Amino acid metabolism and inflammatory burden in ovarian cancer patients undergoing intense oncological therapy

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Summary

**Background & aims:** Cancer and oncological therapy are associated with a progressive physical deterioration, malnutrition, and enhanced inflammatory burden. Our considerable data showing the strong anabolic potential of amino acids (AA) led us to test whether AA can acutely stimulate muscle protein synthesis in cancer patients (CA) undergoing intense chemotherapy.

**Methods:** Mixed muscle fractional synthesis rate (FSR), rates of phenylalanine appearance and disappearance (Ra and Rd), and net phenylalanine balance (NB) were measured during a primed constant infusion of \(\text{L-[ring-}^{2}\text{H}_5\text{]}\text{phenylalanine}\). Blood and muscle tissue samples were collected in the basal state and following ingestion of 40 g of AA given in 30 mL boluses every 10 min for 3 h. Serum and tissue cytokines and NF-\(\kappa\)B expression in skeletal muscle were measured and compared to normative, healthy older controls (OC).

**Results:** Skeletal muscle TNF-\(\alpha\), IL-6, and NF-\(\kappa\)B were elevated in CA. FSR and model-derived protein synthesis (Rd) increased significantly from basal to AA (FSR: 0.052 ± 0.009 vs. 0.120 ± 0.008% h\(^{-1}\), \(P<0.001\); Rd: 23.1 ± 4.1 vs. 36.4 ± 5.0 nmol min\(^{-1}\) 100 mL\(^{-1}\), \(P<0.05\)). Model-derived protein breakdown (Ra) remained unchanged from basal to AA. Phenylalanine NB improved from a negative basal value (−16 ± 2) to zero (0.8 ± 6 nmol min\(^{-1}\) 100 mL\(^{-1}\), \(P<0.05\)) following AA.

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Introduction

Cachexia adds a significant burden to the health of already compromised cancer patients, decreasing quality of life and increasing mortality. Phenotypically, this condition is characterized by a rapid and unwanted loss in lean body mass and fat mass. In a typical individual, a loss of 30% of total body weight reflects a 75% loss of skeletal muscle protein and without immediate therapeutic intervention, often leads to death. Thus, maintaining lean tissue mass and preventing loss of lean muscle tissue is a high priority in the fight against cancer.

The functional impairments characteristic of cancer-related muscle loss are similar, albeit more rapid in onset, to those consistent with age-related sarcopenia. Metabolic abnormalities common to both cancer cachexia and sarcopenia include hormone dysregulation, increased insulin resistance, increased muscle proteolysis, decreased muscle protein synthesis, elevated synthesis of acute phase proteins, and elevated cytokines. Despite these commonalities, the mechanisms behind the rapid loss of muscle mass associated with cachexia, in contrast to the progressive declines seen with sarcopenia, remain poorly understood. Stable isotope tracer methodologies provide a unique opportunity to measure the in vivo rates of skeletal muscle protein synthesis and breakdown in response to physical, nutritional, or pharmacologic interventions. In the case of cancer cachexia, these techniques may yield critical information regarding the mechanisms underlying muscle wasting, and directly assess the efficacy of anti-cachectic interventions.

Considerable debate remains over how host- and tumor-derived factors interact with intracellular signaling mechanisms to disrupt skeletal muscle protein balance. Recently, NF-κB-dependent muscle wasting was found to be activated by late-stage cancer or by chronic inflammation. Further, activation of NF-κB by pathogen associated molecules such as tumor-derived factors and cytokines is an important first step in the intracellular signaling pathway leading to activation of NF-κB mediated muscle protein breakdown. More recently, evidence suggests that expression of pro-cachectic cytokines such as TNF-α and IL-6 may also be responsible for regulating skeletal muscle protein synthesis by down regulating either IGF-1 or inhibiting signaling components of anabolic pathways controlled by IGF-1, Akt and mTOR. However, despite this recent evidence, it is not known how cancer or the concomitant inflammatory process affects in vivo skeletal muscle metabolism. Specifically, we do not know if the skeletal muscle of cancer patients responds to acute nutritional stimuli known to be anabolic in healthy young and older individuals.

Oral amino acids (AA) are acutely anabolic to skeletal muscle in healthy individuals. We recently showed that while age-related differences exist in the time course of the anabolic response, ingestion of AA results in muscle protein anabolism in both young and elderly. We hypothesized that ovarian cancer patients can similarly benefit from oral AA administered in small boluses. In the present study we sought to determine if AA ingestion could acutely stimulate muscle protein synthesis in women with advanced or recurrent ovarian cancer undergoing oncological therapy.

Experimental protocol

Metabolic studies were conducted over an 8 h period following an overnight fast. At 0600 h the morning of the study, polyethylene catheters were inserted into the forearm vein for infusion of labeled phenylalanine, in the wrist vein of the opposite hand for arterIALIZED blood sampling and into the femoral artery and vein of one leg for blood sampling. Blood samples were collected for determination of background phenylalanine enrichment and concentration before starting a primed (2 μmol kg⁻¹) continuous infusion of L-[ring-2H5]phenylalanine (0.05 μmol kg⁻¹ min⁻¹)
from time 0 to 480 min (Figure 1). Muscle biopsy samples (80–100 mg) were collected from the vastus lateralis muscle using a 5 mm Bergström biopsy needle at 120, 300 and 480 min. Samples were frozen in liquid nitrogen and stored at −80°C. At 230 and 420 min a continuous infusion of indocyanine green (ICG) was started into the femoral artery and maintained for 20 min to calculate leg blood flow as previously described. At 230 and 420 min oral amino acids were ingested in small boluses (30 mL) every 10 min from 300 to 480 min (Figure 1) to facilitate gastric accommodation and to prevent bloating and nausea common in this patient population. The mixture contained 40 g of essential and nonessential AA dissolved in 540 mL of distilled water containing sugar-free flavor (Crystal Light, Kraft Foods, White Plains, NY) (Table 1). The proportion of AA in the mixture was selected because it closely resembles that of meat protein and because it had previously been shown to be strongly anabolic in a group of healthy OC.

### Analytical methods

Phenylalanine enrichments and concentrations in blood and muscle samples were analyzed using gas-chromatography mass-spectrometry (GCMS) (Hewlett Packard, Palo Alto, CA) as previously described. Protein-bound phenylalanine enrichment was analyzed with GCMS after protein hydrolysis and AA extraction, using the external curve approach for low enrichments. High sensitivity C-reactive protein (hsCRP) was analyzed using an Immulite 2000 chemiluminescence immunoassay analyzer (DPC, Los Angeles, CA). Skeletal muscle cytokines were determined with commercially available multiplexed biomarker immunoassay kits (Linco Research Inc., St. Charles, MO). All other blood measures were performed by the UTMB Clinical Laboratory.

### Western immunoblot analysis of NF-κB

Cytosolic extracts were prepared from baseline (120 min) muscle tissue for the analysis of NF-κB. Briefly, whole cell extract was prepared from muscle biopsy samples by slicing frozen human muscle into small pieces with a clean razor blade and thawing the tissue in lysis buffer (150 mM NaCl, 10 mM Tris, 1% Triton X-100, 1% Na deoxycholate, 0.1% SDS, 5 mM EDTA) containing protease inhibitors (1 mM phenylmethylsulfonyl fluoride, 1 mM benzamidine, 10 μg/mL pepstatin A) at a concentration of 3 mL of ice cold lysis buffer per gram of tissue. Tissues were homogenized with a Dounce homogenizer and centrifuged at 15,000 g for 20 min, and the supernatants were removed and discarded. Bradford assays were performed for determination of protein concentration of cytosolic extracts. Tissue lysates were electrophoresed on 10% SDS-PAGE gel loaded with 40 μg protein in each well. Immunoblots were incubated with NF-κB p65 (H-268) antibody (Santa Cruz Biotechnology). Chemiluminescence was recorded on Kodak film and analyzed with ImageJ software. Protein loading was evaluated using β-actin.

### Calculations

Phenylalanine was selected to trace muscle protein kinetics because it is neither produced nor metabolized in skeletal muscle. Mixed muscle FSR was calculated by measuring the direct incorporation of L-[ring-2H5]phenylalanine into...
protein, using the precursor-product model:

\[
FSR = [(\text{EP}_2 - \text{EP}_1)/({\text{EM}} \cdot t)] \cdot 60 \cdot 100, (\% h^{-1}),
\]

where \(\text{EP}_1\) and \(\text{EP}_2\) are the enrichments of bound \(^{15}N\)-phenylalanine in consecutive muscle biopsies, \(t\) is the time interval between biopsies and \(\text{EM}\) is the mean \(^{15}N\)-phenylalanine enrichment in the muscle intracellular pool.\(^{21,32}\) Data are expressed as \% h\(^{-1}\).

Protein synthesis (\(\text{Rd}\)) protein breakdown (\(\text{Ra}\)) and net phenylalanine balance (\(\text{NB}\)) were calculated as follows:

\[
\text{Rd} = (\text{Ca} - \text{Ev} - \text{Cv})/\text{BF} \text{ (nmol min}^{-1}\text{100 mL leg}^{-1}),
\]

\[
\text{Ra} = \text{Rd} - \text{NB} \text{ (nmol min}^{-1}\text{100 mL leg}^{-1}),
\]

\[
\text{NB} = (\text{Ca} - \text{Cv}) \cdot \text{BF} \text{ (nmol min}^{-1}\text{100 mL leg}^{-1}),
\]

where \(\text{Ca}\) and \(\text{Cv}\) represent the phenylalanine concentrations in the femoral artery and vein, respectively.\(^{16}\) Data are expressed as \% h\(^{-1}\).

Statistical methods

Statistical analyses were performed using SPSS 13.0 (SPSS Inc.) and Microsoft Excel (Microsoft Corporation). Student \(t\)-tests were used to calculate differences between basal and AA (paired) and between CA and OC (unpaired). \(P \leq 0.05\) was chosen to indicate statistical significance.

Results

Blood chemistry

Laboratory measures were collected to assess nutritional status and cancer inflammatory burden (Table 2). Serum albumin levels, an index of malnutrition, measured in the low normal range in CA. CA-125, a cancer tumor marker was significantly elevated above the normal reference range in CA (\(P = 0.037\)). High sensitivity C-reactive protein levels measured in CA were in the high normal range, suggestive of low-grade systemic inflammation.

Leg blood flow

Leg blood flow in CA remained unchanged from basal to AA (3.55 ± 0.46 vs. 3.54 ± 0.50 mL min\(^{-1}\) 100 mL leg\(^{-1}\)).

Phenylalanine concentration and enrichment

As expected, arterial phenylalanine enrichments fell sharply in the early portion of AA ingestion, while phenylalanine concentrations rose in the expected fashion (Figure 2).

Mixed muscle fractional synthesis rate (FSR)

Mixed muscle FSR increased significantly from a basal value of 0.052 ± 0.009 to 0.120 ± 0.008% h\(^{-1}\) over the 3 h AA period (\(P = 0.001\)) (Figure 3).

Protein synthesis (\(\text{Rd}\))

\(\text{Rd}\), a model-derived index of muscle protein synthesis increased from basal to AA in CA (23.1 ± 4.1 vs. 36.4 ± 5.0 nmol min\(^{-1}\) 100 mL leg\(^{-1}\), \(P \leq 0.05\)) (Figure 4).

Protein breakdown (\(\text{Ra}\))

\(\text{Ra}\), a model-derived index of muscle protein breakdown remained unchanged in CA following AA (38.6 ± 3.7 vs. 35.6 ± 7.2 nmol min\(^{-1}\) 100 mL leg\(^{-1}\) over the last hour of AA ingestion (Figure 5).

Net phenylalanine balance (\(\text{NB}\))

Net phenylalanine balance shifted from net negative to zero (−16 ± 2 vs. 0.8 ± 6 nmol min\(^{-1}\) 100 mL leg\(^{-1}\), \(P \leq 0.05\)) over the last hour of AA ingestion (Figure 5).

Western blot of NF-\(\kappa\)B p65

NF-\(\kappa\)B p65 expression was significantly greater in CA as compared to OC (\(P = 0.0000001\)) (Figure 6).

Discussion

Cancer demands aggressive chemotherapy often resulting in a progressive deterioration in nutritional status, severe protein loss, muscle wasting, and generalized weight loss. Preventing or reversing muscle loss and the cachexia syndrome is a critical part of the intervention process and
requires ongoing vigilance throughout the phases of cancer treatment. In the past few years, we have demonstrated a strong theoretical rationale for the use of AA in clinical populations suffering from muscle wasting.19–21 Our data in both healthy young and old show AA capable of acutely stimulating muscle protein synthesis.19–21 More importantly, we have shown that these acute anabolic benefits translate into the maintenance of lean muscle mass during an otherwise catabolic period of prolonged inactivity.34 Nevertheless, we did not know if a simple AA supplement is capable of acutely stimulating muscle protein synthesis in women undergoing aggressive oncological therapy for advanced and recurrent ovarian cancer.

Several similarities were observed between these cancer patients and healthy older subjects that followed the identical study protocol.21 Leg blood flow and FSR were similar between OC and CA in both the basal state and after AA ingestion. While arterial phenylalanine enrichments were

Figure 2  Arterial phenylalanine concentrations (Ca) and enrichment (Ea) in the basal state ( Ca, Ea) and in response to amino acids ( ■ Ca, ● Ea).

Figure 3  Mixed muscle fractional synthetic rate (FSR) in the basal state (white bars) and in response to amino acids (black bars). **Significantly different than basal, P ≤ 0.001.

Figure 4  Muscle protein synthesis (Rd) and degradation (Ra) in the basal state (white bars) and in response to amino acids (black bars). *Significantly different than basal, P ≤ 0.05.
similar, we observed lower arterial concentrations in CA when compared to the OC in both the basal (48.3 ± 3.8 vs. 59.1 ± 2.1 nmol/mL, CA vs. OC) and AA periods (92.8 ± 3.3 vs. 120 ± 8.3 nmol/mL, CA vs. OC) (P ≤ 0.05). Additionally, the increase in muscle protein synthesis (Rd) from basal to AA was smaller in CA (23.1 ± 4.1 vs. 36.4 ± 5.0 nmol min⁻¹ 100 mL leg⁻¹, P < 0.05), than seen in OC (32.76 ± 7.93 vs. 57.15 ± 8.79 nmol min⁻¹ 100 mL leg⁻¹, P < 0.01).

Muscle protein breakdown (Ra) however was unchanged in response to AA and similar to that previously noted in OC (48.61 ± 9.58 vs. 41.11 ± 8.81 nmol min⁻¹ 100 mL leg⁻¹). Together, these findings resulted in a slightly less positive shift in the NB of CA following AA (−16 ± 2 vs. 0.8 ± 6 nmol min⁻¹ 100 mL leg⁻¹) compared to OC (−16 ± 5 vs. 16 ± 4 nmol min⁻¹ 100 mL leg⁻¹, P ≤ 0.05).

Our findings suggest that AA are acutely anabolic to the skeletal muscle of these metabolically dynamic cancer patients, albeit to a smaller extent than seen in healthy older subjects. Additionally, we showed that this AA-induced anabolism in cancer patients could be attributed to a stimulation of muscle protein synthesis, because muscle protein breakdown remained unchanged. These data are essential in understanding the mechanisms that regulate the loss of muscle mass with cancer and its associated therapies.

In broadest terms, changes in lean muscle mass are a function of the relationship between muscle protein synthesis and breakdown. With cancer, disruption of this kinetic balance is affected not only by the inflammatory burden that accompanies cancer, but also by the various therapeutic interventions. Indeed, some have suggested that the primary factor driving cancer cachexia is enhanced protein degradation occurring through the ubiquitin-proteasome pathway. This pathway is stimulated by inflammatory biomarkers such as TNF-α and NF-κB, both found to be highly elevated and expressed in our cancer group in comparison to healthy OC. Our data show that the anabolic stimulus afforded by AA supplementation occurred despite the catabolic potential of inflammatory biomarkers such as NF-κB p65, TNF-α and IL-6. TNF-α, IL-6 and NF-κB are each known to negatively affect pathways involved in the AA mediated induction of protein synthesis such as the PI3K/Akt/mTOR pathway and activity of initiation factors eIF2 and eIF2B. NF-κB has also been shown to inhibit mTOR activation, which is the pathway by which AA, particularly leucine, stimulate initiation of protein translation. With this evidence, it would seem unlikely that AA could acutely stimulate muscle protein synthesis in cancer patients with an enhanced inflammatory burden. While we did not look at the mTOR signaling pathway directly as it was outside the scope of this study, the fact that FSR and Rd in...
these cancer patients was significantly elevated despite the demonstrated hyperactivation of NF-κB p65 and elevated TNF-α and IL-6 suggests that some portion of the PI3K/Akt/mTOR pathway is intact.

Increased muscle protein breakdown is a major concern in cancer and aging. In cancer cachexia, visceral protein is relatively preserved in comparison to skeletal muscle. This phenomenon is largely due to metabolic recycling of AA liberated from skeletal muscle for splanchnic utilization. While administration of oral AA to cancer patients in the present study resulted in an overall increase in the FSR of skeletal muscle over the 3h absorptive period, the net phenylalanine balance measured in the last hour of AA ingestion failed to achieve a strong positive net balance as seen in previous studies of healthy individuals. This suggests that some portion of these exogenous AA were subjected to an alternative metabolic fate, perhaps toward enhanced acute phase protein synthesis or tumor cell growth. In support of this notion, we showed 20-fold higher concentrations of C-reactive protein among the cancer patients when compared to the healthy OC. While hsCRP remained within normal limits in the cancer patients (0.77 ± 0.13 mg dL⁻¹), these levels were significantly higher than those measured in the older controls (0.04 ± 0.007 mg dL⁻¹). Similar to that observed in the OC, there was no response in protein breakdown (Ra) in the cancer patients following AA ingestion. Taken together, these data indicate that while the cancer patients were able to mount an acute anabolic response to the AA during the absorptive period, anabolism may be attenuated over time as AA are shunted to sustain a low-grade inflammatory response and/or tumor growth. Further outcome-based research is needed to determine if targeted AA such as leucine can provide a more robust stimulation of muscle protein synthesis. Leucine, and to a lesser extend the two other branched chain amino acids valine and isoleucine, is known to stimulate protein synthesis through increased phosphorylation of mTOR, S6K1, and 4E-BP1. While our balanced AA supplement contained 3.2 ± 0.0g leucine, higher concentrations may be required to produce a sufficiently strong anabolic signal in cancer patients. Enhanced tumor growth is a potential risk involved with anabolic interventions. Future outcome-based studies are needed to establish whether the use of AA supplements as an anticachectic intervention increases tumor growth. While the results of this study are encouraging with the finding that AA are acutely anabolic to cancer patients, these patients demonstrated a mild inflammatory burden. Conversely, in patients with more pronounced inflammatory states, such as progressive cachexia and unstable disease, a less anabolic response may be expected.

In conclusion, we have demonstrated that AA ingestion acutely stimulates muscle protein synthesis in ovarian cancer patients with advanced disease and inflammatory syndrome undergoing oncological therapy. The efficacy and potential clinical benefit of AA supplementation in cancer patients was highlighted by the similar anabolic response observed in a cohort of healthy individuals who received the same AA supplement. These data represent a positive first step toward preservation of nutritional status and prevention of cancer cachexia. Additional studies are needed to determine if long-term administration of targeted nutritional therapies such as with leucine are capable of preventing the skeletal ravages of cancer and its associated therapies.

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M.S.-M. was the principal investigator. M.S.-M., E.V., R.R.W. and R.J.U. were responsible for study design and implementation. E.L.D., D.P.-J., and M.S.-M. performed data analysis and interpretation. S.L.C. and S.S. conducted sample processing and analyses. C.D.A. was the physician for the gynecological cancer patients. A.P.S. and E.V. performed the surgical procedures involved in the experiment.


References


