Effects of gender and body weight on fibroblast growth factor 23 responsiveness to estimated dietary phosphorus

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Abstract : Fibroblast growth factor 23 (FGF23) is a molecule involved in regulating phosphorus homeostasis. Although some studies indicated an association between serum FGF23 levels and sex, the association has not been fully investigated. The purpose of this study was to evaluate whether sex could influence FGF23 responsiveness to dietary phosphorus intake in healthy individuals. Thirty two healthy subjects between 21 and 28 years were recruited for this study. Subjects performed 24-hour urine collection and blood samples were collected. We estimated phosphorus intake (UC-P) from the urine collection (UC), and evaluated any association between UC-P and serum FGF23 levels. Subsequently, we compared serum FGF23 levels between males and females. Positive correlation was observed between UC-P and serum FGF23 levels. Serum FGF23 levels were significantly higher in males than in females. Serum FGF23 levels/UC-P was significantly higher in females than in males. There was no significant difference in serum FGF23 levels between UC-P and serum FGF23 levels. Serum FGF23 levels were significantly higher in males than in females. Our results indicate that there was no gender difference between FGF23 responsiveness to phosphorus intake per body weight. J. Med. Invest. 63 : 58-62, February, 2016

Keywords : dietary phosphorus intake, serum FGF23 levels, sex, body weight, 24-hour urine collection

INTRODUCTION

Phosphorus is an important constituent of nucleotides and is essential for bone mineralization, muscle function, cellular signal transduction, and energy storage (1). Serum phosphorus levels are usually maintained in the normal range of 2.5 mg/dl to 4.5 mg/dl, however, disorders of the kidney decreases phosphorus excretion in the urine and increases serum phosphorus levels.

It has been reported that an increase in serum phosphorus levels induces vascular calcification, arterial sclerosis, cardiovascular disease and mortality among renal patients (2-4). It was recently reported that a high phosphorus intake increases serum phosphorus levels and impairs endothelial function, even in healthy individuals (5). Therefore, it is advisable that both kidney failure patients and healthy individuals avoid having too much phosphorus. In healthy individuals with normal renal function, phosphorus homeostasis is tightly controlled by several factors, such as parathyroid hormone (PTH), fibroblast growth factor 23 (FGF23) and 1,25-dihydroxyvitamin D (1,25(OH)2D) (6-10). These factors maintain serum phosphorus levels through the control of renal phosphorus excretion and intestinal phosphorus absorption.

FGF23 is a recently identified molecule, involved in regulating phosphorus homeostasis (11-13). Secretion of FGF23 is stimulated by an increase in serum phosphorus levels, and principally acts in the kidney to regulate phosphorus metabolism. FGF23 reduces serum phosphorus levels by at least two mechanisms: 1) it decreases renal phosphorus reabsorption by lowering renal sodium-phosphorus transporter (NaPi) 2a and NaPi 2c expression and 2) it decreases intestinal phosphorus absorption by diminishing plasma 1,25(OH)2D levels which inhibiting 1α-hydroxylase and stimulating 24-25hydroxylase activities. It has been reported that high phosphorus loading increases serum FGF23 levels (6-8). FGF23 is synthesized in osteocytes, and it has been reported that various factors involved in bone metabolism, such as phosphaturic regulating genes with homologies to endopeptidases on the X chromosome (PHEX), dentin matrix protein 1 (DMP1), and estrogen, influence its synthesis (14).

Estrogen is one of the factors affecting bone metabolism and recent studies found that administration of estrogen increased serum FGF23 levels in ovariectomized rats (15). In the Heart and Soul Study, serum FGF23 levels were significantly higher in post-menopausal females compared with males, and were decreased by estrogen therapy (16). Although some studies have indicated at an association between serum FGF23 levels and sex, the association has not been fully investigated. It remains unclear whether secretion of FGF23 in response to dietary phosphorus intake is affected by sex. The purpose of this study was to evaluate whether sex influences FGF23 responsiveness to dietary phosphorus intake in healthy individuals.

MATERIALS AND METHODS

Subjects

Thirty two healthy subjects (14 males and 18 females) between 21 and 28 years were recruited for this study. We performed anthropometric measurements and biochemical examinations of blood to show that subjects had no health problems. The clinical and biological characteristics of the subjects are shown in Table 1. All subjects gave written informed consent and the ethical committee of the University of Shizuoka approved this study. The protocol conformed to the Helsinki Declaration.

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Table 1. Characteristics of the subjects and sex comparison.

<table>
<thead>
<tr>
<th></th>
<th>Total (n=32)</th>
<th>Male (n=14)</th>
<th>Female (n=18)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td>22.8 ± 2.0</td>
<td>22.8 ± 2.0</td>
<td>21.5 ± 0.7</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>HT (cm)</td>
<td>164.4 ± 9.6</td>
<td>172.5 ± 6.7</td>
<td>158.1 ± 6.0</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>BW (kg)</td>
<td>57.3 ± 10.8</td>
<td>65.7 ± 10.8</td>
<td>50.8 ± 4.7</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>BFP (%)</td>
<td>21.4 ± 5.8</td>
<td>18.6 ± 4.9</td>
<td>24.3 ± 4.7</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>LBM (kg)</td>
<td>44.5 ± 9.2</td>
<td>53.1 ± 6.7</td>
<td>37.7 ± 3.6</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>21.1 ± 2.7</td>
<td>22.0 ± 2.7</td>
<td>20.4 ± 2.5</td>
<td>0.11</td>
</tr>
<tr>
<td>TG (mg/dL)</td>
<td>62.3 ± 29.7</td>
<td>72.4 ± 35.4</td>
<td>54.5 ± 22.4</td>
<td>0.16</td>
</tr>
<tr>
<td>LDL-c (mg/dL)</td>
<td>93.3 ± 24.2</td>
<td>87.0 ± 26.4</td>
<td>98.2 ± 21.8</td>
<td>0.20</td>
</tr>
<tr>
<td>HDL-c (mg/dL)</td>
<td>63.4 ± 12.4</td>
<td>58.3 ± 8.2</td>
<td>63.4 ± 12.4</td>
<td>0.05</td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td>4.9 ± 0.2</td>
<td>4.9 ± 0.2</td>
<td>5.0 ± 0.2</td>
<td>0.38</td>
</tr>
<tr>
<td>TP (g/dL)</td>
<td>7.4 ± 0.4</td>
<td>7.4 ± 0.5</td>
<td>7.5 ± 0.4</td>
<td>0.32</td>
</tr>
<tr>
<td>Alb (g/dL)</td>
<td>4.8 ± 0.3</td>
<td>4.8 ± 0.3</td>
<td>4.7 ± 0.3</td>
<td>0.70</td>
</tr>
<tr>
<td>UN (mg/dL)</td>
<td>12.8 ± 2.8</td>
<td>13.3 ± 3.2</td>
<td>12.4 ± 2.5</td>
<td>0.34</td>
</tr>
<tr>
<td>Cre (mg/dL)</td>
<td>0.73 ± 0.14</td>
<td>0.86 ± 0.09</td>
<td>0.63 ± 0.06</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Na (mEq/L)</td>
<td>140.7 ± 1.4</td>
<td>140.9 ± 1.4</td>
<td>140.5 ± 1.3</td>
<td>0.28</td>
</tr>
<tr>
<td>K (mEq/L)</td>
<td>4.2 ± 0.3</td>
<td>4.1 ± 0.3</td>
<td>4.2 ± 0.3</td>
<td>0.46</td>
</tr>
<tr>
<td>Cl (mEq/L)</td>
<td>104.4 ± 1.5</td>
<td>104 ± 1.4</td>
<td>104.7 ± 1.4</td>
<td>0.17</td>
</tr>
<tr>
<td>Ca (mg/dL)</td>
<td>9.5 ± 0.3</td>
<td>9.6 ± 0.3</td>
<td>9.5 ± 0.2</td>
<td>0.20</td>
</tr>
<tr>
<td>Pi (mg/dL)</td>
<td>4.0 ± 0.4</td>
<td>3.7 ± 0.5</td>
<td>4.2 ± 0.3</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>PTH (ng/mL)</td>
<td>42.8 ± 12.4</td>
<td>44.9 ± 14.1</td>
<td>41.2 ± 2.6</td>
<td>0.41</td>
</tr>
</tbody>
</table>

Data are means ± SD. HT, height; BW, body weight; BFP, body fat percentage; LBM, lean body mass; BMI, body mass index; TG, triglyceride; LDL-c, low density lipoprotein-cholesterol; HDL-c, high density lipoprotein-cholesterol; HbA1c, hemoglobin A1c; TP, total protein; Alb, albumin; UN, urea nitrogen; Cre, creatinine; Na, sodium; K, potassium; Cl, chloride; Ca, calcium; Pi, phosphorus; PTH, intact parathyroid hormone.

Two sample t-test or Mann-Whitney test was used for compare male and female.

Protocol

This study was implemented according to a previously published method (17). Subjects performed 24-hour urine collection and weighed dietary records. Subjects were asked to avoid participation in heavy exercise during the study. Subjects weren’t limited in any other activities or dietary intake and were prescribed to spend a day as normal. On the following morning, dietary records, pictures of foods consumed and 24-hour urine samples were collected, in addition, blood was taken after sufficient rest. Subjects participated three times in these exams with at least one day intervals.

24-hour urine collection method

Subjects performed the 24-hour urine collection method using URIN-MATEP (Sumitomo Chemical Co. Ltd., Tokyo, Japan), which could accumulate 1/50th of whole urine, over a 24-hour monitoring period. Subjects were instructed to discard the first morning void and to collect all urine over the following 24-hours, including the first void on the next morning. After weighing the total volume of urine from each individual, samples were dispensed them into vessels for storage at 4°C or -30°C until use. The analysis company, SRL Inc. (Tokyo, Japan) measured urine nitrogen levels, creatinine levels and inorganic phosphorus levels. Weighed dietary records often over- or underestimate phosphorus intake because it does not add phosphorus derived from food additive, fluctuations in phosphorus content due to cooking loss and production region and picking seasons are not considered. We reported that the 24-hour urine collection method can estimate the amount of dietary phosphorus intake, and were superior to estimation by weighed dietary record (17). So, we calculated the daily dietary intake, estimated from the urine collection (UC), of phosphorus (UC-P).

We calculated UC-P based on a report that phosphorus absorption was about 65% (17).

\[
\text{UC-P} (\text{mg/day}) = \text{urine phosphorus excretion (mg/day)} / 0.65 \quad \text{(17)}
\]

Anthropometric and biochemical examinations of blood

Height, body weight, body fat percentage and lean body mass were measured using a bioelectrical impedance analysis method (TANITA TBF-215; TANITA Corporation, Tokyo, Japan). Body Mass Index (BMI) was calculated using the following formula:

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

Blood samples were dispensed into vacuum blood collection tubes and centrifuged (4°C, 3000 rpm, 10 min) immediately. Serum and plasma samples were separated and stored at -30°C until use.

Triglyceride (TG), low-density lipoprotein cholesterol (LDL-Ch), high-density lipoprotein cholesterol (HDL-Ch), hemoglobin A1c (HbA1c), total protein (TP), albumin (Alb), urea nitrogen (UN), creatinine (Cre), sodium (Na), potassium (K), chloride (Cl), calcium (Ca), phosphorus (Pi), intact parathyroid hormone (PTH) and intact FGF23 (FGF23) levels were measured by SRL Inc.

Statistical analysis

Data were described as means ± SD. The data were tested for normality using the Shapiro-Wilk test. All statistical analyses were performed with SPSS Statistic, version 19.0 for Windows (1989-2010, SPSS, Inc., an IBM Company, US) and were considered statistically significant at P < 0.05. Pearson’s correlation coefficient was used to evaluate association between UC-P and serum FGF23 levels. Two-sample t-tests and Mann-Whitney tests were used to compare serum FGF23 levels, serum FGF23 levels/UC-P and serum FGF23 levels/UC-P/BW between the male and female groups.
RESULTS

Subject characteristics

The characteristics of the subjects are shown in Table 1. Mean age was 22.1±1.5 years and body mass index was 21.0±2.7 kg/m². Subjects had normal glucose and lipid metabolism, hepatic function and renal function.

The association between UC-P and serum FGF23 levels

Positive correlation was observed between UC-P and serum FGF23 levels (r=0.35; P<0.01) (Figure 1).

Serum FGF23 levels comparisons between males and females

To evaluate whether sex could influence FGF23 responsiveness to dietary phosphorus intake, serum FGF23 levels were compared between male and female subjects (Figure 2). Serum FGF23 levels were significantly higher in males than in females (37.4±8.2 pg/ml and 33.1±7.5 pg/ml, respectively) (P<0.05) (Figure 2a). UC-P was also significantly higher in males compared to females (1141±308 mg/day and 788±264 mg/day, respectively) (P<0.01) (Figure 2b). However, serum FGF23 levels/UC-P was significantly higher in females than in males (P<0.01). Serum levels/UC-P in males was 0.035±0.012 pg/ml and in females was 0.047±0.025 pg/ml (Figure 2c). UC-P/BW was significantly higher in males than in females (17.4±4.3 mg/kg and 15.6±4.9 mg/kg, respectively) (P<0.05) (Figure 2d). To examine the effect of UC-P/BW on serum FGF23 levels, we analyzed sex differences in serum FGF23 levels/UC-P/BW. There was no significant difference in serum FGF23 levels/UC-P/BW between males and females (2.3±0.7 pg/ml and 2.4±1.5 pg/ml, respectively) (P=0.95) (Figure 2e).

DISCUSSION

This study evaluated whether sex influenced FGF23 responsiveness to dietary phosphorus intake. UC-P and serum FGF23 levels were positively associated. FGF23 contributed to phosphorus homoeostasis through suppression of renal phosphorus reabsorption and intestinal phosphorus absorption (18, 19). In a previous study, using a phosphorus loading test with three test meals containing different phosphorus levels (400 mg/meal: P400, 800 mg/meal: P800, and 1200 mg/meal: P1200), serum FGF23 levels were little changed after P400 and P800 loading, but significantly increased at 8 h after P1200 loading (6). Another study reported that serum FGF23 levels were significantly higher after 2500 mg phosphorus...
loading than after 500 mg phosphorus loading (20). Therefore, the association between UC-P and serum FGF23 levels observed in this study suggests that FGF23 was secreted in response to the amount of phosphorus ingested in order to maintain phosphorus homeostasis, however, the correlation was weak (r = 0.35; P < 0.01). We previously reported that the 24-hour urine collection method could be used to estimate diurnal dietary phosphorus intake (17). It is considered that serum FGF23 levels might be affected by prolonged phosphorus intake, PTH (21), 1,25(OH)2D (21) and genetic factor such as single nucleotide polymorphism (22, 23), which might be one of the reasons for a weak association between UC-P and serum FGF23 levels. In this study, associations between serum FGF23 levels and these factors were not investigated, so we needed to investigate these associations in the future study.

We compared UC-P and serum FGF23 levels between male and female subjects to evaluate whether sex could influence FGF23 responsiveness to dietary phosphorus intake. Serum FGF23 levels were significantly higher in males compared to females. However, serum FGF23 levels/UC-P was significantly higher in females than in males. There was no significant difference in serum FGF23 levels/UC-P/BW between male and female subjects. Male subjects had significantly higher UC-P/BW compared with females. These results suggest that there is no difference in FGF23 responsiveness to phosphorus intake per body weight between males and females, but UC-P/BW in males was higher than in females and as a result, serum FGF23 levels was significantly higher in males. The recommended intake amounts of phosphorus are 1000 mg/day for adult males and 900 mg/day for adult females, and the tolerable upper limit of phosphorus intake is 3000 mg/day for both adult males and females in Japan (24). The results of this study suggest that recommended intake and tolerable upper limit of phosphorus intake needs to be established according to body weight.

A previous study reported that serum FGF23 levels in postmenopausal females were significantly higher than in males, and were decreased by estrogen therapy (18). Since the estrogen receptor is present at the proximal tubule which regulates urinary phosphorus excretion, it has been suggested that serum phosphorus levels are increased by hormonal imbalance with menopause, and FGF23 secretion is increased. A previous study suggested that serum phosphorus levels are higher in postmenopausal females compared with menstruating females (25). It is also reported that postmenopausal status is associated with higher serum phosphorus levels (26). The female subjects in this study were premenopausal, suggesting there would be no difference in serum FGF23 levels/UC-P/BW between male and female subjects, however, estrogen was not measured in this study and it is unclear whether estrogen influenced FGF23 responsiveness to phosphorus intake.

Phosphorus intake and serum FGF23 levels correlated in all subjects in this study however, the correlation was weaker in female subjects (data not shown). There are two possible reasons for these results: the female subjects in this study may have paid more attention to their diet and body habitus compared to male subjects (27). The variation in body weight, body fat mass and percent body fat was distributed in a narrow range, and as a result, serum FGF23 levels was significantly higher in males compared to females. However, UC-P/BW was significantly higher in females compared with males. The recommended intake amounts of phosphorus are 1000 mg/day for adult males and 900 mg/day for adult females, and the tolerable upper limit of phosphorus intake is 3000 mg/day for both adult males and females in Japan (24). The results of this study suggest that recommended intake and tolerable upper limit of phosphorus intake needs to be established according to body weight.

CONFLICT OF INTEREST

All authors declare that they had no conflict of interest.

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