increase following ingestion of 26% Leu EAA (basal: 0.044 ± 0.003%/h; post-EAA: 0.049 ± 0.006%/h; P > 0.05) but did increase following ingestion of 41% Leu EAA (basal: 0.038 ± 0.007%/h; post-EAA: 0.056 ± 0.008%/h; P < 0.05). Similar to the FSR responses, the mean response of muscle phenylalanine net balance, a reflection of muscle protein balance, was improved (P < 0.05) in all groups, with the exception of the 26% Leu elderly group. We conclude that increasing the proportion of leucine in a mixture of EAA can reverse an attenuated response of muscle protein synthesis in elderly but does not result in further stimulation of muscle protein synthesis in young subjects.

Ingestion of an amino acid mixture containing extra leucine has the potential to affect muscle protein metabolism in several ways. In addition to providing leucine and other amino acids as precursors for protein synthesis, the extra leucine may stimulate specific intracellular pathways associated with muscle protein synthesis. Specifically, there is evidence implicating a leucine-mediated increase in plasma insulin, resulting in a regulation of the ribosomal protein S6 kinase (S6K1) and the eukaryotic initiation factor (eIF)4E-binding protein-1 (1), which are involved in the initiation of muscle protein synthesis. There is also evidence suggesting that plasma leucine can regulate muscle protein synthesis by insulin-independent mechanisms (2).

The importance of an improved response of skeletal muscle protein synthesis to the ingestion of amino acids is obvious for individuals across the age spectrum, and particularly the elderly, because skeletal muscle mass declines with advancing age (22). We have recently shown that elderly are less responsive than young individuals to the ingestion of a small bolus of EAA (20). Ingestion of extra leucine may be particularly important for the stimulation of skeletal muscle protein synthesis in the elderly, because evidence from animal studies indicates that skeletal muscle protein synthesis becomes less responsive to the stimulatory effects of leucine with aging (8). Additional evidence indicates that meals supplemented with leucine improve the postprandial muscle protein synthesis in old rats (29).

The purpose of this study was to determine the acute effects of two different EAA mixtures on skeletal muscle protein metabolism in elderly and young subjects, a mixture that is based on the distribution of EAAs in whey protein (26% leucine) and a similar mixture that is enriched in leucine (41% leucine). We determined muscle protein synthesis by calculating the incorporation rate of 1-[ring-2H5]phenylalanine in the skeletal muscle protein pool and muscle protein retention by measuring the leg arteriovenous net balance of phenylalanine.

METHODS

Subjects. Elderly subjects were recruited through the Sealy Center on Aging Volunteers Registry of the University of Texas Medical Branch at Galveston. Young subjects were recruited through newspaper advertisements. Subject characteristics are presented in Table 1. Leg volume was determined using an anthropometric method (15, 18).

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Table 1. Subject characteristics

<table>
<thead>
<tr>
<th>Age, yr</th>
<th>Weight, kg</th>
<th>Height, cm</th>
<th>Body fat, %</th>
<th>Leg volume, liters</th>
<th>LLM, kg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Elderly</td>
<td>66.7±2.0</td>
<td>81.7±3.6</td>
<td>171.5±5.2</td>
<td>31.3±2.8</td>
<td>8.2±0.5</td>
</tr>
<tr>
<td>Young</td>
<td>26% Leu</td>
<td>41% Leu</td>
<td>26% Leu</td>
<td>41% Leu</td>
<td>26% Leu</td>
</tr>
<tr>
<td>3 F, 7 M</td>
<td>66.5±2.2</td>
<td>74.5±4.7</td>
<td>165±2.3</td>
<td>30.8±2.7</td>
<td>7.8±0.8</td>
</tr>
<tr>
<td>5 F, 5 M</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20% Leu</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6 F, 4 M</td>
<td>70±4.7</td>
<td>76.6±7.7</td>
<td>170±2.3</td>
<td>25.2±3.3</td>
<td>8.0±0.6</td>
</tr>
<tr>
<td>165±2.0</td>
<td></td>
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<td></td>
<td></td>
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<tr>
<td>20% Leu</td>
<td></td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>6 F, 4 M</td>
<td>28.8±2.6*</td>
<td>11.7±1.0</td>
<td>29.8±2.0</td>
<td>8.5±1.1</td>
<td></td>
</tr>
<tr>
<td>165±2.0</td>
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</tbody>
</table>

Values are means ± SE. F, females; M, males; LLM, leg lean mass; 26% (26 Leu) and 41% (41 Leu) leucine in 6.7 g of essential amino acids. *Significantly different from the elderly groups (P < 0.05).

Leg lean mass and percentage of body fat were determined using dual-energy X-ray absorptiometry (DEXA). Subjects in the same age category were randomly assigned into two groups. Subjects in one group ingested 6.7 g of EAAAs containing 1.7 g of leucine (26% Leu; percentage of leucine found in whey protein), whereas subjects in the other group ingested 6.7 g of EAAAs containing 2.8 g of leucine (41% Leu). The latter EAA mixture was developed with the purpose of avoiding substantially decreasing the availability of the other EAA while increasing the proportion of leucine (Table 2). Amino acids were dissolved in 250 ml of a noncaloric/noncaffeinated soft drink.

Subjects were determined to be healthy on the basis of medical history, physical examination, resting electrocardiogram (ECG), and determination of the leg blood flow. All subjects that qualified for the study were instructed to eat their usual diet for the week before the study and avoid substantially decreasing the availability of the other EAA as well as increasing the proportion of leucine (Table 2). Amino acids were dissolved in 250 ml of a noncaloric/noncaffeinated soft drink.

Table 2. Composition of the essential amino acid mixtures (in g)

<table>
<thead>
<tr>
<th>E382 LEUCINE AND MUSCLE PROTEIN METABOLISM</th>
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<table>
<thead>
<tr>
<th>Essential Amino Acid</th>
<th>26% Leu</th>
<th>41% Leu</th>
</tr>
</thead>
<tbody>
<tr>
<td>Histidine</td>
<td>0.304</td>
<td>0.239</td>
</tr>
<tr>
<td>Isoleucine</td>
<td>0.781</td>
<td>0.614</td>
</tr>
<tr>
<td>Leucine</td>
<td>1.721</td>
<td>2.790</td>
</tr>
<tr>
<td>Lysine</td>
<td>1.360</td>
<td>1.069</td>
</tr>
<tr>
<td>Methionine</td>
<td>0.362</td>
<td>0.284</td>
</tr>
<tr>
<td>Phenylalanine</td>
<td>0.506</td>
<td>0.398</td>
</tr>
<tr>
<td>Threonine</td>
<td>0.955</td>
<td>0.751</td>
</tr>
<tr>
<td>Valine</td>
<td>0.738</td>
<td>0.580</td>
</tr>
<tr>
<td>Total</td>
<td>6.726</td>
<td>6.726</td>
</tr>
</tbody>
</table>

The 26% Leu mixture was based on the amounts of essential amino acids in 15 g of whey protein.
place the tubes in a heating block (110°C) for 24 h. The hydrolysate was passed over a cation exchange column (AG 50W-8X 200–400 mesh H⁺ form cation resin; Bio-Rad Laboratories, Hercules, CA) and then processed the same way as the blood. Muscle protein-bound phenylalanine enrichment was determined on GC-MS using selected ion monitoring (m/z 234, 237, and 239) and the standard curve method (7, 26).

**Calculations.** The concentrations of phenylalanine and leucine in the blood were determined on the basis of the weight of the blood sample, the amount of internal standard added, and the t/T for L-[U-13C9-15N]phenylalanine and L-[U-13C6]leucine, respectively (4).

The muscle free phenylalanine was determined similarly, and the intracellular concentration was calculated using the chloride method (3). The net balance (NB) of phenylalanine (NBPhe) across the leg at each time point was calculated as follows:

$$NB = (C_a - C_v) \times BF$$

where $C_a$ and $C_v$ are the phenylalanine concentrations in the femoral artery and vein, respectively, and BF is the leg blood flow. The NBPhe was determined for the basal period and the post-EAA period by calculating the average NBPhe during the respective period (19). At any given period where the intracellular free phenylalanine concentration remains constant, the rates of phenylalanine disappearance from the artery ($R_d$) and appearance to the vein ($R_a$) reflect the rates of incorporation of blood phenylalanine into muscle proteins ($R_d$) and release from muscle proteins breakdown ($R_a$), because phenylalanine is not metabolized in muscle (35). These parameters were calculated by using the following equations:

$$R_d = [(E_v \times C_a) - (E_a \times C_v)] \times BF/E_a$$

$$R_a = R_d - NB$$

where $E_a$ and $E_v$ are the blood phenylalanine enrichments, expressed as $t/T$ in the femoral artery and vein, respectively. $R_d$ and $R_a$ values during the basal and the post-EAA periods were averaged to calculate mean $R_d$ and $R_a$ responses during the respective period. The fractional synthetic rate (FSR, %/h) of mixed-muscle protein was calculated as follows (31):

$$FSR = \frac{\Delta E_p}{E_a \times t} \times 60 \times 100$$

where $\Delta E_p$ defines the increment in the muscle protein-bound phenylalanine $t/T$ between two biopsies, $E_a$ is the average arterial phenylalanine $t/T$ during the isotopic steady state between the two biopsies, and $t$ is the time interval (min) between the biopsies. The factors 60 and 100 are used to express the FSR values in percentage per hour.

**Statistical analyses.** One-way analysis of variance (ANOVA) was used to compare subject characteristics across groups. Two-way (group × time) repeated-measures ANOVA was used to compare differences between and within groups. When appropriate, statistically significant F values were followed by Tukey’s tests. All data are expressed as means ± SE, and a P value ≤0.05 was considered statistically significant.

**RESULTS**

**Blood flow.** No differences were found in the blood flow measurements performed before and after the EAA ingestion for either group ($P > 0.05$). For each subject, an average from the two blood flow values measured before and after the EAA ingestion was used to calculate study parameters to reduce variability. The mean leg blood flow in the elderly was 3.5 ± 0.8 ml·min⁻¹·100 ml leg volume⁻¹ (26% Leu) and 3.4 ± 0.4 ml·min⁻¹·100 ml leg volume⁻¹ (41% Leu), whereas in the young it was 3.4 ± 0.3 ml·min⁻¹·100 ml leg volume⁻¹ (26% Leu) and 2.7 ± 0.3 ml·min⁻¹·100 ml leg volume⁻¹ (41% Leu). There were no differences in the mean leg blood flow values between groups ($P > 0.05$).

**Arterial blood leucine and phenylalanine concentrations and phenylalanine enrichment.** Mean arterial blood leucine response to the EAA ingestion in the four groups is presented in Fig. 2. There was a significant time effect for the leucine response ($P < 0.05$), as well as a group effect, with the 41% Leu groups having a higher leucine concentration than the 26% Leu groups ($P < 0.05$). Within groups, arterial blood leucine concentration increased immediately (15 min) and remained significantly ($P < 0.05$) elevated compared with basal for 105 min in the 26% Leu elderly group, 120 min in the 41% Leu elderly group, 90 min in the 26% Leu young group, and 150 min in the 41% Leu young group.
min in the 41% Leu young group. With respect to the arterial blood phenylalanine concentration response there was a significant time effect (P < 0.05), but there was no group effect. Arterial blood phenylalanine concentration increased significantly (P < 0.05) by 15 min and returned to a value that was not different (P > 0.05) than basal by 90 min in all groups, with the exception of the 41% Leu young group, where arterial blood phenylalanine concentration returned to basal by 75 min.

Mean arterial L-[ring-2H3]phenylalanine enrichment of phenylalanine (t/T) in the 26% Leu elderly group was 0.081 ± 0.004 for the period before and 0.092 ± 0.005 for the period after the EAA ingestion, whereas in the 41% Leu elderly group it was 0.081 ± 0.005 before and 0.092 ± 0.005 after the EAA ingestion. In the young, the corresponding values for the 26% Leu group were 0.074 ± 0.004 for the period before and 0.083 ± 0.004 for the period after the EAA ingestion, whereas for the 41% Leu group they were 0.089 ± 0.007 before and 0.091 ± 0.007 after the EAA ingestion. The arterial blood phenylalanine enrichment during the course of the study in the four groups is shown in Fig. 3.

**Muscle free phenylalanine concentration.** Muscle free phenylalanine concentrations were determined at the muscle biopsy sampling time points. Basal muscle free phenylalanine concentrations were: 26% Leu elderly, 79 ± 6 nmol/ml; 41% Leu elderly, 75 ± 6 nmol/ml; 26% Leu young, 68 ± 7 nmol/ml; 41% Leu young, 76 ± 10 nmol/ml. There was no significant group effect, but there was a significant time effect (P ≤ 0.05) after the amino acids ingestion. As expected, muscle free phenylalanine concentration showed an increase at 60 min after the EAA ingestion compared with basal in all groups (values not shown). However, by the end of the study, muscle free phenylalanine concentrations had returned to values that were not different (P > 0.05) from basal (26% Leu elderly, 88 ± 5 nmol/ml; 41% Leu elderly, 69 ± 6 nmol/ml; 26% Leu young, 63 ± 8 nmol/ml; 41% Leu young, 69 ± 11 nmol/ml).

**Plasma insulin.** Arterial plasma insulin concentrations in response to the ingestion of EAAAs containing either 26 or 41% leucine are shown for all groups in Fig. 4. There were no significant group effects, but there was a significant time effect (P < 0.05). Mean plasma insulin concentrations increased significantly (P < 0.05) in all groups in response to the EAAAs and remained significantly different from basal until 30 min in the young groups and until 45 min in the elderly groups. Mean arterial plasma insulin values appeared to peak at a later time in the elderly groups than in the young groups (i.e., 30 vs. 15 min).

**Leg phenylalanine net balance.** Figure 6 depicts the leg phenylalanine net balance response in all four groups during the basal period and the period after the EAA ingestion. There was no significant group effect (P > 0.05), but there was a significant time effect. Within each group, the leg phenylalanine net balance increased immediately after the EAA inges-
tion and remained significantly different from basal until 30 min in the 26% Leu elderly and 41% Leu young groups and until 45 min in the 26% Leu young and 41% Leu elderly groups \((P < 0.05)\).

Figure 7 shows average values of the response of the leg phenylalanine net balance at the basal period and the period after the EAA ingestion for all four groups. There were no differences between groups in the basal period \((P > 0.05)\).

After the EAA ingestion there was a significant improvement in the mean leg phenylalanine net balance in all groups \((P < 0.05)\), with the exception of the 26% Leu elderly group \((P > 0.05)\). There were no significant differences between groups for the post-EAA period \((P > 0.05)\).

**DISCUSSION**

We investigated muscle protein metabolism after the ingestion of two different mixtures composed of \(~7\) grams of EAA: one mixture mimicked the distribution of the EAA in whey protein, whereas the other had higher leucine content. The results suggest that the EAA leucine has a unique role in the stimulation of muscle protein synthesis by EAAs in elderly humans. Specifically, in the elderly, the leucine-enriched EAA mixture stimulated postprandial muscle protein synthesis and resulted in postprandial accretion of muscle proteins, reversing the lack of response following the whey protein-based EAA mixture. In contrast, in the young, both EAA mixtures stimulated muscle protein synthesis, and no unique advantage of extra leucine was evident.

It is now well established that among all the plasma amino acids, the EAAs are the most important for the stimulation of muscle protein synthesis \((12, 36)\). Previous studies have shown that EAAs stimulate muscle protein synthesis in both elderly \((25, 34)\) and young \((25, 32)\) individuals. We have previously found that stimulation of muscle protein synthesis is not different between young and elderly when large amounts of EAAs are ingested \((25)\), but the elderly have reduced muscle protein synthesis when small amounts of EAAs are ingested \((20)\). The latter is evident in the present study, which further underscores the importance of the leucine content in the formulation of any amino acid supplement for the stimulation of muscle protein synthesis in the elderly. Leucine content becomes particularly important when decreasing the overall amount of EAAs in an amino acid supplement, since this decreases the availability of leucine.

It has long been known that EAAs stimulate insulin secretion \((23)\), and among them, leucine appears to be one of the most potent stimuli \((21)\). Following the EAA ingestion there was a transient, but significant, increase in the circulating insulin in all four groups (Fig. 4). An increase in plasma insulin concentration increases net muscle protein balance during hyperaminoacidemia \((10, 13)\), and therefore, it would be expected to have played a role in the improved leg protein retention in the present study. However, the overall response was not different between the 26% Leu and 41% Leu groups, and therefore, the greater muscle protein retention in the elderly following the leucine-enriched EAA mixture cannot be attributed to differences in plasma insulin concentration. Therefore, the insulin data in the present study do not support an insulin-mediated but rather a direct effect of blood leucine on stimulating muscle protein synthesis in the elderly.

Because improvement in muscle protein balance following EAA ingestion (Fig. 7) was observed, together with the increase in muscle protein FSR (Fig. 5), the improved muscle protein balance can be attributed to the stimulation of muscle protein synthesis by the EAA mixtures. The lack of a response in muscle protein synthesis in the 26% Leu elderly group may...
reflect blunted responsiveness to one or more of the ingested amino acids. However, the improved protein synthesis following the leucine-enriched EAAs in the same age group can only be attributed to the increased leucine content in the EAA mixture, since the content of the rest of the EAA in the mixture was decreased. Although it has long been known that the branched-chain amino acids (12), and more specifically leucine, are unique among the amino acids in the stimulation of muscle protein synthesis (6, 14), only recently have the mechanisms of the regulation of muscle protein synthesis by leucine started to be understood. On the basis of this evidence, it can be speculated that, in the present study, activation of S6K1 might have been implicated in the stimulation of muscle protein synthesis by higher plasma leucine concentration. Relative to that, it has been shown that S6K1 is regulated in vitro by leucine and requires higher levels of leucine concentration to be activated in aged rats (8). However, there is also evidence suggesting that high leucine concentrations increase protein synthesis without altering the activity of S6K1, but this response involves other mechanisms, such as an enhanced binding of eukaryotic initiation factor (eIF)4E to eIF4G (5).

Initiation of muscle protein synthesis by leucine may have not been optimally activated following ingestion of the 26% Leu mixture in the elderly, because no significant change in muscle protein FSR was observed. The findings of the present study suggest that ingestion of extra leucine in an amount approximate to that provided in the 41% Leu group may be required to activate the initiation of protein synthesis in the elderly. On the other hand, the young individuals showed improved muscle protein synthesis following the EAA ingestion regardless of the magnitude of changes in blood leucine concentration. It is possible that mechanisms associated with the activation of muscle protein synthesis by leucine in the young are either sufficiently active at basal leucine concentration or are highly sensitive to even small changes in blood leucine concentration. The findings of the present study indicating decreased sensitivity of muscle protein synthesis to leucine in the elderly are consistent with a recently reported study in rats (8). Extra leucine may have failed to further enhance the rate of muscle protein synthesis in the young because of a corresponding decrease in the overall availability of the nonleucine EAA component of the 41% Leu mixture. Support for this argument comes from one of our previously published reports (25), where the same amount of leucine as in the 41% Leu mixture in the present study approximately doubled the net muscle protein synthesis in the young when ingested as a component of 15 g of EAAs.

In addition to stimulation of muscle protein synthesis, inhibition of muscle protein breakdown may have contributed to the observed responses. Muscle protein breakdown can be estimated from the calculated value for $R_d$, although it is recognized that the accuracy of both the $R_d$ and $R_a$ values may be limited by the perturbation of the steady state in the concentration of muscle free phenylalanine after the amino acid ingestion. Nonetheless, an average value during the entire post-EAA period can provide an integrated response because the concentration of muscle free phenylalanine was not different from basal at the end of the study. On the basis of the calculated phenylalanine $R_a$, inhibition of muscle protein degradation by leucine may be more effective at higher blood leucine concentrations in the young, since among all groups the 41% Leu young group showed the largest decrease in phenylalanine $R_a$. Although not significant ($P = 0.09$), this decrease in phenylalanine $R_a$ is in line with previous data when blood leucine increased at similar levels (24).

Animal studies indicate that meals supplemented with leucine can, both acutely (9) and over a period of at least 10 days (29), beneficially affect muscle protein anabolism. The present study provides for the first time in vivo evidence in elderly humans that a relatively small bolus of ingested leucine (~3 g) can acutely improve muscle protein retention and reverse a lack of stimulation of muscle protein synthesis following the ingestion of a small amount of EAAs. Whether these effects of leucine on muscle protein anabolism can be sustained over longer periods of time in conjunction with leucine-supplemented meals remains to be shown. It is important to note that the increase in plasma leucine concentration in the 26% Leu mixture in the present study was similar to that expected following consumption of a meal of average protein content (~15 g of protein) and that the anabolic effect of extra leucine on muscle protein in the present study was observed in the elderly at less than double peak blood leucine concentration relative to the whey protein-based mixture. The present findings do not argue against greater stimulation of muscle protein synthesis by larger increases in peak blood leucine concentration in either elderly or young individuals. Any such hypothesis, however, should be evaluated in the context of blood availability of other amino acids, because limited supply of such amino acids may compromise the potential for muscle protein anabolism under conditions of stimulated muscle protein synthesis (12).

In conclusion, this study demonstrates for the first time in elderly humans that attenuated response of muscle protein synthesis following ingestion of small amounts of amino acids can be reversed by ingestion of extra leucine. The present data emphasize the important role of leucine in the formulation of any amino acid/protein supplement for reversing attenuated response of muscle protein synthesis to nutritional supplementation in the elderly.

ACKNOWLEDGMENTS

We thank the nurses and the staff at the General Clinical Research Center (GCRC) at University of Texas Medical Branch in Galveston, TX, as well as Dan Creson, Susan Minello, and Roxana Hirst. We gratefully acknowledge...
Stephaine J. Blasé, Melissa Bailey, Christopher Danesi, Gaurang K. Jariwala, and Ming-Qian Zheng for skillful technical assistance.

GRANTS

The work was sponsored by National Institute of Arthritis and Musculo-skeletal and Skin Diseases Grant AR-49038 and Shriners Hospital Grant 8490. Studies were conducted at the GCRC at the University of Texas Medical Branch at Galveston, funded by Grant M01-RR-00073 from the National Center for Research Resources, National Institutes of Health, United States Public Health Service. Support was also provided by AminoScience Laboratories, Ajinomoto.

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