Thermogenic effect and substrate oxidation of protein from animal and plant sources in adults

A Fahmy Arif Tsani, Lee Myung Joo, Kim Eun Kyung

ABSTRACT

Background: Changing nutrient source is one of the efforts to increase thermogenic effect (TEF) which may be significant for body weight reduction. Objective: The aim of this study was to investigate the effects of high protein diets using animal (chicken) and plant (tofu) sources on thermogenic effect (TEF) and substrate oxidation. Method: Ten female adults (mean age 20.8±1.2 y) participated in two isocaloric diet ingestions. Each meal provided 30% of the daily basal energy need (32/26/42% as protein/fat/carbohydrates, respectively). Postprandial energy expenditure was measured by indirect calorimetry. Results: There were no significant differences in TEF and substrate oxidation. The postprandial fat oxidation rate was higher than that at the preprandial state, while carbohydrate and protein oxidation rates were lower. Conclusion: No differences were observed in TEF and substrate oxidation in animal- and plant-based diets. A high protein diet could be beneficial for weight loss, but animal protein does not appear to offer superior benefits compared to plant protein.

KEY WORDS: thermic effect of food, high-protein diet, substrate oxidation, female adult

INTRODUCTION

Obesity has become one of the primary health concerns in the world because its prevalence has increased dramatically. The increase in the prevalence of the associated comorbidities is a significant burden on health care systems worldwide (1). Increased energy intake and decreased energy output are thought to be the two main causes of the development of a positive energy balance, which is related to obesity (2). The thermic effect of food (TEF) is one of the contributors to energy output, besides energy used for basal metabolism, physical activity, and non-exercise activity thermogenesis (3). TEF is the increase in energy expenditure above the resting state, which is required for the digestion and absorption of ingested food. It is the major form of thermogenesis in humans, accounting for 5% to 15% of the total daily energy expenditure (4).

The TEF of separate nutrients is highest for protein, followed by carbohydrates and fat. It has been hypothesized that different protein sources can...
differentially affect TEF (5), but very limited data from human studies on this topic are available. Differential substrate oxidation has been reported in casein (complete protein) and gelatin (incomplete protein) (6). Conversely, protein increases satiety to a greater extent than carbohydrates or fat, and high protein meals induce a greater acute appetite suppressive effect than normal protein meals (7). Considering the possible effects of the protein source, the present study aimed to investigate the thermic effects and substrate oxidation in high protein diets with different animal- and plant based-diets.

METHODS

Subjects

The subjects were 10 healthy female university students who were enrolled at Gangneung-Wonju National University (GWNU), Gangneung city, Gangwon Province, South Korea. The inclusion criteria were as follows: age > 18 years, non-smoker, not pregnant or lactating, and no food allergies. Exclusion criteria included a history of metabolic or endocrine disease, taking medications regularly, and reporting more than moderate physical activity prior to measurements. The subjects were fully informed about the procedures of this study, and informed consent was obtained from each subject. This study was approved by Gangneung Wonju National University prior to implementation.

Study protocols

Two different isocaloric breakfast meals including animal- and plant protein-based diets were tested in a randomized crossover design. After an overnight fast of \( \geq 9 \) hours and a minimal amount of activity, the subjects were randomly assigned to the sequence of test meals. There was a washout period of \( \geq 5 \) days between test days, and each test day lasted \( 7 \) h. On the morning of the test days, the subject travelled to the Department of Food and Nutrition by car, bus, or by walking slowly and arrived at 8.30 a.m. After arriving at the laboratory, the anthropometry of the subjects was measured. Resting energy expenditure was measured after 10 min of bed rest. The subjects were instructed to consume the test meal within 15 min. Over the following 6 h, resting energy expenditure (REE) was measured in periods of 20 min with a 10 min break between each measurement. No additional foods were permitted for the following 6 h. For all REE measurements, the subjects remained in a supine position in an adjustable bed with their head placed under a transparent ventilated hood connected to a monitor by a tube. A documentary video was provided that could be quietly watched by the subjects.

Test meals

The two isocaloric test breakfast meals were provided to the subjects on two different days. Both of the meals were high protein diets with different sources (animal- and plant protein-based). Animal protein was represented by chicken meal, and plant protein was represented by tofu meal. Both of them were cooked and served with rice, kimchi, and a cup of water. Each meal provided 30\% of each individual’s basal energy need, which was determined by weight, height, and age. Table 1 describes the portion of the meal, which was composed of 32\%, 26\%, and 42\% energy as protein, fat, and carbohydrates, respectively. Meals were prepared at the Nutrition Laboratory Department of Food and Nutrition GWNU. Food ingredients were weighed to the nearest 0.1 g. Macronutrient analysis of the meal was performed using the computerized nutrient composition program Computer Aided Nutritional Analysis Program Version 3.0 (CAN Pro 3.0).

Anthropometry

Anthropometric measurements were carried out by the same investigator. The data included weight, height, body mass index, and body composition. Weight and height were measured using a digital scale and stadiometer. Weight was obtained with the subject wearing light clothing and measured to the nearest 0.1 kg.
Height was measured to the nearest 0.1 cm without shoes and socks. Body fat and skeletal muscle were measured by bioelectrical impedance analysis (Inbody720, Biospace Corp., Korea). The waist-hip ratio was determined by measurement using a tape line, while TSF was measured using a skin-fold caliper. Body mass index (BMI), fat free mass (FFM), and body surface area were calculated by the following formulae:

\[
\text{BMI (kg/m}^2\text{)} = \frac{\text{body weight (kg)} }{[\text{height (m)}]^2}
\]

\[
\text{FFM (kg)} = \text{body weight (kg)} - \text{fat mass (kg)}
\]

\[
\text{BSA (m}^2\text{)} = \sqrt{\frac{\text{body weight (kg)} \times [\text{height (cm)}]}{0.007184}}
\]

**Resting energy expenditure and substrate oxidation**

Measurements of fasting, 6h postprandial REE, and substrate oxidation were conducted with indirect calorimetry of a TrueOne 2400 metabolic cart (Model QMC, ParvoMedics Corp., USA), which recorded the amount of O\(_2\) consumed and CO\(_2\) produced. The ventilated hood system was automatically recalibrated every five minutes during measurement.

Upon arrival at the laboratory, subjects were asked to relax for 10 minutes before measurement. The REE was measured for 20 minutes before consumption of each meal. The 6h postprandial energy expenditure was measured every 30 minutes. No additional foods were permitted for the following 6 hours. For all measurements, the subjects remained in a supine position in an adjustable bed with their head placed under a transparent ventilated hood connected to a monitor by a tube. A documentary video was provided that could be quietly watched by subjects.

The REE represents an average of the full collection period, and it was calculated from the oxygen consumption and carbon dioxide production (9). The TEF of test meals was calculated as the difference of the postprandial and the fasting REE (10). The carbohydrate and fat oxidation rate were calculated using the following formula:

\[
\text{Protein oxidation (g/min) = REE (kJ/min) x 0.15/16.74 (11)}
\]

\[
\text{Carbohydrate oxidation (g/min) = } 4.585 \times \text{VCO}_2\text{(L/min)} - 3.2255 \times \text{VO}_2\text{(L/min)}
\]

\[
\text{Fat oxidation (g/min) = } 1.6946 \times \text{VCO}_2\text{(L/min)} - 1.7012 \times \text{VO}_2\text{(L/min)}
\]

**Statistical analysis**

Statistical analysis was performed using the Statistical Analysis System (version 9.2, SAS Institute Inc, Cary, NC). Analysis of variance (ANOVA) for repeated measurements was performed to test the timing effect for differences along the experiment in REE, TEF, and substrate oxidation. A paired t-test was used to compare the differences of the measured parameters between two meals. The postprandial response of the serial measurements of TEF were summarized, and the areas under the curve (AUC) or above the curve (AAC) were calculated using the trapezoid rule (12). The AUCs adjusted for baseline values were calculated by subtracting the values in the fasting state from each postprandial value. The level of significance was set at \(P\)-values less than 0.05.

**RESULTS**

Ten subjects participated in and completed the study. All data concerning anthropometric measurements are summarized in Table 2. The mean age and BMI were 20.8±1.2 years and 21.3±3.4 kg/m\(^2\), respectively. Based on the World Health Organization’s criteria (2), one subject was classified as obese.

The fasting and postprandial energy expenditures are shown in Table 3. Repeated ANOVA measurements showed significant differences in REE between before and after the ingestion of two meals. Postprandial energy expenditure in chicken meals decreased relatively similar to the fasting state at 330 min, which was faster than those who received a tofu meal (at 360 min). Thermic effect of food was different between the two meals at the early measurement time. The chicken meal was significantly more thermogenic at 30 min (kcal/min) than the tofu meal (kcal/min) (Figure 1). Both of the meals reached the maximum point of TEF at 60 min.

Figure 2 shows the amount of TEF in different analyses. The area under the curve representing the total TEF over 6 hours was higher (not significant) in chicken in the tofu meals (47.79 kcal vs. 44.78 kcal; \(P=0.51\)). The percentage of energy intake as TEF in both meals was not different (\(P=0.51\)); it was 12.6% and 11.39% for chicken and tofu meals, respectively. Similarly, the percentage of
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Table 2. Characteristics of the subjects

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Healthy female adults (n=10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>20.8 ± 1.23 (20 ~ 23)</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>159.21 ± 4.78 (152 ~ 168)</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>54.8 ± 11.5 (42.2 ~ 80.9)</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>21.43 ± 3.4 (16.9 ~ 29.1)</td>
</tr>
<tr>
<td>Waist-hip ratio</td>
<td>0.8 ± 5.66 (0.69 ~ 0.9)</td>
</tr>
<tr>
<td>Fat mass (kg)</td>
<td>18.22 ± 4.16 (12.9 ~ 32)</td>
</tr>
<tr>
<td>Fat mass (%)</td>
<td>32.94 ± 6.63 (26.3 ~ 40.4)</td>
</tr>
<tr>
<td>Fat free mass (kg)</td>
<td>36.58 ± 0.06 (29.3 ~ 48.9)</td>
</tr>
</tbody>
</table>

Table 3. Preprandial and postprandial energy expenditures of two meals

<table>
<thead>
<tr>
<th>Time (min)</th>
<th>Chicken meal (animal-based)</th>
<th>Tofu meal (plant-based)</th>
<th>Difference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Preprandial state</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Postprandial state</td>
<td></td>
<td></td>
</tr>
<tr>
<td>30 min</td>
<td>1533.93 ± 229.87**</td>
<td>1459 ± 154.23*</td>
<td>0.0176(S)²</td>
</tr>
<tr>
<td>60 min</td>
<td>1548.52 ± 214.86**</td>
<td>1514 ± 165.36*</td>
<td>0.466 (NS)³</td>
</tr>
<tr>
<td>90 min</td>
<td>1530.12 ± 193.58**</td>
<td>1507 ± 178.39**</td>
<td>0.801 (NS)³</td>
</tr>
<tr>
<td>120 min</td>
<td>1523.83 ± 199.24**</td>
<td>1495 ± 202.64**</td>
<td>0.6652 (NS)³</td>
</tr>
<tr>
<td>150 min</td>
<td>1512.54 ± 219.21**</td>
<td>1475 ± 184.48**</td>
<td>0.507 (NS)³</td>
</tr>
<tr>
<td>180 min</td>
<td>1493.35 ± 216.01**</td>
<td>1462 ± 171.88*</td>
<td>0.7001 (NS)³</td>
</tr>
<tr>
<td>210 min</td>
<td>1492.83 ± 232.55**</td>
<td>1464 ± 186.6*</td>
<td>0.7549 (NS)³</td>
</tr>
<tr>
<td>240 min</td>
<td>1469.26 ± 241.48**</td>
<td>1459 ± 175.35**</td>
<td>0.6995 (NS)³</td>
</tr>
<tr>
<td>270 min</td>
<td>1455.38 ± 216.42*</td>
<td>1446 ± 169.51**</td>
<td>0.7074 (NS)³</td>
</tr>
<tr>
<td>300 min</td>
<td>1436.09 ± 218*</td>
<td>1426 ± 184.06*</td>
<td>0.7469 (NS)³</td>
</tr>
<tr>
<td>330 min</td>
<td>1447.66 ± 214.94</td>
<td>1419 ± 192.26*</td>
<td>0.643 (NS)³</td>
</tr>
<tr>
<td>360 min</td>
<td>1416.35 ± 194.85</td>
<td>1402 ± 179.54</td>
<td>0.9097 (NS)³</td>
</tr>
</tbody>
</table>

¹Significantly different by repeat ANOVA t-test between preprandial and postprandial state; ²S = significant; NS = not significant based on Paired t-test between chicken and tofu meal; ³* = p< 0.05; ** = p< 0.01

Figure 1. Thermic effect curve of chicken and tofu meal

Figure 2. Area under curve (AUC) of TEF during 6 hours measurements
resting energy expenditure as TEF(14.62% vs. 13.97%, \( P=0.71 \)), TEF per weight (0.87 vs. 0.82 kcal/kg, \( P=0.53 \)), and TEF per fat-free mass (1.3 vs. 1.21 kcal/kg, \( P=0.5 \)) were also not significantly different in the two meals.

Substrate oxidation is described in Figure 3. No differences were observed in the carbohydrate oxidation (\( P=0.45 \)), fat oxidation (\( P=0.75 \)), and protein oxidation (\( P=0.24 \)) of both meals. The postprandial carbohydrate oxidation rate was lower than before the consumption of the meals. However, the decreased value of the chicken meal was not significant (\( P=0.07 \)). Conversely, postprandial fat and protein oxidation were significantly higher than in the fasting state in both meals (\( P<0.01 \) and \( P<0.001 \) in the chicken and tofu meals, respectively). Repeated ANOVA analyses showed that postprandial carbohydrate and fat oxidation were relatively similar to the fasting state at 330 min in both meals, while the postprandial protein oxidation reached a similar point as the fasting state at 360 min.

**DISCUSSION**

In the present study, animal- and plant-based diets induced a relatively similar level of energy expenditure. The thermic effect of both meals were determined to be similar based on total measurement, energy intake-adjusted, REE-adjusted, body weight-adjusted, and fat-freemass-adjusted values. This result could be due to both meals having a high protein content, which induced higher TEF compared to other macronutrients. Studies of human energy metabolism consistently report higher energy expenditure following the consumption of meals with high protein content (13). Contrary to this finding, observed that a high animal protein diet produced a greater increase (1.9%) in total energy expenditure thana high plant protein diet (5). This could be caused by the different sources of animal protein that were used. They used pork meal as the animal protein-based diet instead of chicken meal.

Chicken and tofu are two foods with a high protein composition. This study hypothesized that the use of chicken could stimulate higher effects on energy expenditure. However, we did not observe any not significant differences in energy expenditures between the two meals due to a similar effect of thermogenesis. This insignificant postprandial thermogenesis could be caused by a similar content of amino acids between the two meals. A study reported that a well-balanced amino acid mixture produced a higher thermogenic response than an amino acid mixture with a lower biological value (5). Furthermore, the results from animal studies suggested that leucine is particularly thermogenic compared with other amino acids (14).

This study hypothesized that only a small contribution from the fat content in the meals could influence the thermic effect due to the low percentage of fat in both meals, which was 27% of the energy intake.
Furthermore, fat has the lowest thermic effect for separate macronutrients, which ranged from 0% to 3% (15).

The thermic effect of the tofu meal stopped at 360 min, while thermic effect of the chicken meal disappeared at 360 min. Conversely, the thermic effect of the chicken meal was significantly higher than the tofu meal at 30 min. This could be evidence that although both of the meals had similar effect on energy expenditures, animal protein had a more acute effect than that of a plant protein-based diet.

This study showed that chicken and tofu meals did not differ in substrate oxidation. Fat and protein oxidation increased after ingestion of the meals. Protein oxidation lasted longer (at 360 min) than carbohydrate and fat oxidation (at 330 min). This result suggests that a high protein diet influenced the higher macronutrient oxidation rate (11). The differences were only observed in the pattern of increased substrate oxidation after ingestion. The increased carbohydrate, fat, and protein oxidation rates of the chicken meal were relatively faster than those of the tofu meal. This could be caused by a more acute thermic effect of the chicken meal than the tofu meal.

CONCLUSIONS

In conclusion, the present study has shown that TEF and substrate oxidation are not significantly different between animal- and plant-based protein diets. However, animal-based protein diets exerted a more acute thermic effect than plant protein-based diets. These similar effects could be strongly caused by the same texture and amino acid compositions of the diets.

Our future work entails exploring additional studies that are related to this finding. Exploring various age groups and genders with adjustments based on body composition and physical activity could be an alternative for obtaining the true thermic effect of a high protein diet, which could be relevant for weight loss programs.

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Declaration of Interest Statement

The authors declare that there are no known conflicts of interest associated with this publication.

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