Investigation of Metacarpal Bone Morphology in Normal-weight, Overweight and Obese Adolescent Subjects

Normal Kilolu, Aşırı Kilolu ve Obez Adölesan Bireylerde Metakarpal Kemik Morfolojisinin İncelenmesi

Objective: The prevalence of childhood and adolescent overweight and obesity is increasing in most developed countries. This study aimed to investigate the metacarpal bone morphology in normal-weight, overweight and obese adolescent subjects at different pubertal stages.

Materials and Methods: This radiographic study was performed in 124 subjects at different pubertal stages. The subjects were divided into three groups based on body mass index percentile: normal-weight, overweight and obese. The second and fourth metacarpal bone cortical thickness, width and metacarpal index (MCI) were measured on left hand-wrist radiographs.

Results: The values of the second and fourth metacarpal bone cortical thickness and width were significantly different among the groups. The second metacarpal bone MCI was significantly different among the groups. Moreover, the values of the second and fourth metacarpal bone cortical thickness, width and MCI in obese and overweight subjects were greater than those in normal-weight subjects. The values of the second and fourth metacarpal bone cortical thickness and width were significantly different in subjects before and after their pubertal growth peak period.

Conclusion: The metacarpal bone parameter in overweight and obese adolescent subjects was significantly greater than that in normal-weight subjects.
Introduction

Obesity prevalence is increasing rapidly and it is one of the most widespread metabolic diseases in developed countries. The terms overweight and obesity are defined as a condition of abnormal or excessive fat accumulation in adipose tissue at a level that could damage health. It is related to excess food consumption, poor dietary intake, and insufficient physical activity. According to the World Health Organization, childhood obesity is one of the most important health problems due to its rapid rate of increase. Obesity development in childhood is associated with the following obesity in adulthood (1). Body mass index (BMI) is a convenient and easy way of classifying obesity and overweight. It is formulized as the weight (kilogram) divided by the square of height (meters) (2). However, BMI is not as reliable in childhood because of the growth potential of the patient which enhances BMI through the growth of fat-free tissues (3). BMI percentile which is age and gender-specific is essential to provide a correct classification of BMI in childhood, and it is referred to as BMI-for-age (4). According to this classification, BMI under the 5th percentile describes underweight, BMI between 5th-84th percentile describes normal, BMI between 85th-94th percentile describes overweight and BMI upper 95th percentile describes obese (5).

The effect of obesity on the skeletal system has been a topic of interest in the literature. Mechanical loading caused by increased body weight in overweight and obese subjects leads to a beneficial effect on bone formation and thus, acts as a protector against osteoporosis (6). It has been indicated that obesity enhances bone mass and reduces fracture risk (7). On the contrary, it has also been shown that overweight patients have an increased risk of fracture (5). Different mechanisms play a role in bone metabolism in obese patients. Increased proinflammatory cytokines, alterations in bone turnover, and hormonal and mechanical loading on bone in obese patients are among these mechanisms (8). Leonard et al. (9) concluded that obesity during childhood and adolescence was associated with increased whole-body bone dimensions and mass. Studies have shown that obesity affects cortical bone rather than trabecular bone (10).

One of the important factors that affect childhood growth is nutrition (11). While excess body weight is related to early pubertal growth, poor nutrition is associated with delayed pubertal growth spurt (12). The effect of childhood obesity on skeletal maturation was previously investigated in the literature and it was shown to be associated with early pubertal development and taller stature (13). Akridge et al. (4) used Fishman’s hand-wrist analysis to assess whether increased BMI results in accelerated skeletal age and their results showed that overweight and obese children did not have significantly accelerated skeletal maturation after age and gender adjusting.

Skeletal maturation is an important factor in the planning of an orthodontic treatment which may contain growth modulation to correct skeletal discrepancies of the jaws. Compared to chronological age, skeletal age determination is more secure to determine physical maturation. Different methods have been described to assess maturational status. The cervical vertebrae, the elbow, the foot, the ankle, and the hip are the skeletal parts that can provide information about maturation (14). Nevertheless, the hand-wrist radiographs are the most commonly used ones to determine skeletal maturation by orthodontists in many ways (15). The hand-wrist radiographs are highly correlated with chronological age and the effects on other skeletal parts (16). Besides different methods of skeletal age assessment, hand-wrist radiographs could also be used to assess the metacarpal index (MCI).

MCI is a measure of bone mass quantification that stands for the cortical thickness of long bones relative to the outer bone diameter at the measuring site. It could be accepted as a relative evaluation of...
the cortical bone thickness. Barnett and Nordin (17) firstly described MCI in 1960 which represents the cortical thickness of the mid-second metacarpal at the radial plus ulnar sides divided by the outer width of the same bone. It is considered as one of the evaluation methods of osteoporosis and fracture risk and digital X-ray radiogrammetry offers the advantage of practical MCI measurement (18). Cortical thickness of the metacarpals and accordingly the MCI decreases with age like in other long bones, especially after menopause in women, and it represents a risk factor for fracture (19,20).

Considering the effect of obesity on bone metabolism, the aim of this study was to investigate metacarpal bone geometry in normal-weight, overweight and obese orthodontic children and adolescents at different pubertal stages.

**Materials and Methods**

The local research ethics committee of the Ordu University Granted Ethical Approval for the study (no: 2018/23). Patients with general health problems, bone diseases, syndromes, endocrine diseases, metabolic disorders and trauma or surgery history to the hand-wrist were not included. The hand-wrist radiographs of low quality were excluded. This cross-sectional radiographic study was performed in 124 subjects who referred to the Ordu University, Department of Orthodontics. Informed consents were taken from the parents of the subjects that were below the age of 18. According to the BMI percentile, three groups were generated. Normal-weight, overweight and obese groups comprised 48 (18 males, 30 females; mean age, 13.92±2.43 years), 37 (10 males, 27 females; mean age, 13.98±2.59 years) and 39 (16 males, 23 females; mean age, 12.82±2.38 years) subjects, respectively. Distribution of the groups according to BMI, gender and chronologic age were given in Table 1. No statistically significant differences in age and gender were observed between the groups. The sample size was determined by G*Power Software version 3.1.9.2 (Universität Düsseldorf, Germany) for the second MCI at alpha error probability of 0.05 and a power of 90%. The power analysis showed that 80 samples were required. To strengthen the study, a total of 124 subjects were included in the present study.

With the use of a mechanical weighing scale with 0.1 kg accuracy and a wall-mounted stadiometer with the graduation of 1 mm, the patient’s body weight and height were obtained and BMI calculation was carried out by dividing weight (kilogram) to the square of height (meters) at the patient’s first visit to the clinic. Age- and gender-specific BMI percentiles were calculated using the Centers for Disease Control and Prevention (CDC) guidelines (21). Patients were classified according to their BMI as follows: 5th-84th percentiles were characterized as normal-weight, 85th-94th percentile were characterized as overweight, and above 95th percentile were characterized as obese.

All left hand-wrist radiographs were obtained by the same roentgraphic film device (Kodak 8000C Digital Panoramic and Cephalometric System, Cephalostat, Corestream Health Inc, Rochester NY, US). Measurements of the second and fourth metacarpal bones’ cortical thickness and width were performed at the midpoint of the bones, using a cephalometric software program (Foxit Reader software, Foxit Corporation, Fremont, CA, USA) by the same investigator after calibration and after that, MCI values were calculated (Figure 1A, B).

To also analyze the effect of pubertal growth stage on the MCI of overweight and obese patients, the

<table>
<thead>
<tr>
<th>Groups</th>
<th>Male/female</th>
<th>Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal-weight</td>
<td>18/30</td>
<td>13.92 (2.43)</td>
</tr>
<tr>
<td>Overweight</td>
<td>10/27</td>
<td>13.98 (2.59)</td>
</tr>
<tr>
<td>Obese</td>
<td>16/23</td>
<td>12.82 (2.38)</td>
</tr>
<tr>
<td>p</td>
<td>0.414a</td>
<td>0.066b</td>
</tr>
</tbody>
</table>

| a: Results of chi-square test, b: Results of One-Way ANOVA test |
patients were classified according to their growth stage by Fishman skeletal maturation stages on the hand-wrist radiographs (22). However, when the patients were divided into 11 groups using Fishman’s method, the groups did not have a sufficient number of patients. Therefore, the patients were divided into 2 growth stages to represent before and after pubertal growth peak periods. Growth stage 1 (GS1) comprised patients that were in stages 1-5 and growth stage 2 (GS2) contained patients with stages of 6-11 according to Fishman’s method.

**Statistical Analysis**

All statistical analyses were performed by using SPSS (SPSS for Windows version 20.0; SPSS Inc, Chicago, IL) program. After performing the normal distribution test, non-parametric tests were performed to the parameters with non-normal distributions, while parametric tests were applied to the parameters having a normal distribution. Comparison of the groups according to BMI percentile, gender and chronologic age were performed by chi-square and One-way ANOVA tests, respectively. Comparisons between second and fourth metacarpal bone cortical thickness, width and MCI measurements among the different BMI percentile groups were performed using the One-way ANOVA, Kruskal-Wallis and post-hoc tests. In all statistical tests, values of p<0.05 were considered statistically significant.

**Results**

The second metacarpal bone’s cortical thickness, width and MCI values were significantly different among normal-weight, overweight and obese patients, while only cortical thickness and width values of the fourth metacarpal bone were significantly different (p<0.05). No significant difference was observed for the fourth metacarpal bone MCI values. Post-hoc comparisons showed that the significant differences were between normal weight-overweight and normal weight-obese patient groups, with higher values in overweight and obese groups compared to normal-weight group. Overweight-obese groups did not differ from each other significantly for the measured parameters (Table 2).

Table 3 displays the cortical thickness, width and MCI values of second and fourth metacarpal bones of the different BMI percentile groups at different pubertal growth stages. Significant differences were found in second and fourth metacarpal bones’ cortical thickness and width between normal-weight, overweight and obese patients before pubertal GS1 (p<0.05). All of the measured parameters of second and fourth metacarpal bones were significantly different among the weight groups after the pubertal peak period (GS2) (p<0.05). Statistically significant differences were found at some of the investigated parameters between normal-weight and obese subjects at GS1, while more parameters differed significantly between normal weight-overweight and normal weight-obese subjects at GS2 using post-hoc tests (p<0.05). Overweight and obese patients have significantly higher values in these parameters, compared to normal-weight patients. No significant differences were observed between overweight-obese groups according to the pubertal growth stage.

**Discussion**

Childhood and adolescent overweight and obesity prevalence are rising in almost all industrialized
countries. Childhood obesity has been associated with multiple general health problems such as endocrine diseases, metabolic and cardiovascular issues (12, 23), obstructive sleep apnea, gastrointestinal diseases, orthopedic complications (24), low self-esteem (25), musculoskeletal complaints (26) and fractures (27).

Controversial results were found in the studies analyzing the effect of obesity on bone metabolism. Several studies demonstrated that overweight and obese children have a significantly higher bone mineral density (BMD) and bone mineral content (BMC) (28-30). Enhanced adipose tissue in childhood obesity is considered as the ground for enhanced total BMD through increased mechanical load on the bone (9, 30). Leonard et al. (9) investigated the effect of childhood obesity on skeletal mass and dimensions in 132 non-obese and 103 obese subjects and their findings suggested that obesity in children and adolescents resulted in enhanced vertebral bone density and enhanced whole-body bone dimensions and mass. In their review that was conducted on twenty-seven studies and a total of 5958 children, van Leeuwen et al. (31) showed that overweight and obese children had a higher peak bone mass, BMD and BMC than normal-weight children. They also indicated that being obese compared with overweight does not have as much difference as being obese or overweight compared with being of normal weight in terms of BMD and BMC. In contrast with the studies that identify a positive relationship between obesity and bone mass, there are also a few studies in the literature showing a negative relationship between these two (10, 32).

In accordance with the findings of the studies showing a positive association between obesity and bone mass and dimensions, our results showed an enhanced metacarpal width and cortical thickness in overweight and obese adolescents than normal-weight patients and no statistically significant difference was observed between overweight and obese subjects.

Oestrogen is a substantial factor in bone metabolism. Oestrogen deficiency results in an increased osteoclast formation and an imbalance in bone remodeling and accordingly, bone mass reduction (33). Fat tissue plays an important role in oestrogen metabolism. Therefore, overweight and obese patients have enhanced oestrogen levels that cause a favorable effect on bone formation (34). Other than oestrogen, leptin is also a significant determinative in bone metabolism. With its effect on regulating appetite, leptin is a mediator that is

Table 3. Metacarpal index, cortical thickness and width of second and fourth metacarpal bones of the different body mass index percentile groups according to pubertal growth stage

<table>
<thead>
<tr>
<th>Growth stage</th>
<th>Metacarpal bone</th>
<th>Parameters</th>
<th>Normal-weight</th>
<th>Overweight</th>
<th>Obese</th>
<th>p</th>
<th>Post-hoc test*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>N-OW</td>
<td>N-O</td>
<td>OW-O</td>
<td></td>
<td>N-OW</td>
</tr>
<tr>
<td>GS1</td>
<td>2nd Metacarpal</td>
<td>Cortical thickness</td>
<td>1.52 (0.17)</td>
<td>1.74 (0.42)</td>
<td>1.91 (0.16)</td>
<td>0.002*</td>
<td>0.153</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Width</td>
<td>5.90 (0.21)</td>
<td>6.33 (0.99)</td>
<td>6.84 (0.34)</td>
<td>0.001*</td>
<td>0.153</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Metacarpal index</td>
<td>0.52 (0.06)</td>
<td>0.54 (0.05)</td>
<td>0.56 (0.04)</td>
<td>0.135*</td>
<td>0.459</td>
</tr>
<tr>
<td></td>
<td>4th Metacarpal</td>
<td>Cortical thickness</td>
<td>1.34 (0.16)</td>
<td>1.45 (0.29)</td>
<td>1.56 (0.20)</td>
<td>0.044*</td>
<td>0.472</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Width</td>
<td>5.32 (0.45)</td>
<td>5.78 (0.78)</td>
<td>5.80 (0.33)</td>
<td>0.048*</td>
<td>0.219</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Metacarpal index</td>
<td>0.51 (0.07)</td>
<td>0.50 (0.04)</td>
<td>0.54 (0.06)</td>
<td>0.321*</td>
<td>0.984</td>
</tr>
<tr>
<td>GS2</td>
<td>2nd Metacarpal</td>
<td>Cortical thickness</td>
<td>1.96 (0.28)</td>
<td>2.33 (0.36)</td>
<td>2.23 (0.29)</td>
<td>0.000*</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Width</td>
<td>6.47 (0.70)</td>
<td>7.07 (0.76)</td>
<td>7.06 (0.89)</td>
<td>0.003*</td>
<td>0.007</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Metacarpal index</td>
<td>0.61 (0.07)</td>
<td>0.66 (0.08)</td>
<td>0.64 (0.08)</td>
<td>0.027*</td>
<td>0.020</td>
</tr>
<tr>
<td></td>
<td>4th Metacarpal</td>
<td>Cortical thickness</td>
<td>1.49 (0.18)</td>
<td>1.72 (0.29)</td>
<td>1.72 (0.28)</td>
<td>0.000*</td>
<td>0.001</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Width</td>
<td>5.33 (0.51)</td>
<td>5.74 (0.66)</td>
<td>5.64 (0.81)</td>
<td>0.034*</td>
<td>0.036</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Metacarpal index</td>
<td>0.56 (0.05)</td>
<td>0.60 (0.09)</td>
<td>0.62 (0.10)</td>
<td>0.021*</td>
<td>0.115</td>
</tr>
</tbody>
</table>

*p*: Results of One-way ANOVA test, p*: Results of Kruskal-Wallis test, §: Post-hoc Tukey's honestly significant difference test for parametric data and Mann-Whitney U test for non-parametric data, GS1: Before pubertal growth pick period, GS2: After pubertal growth pick period; N: Normal-weight, OW: Overweight, O: Obese groups.
secreted by adipocytes. It plays a critical role in skeletal development (35). Obese patients have increased proinflammatory leptin levels (36). As the amount of fat in the body increases, the leptin concentration also increases. Acting as a growth factor on skeletal growth centers’ chondrocytes, increased leptin levels in obese and overweight children result in enhanced bone mass than normal-weight children (37,38). Potentially increased leptin and oestrogen levels in overweight and obese subjects in this study may have an effect on increased metacarpal bone cortical thickness and width.

The revised CDC BMI percentile charts which are considered as a standard for children and adolescents aged between 2 and 20 were used in the present study. Clinical use of BMI percentile is the most rapid and practical method to assess weight status. However, the separation between fat tissue and muscle is not possible when defining a subject as overweight or obese using the BMI percentile and it should not be taken into account as a definitive marker of overweight and obesity. Subjects considered as obese using BMI percentile might not have excess adipose tissue. Diagnosis of overweight and obesity could be supported with other methods of adiposity analysis such as dual energy X-ray absorptiometry, skinfold thickness, air-displacement plethysmography, hydrostatic weighing, arm and waist circumferences, isotope dilution, bioelectrical impedance analysis, computerized tomography and magnetic resonance imaging (39,40).

The effect of fat mass on bone growth is shown to be related to the stage of puberty (41). The strength of this study is that the effect of growth stages and pubertal growth peak was also taken into consideration while assessing the effect of obesity on metacarpal bone geometry. The lack of prospective data to analyze the long-term effect of obesity on metacarpal cortical thickness, width and MCI can be considered a limitation of this study due to its cross-sectional design. Further long-term prospective researches should be designed.

Conclusion

Overweight and obese adolescents have significantly greater metacarpal bone parameters than normal-weight patients. Orthodontists should be aware of these metacarpal bone changes whilst evaluating hand-wrist radiographs for treatment timing and planning in overweight and obese patients.

Ethics

Ethics Committee Approval: The local research ethics committee of the Ordu University granted ethical approval for the study (decision no: 2018/23, date: 01/02/2018).

Informed Consent: Informed consents were taken from the parents of the subjects that were below the age of 18.

Peer-review: Externally peer-reviewed.

Authorship Contributions


Conflict of Interest: No conflict of interest was declared by the authors.

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