The bone densitometry is normal in Turner syndrome prepubertal patients after height age correction

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Abstract

Objectives: to evaluate the bone mass in prepubertal patients with Turner Syndrome (TS) according to height age (HA) and verify the influence of karyotype and adiposity.

Methods: retrospective and analytical study of prepubertal TS patients. The variables analyzed were: karyotype, age at bone densitometry (BD), height, body mass index (BMI) and BD result. The result of the BD was corrected using HA. BMI and BD were calculated on Z score for chronological age (CA) and for HA.

Results: thirty-seven prepubertal patients were selected and after exclusion criteria, 13 cases between 10 and 13 years old were included in the study. The BD for HA was significantly higher than for CA (0.39 ± 1.18 x -1.62 ± 1.32), without karyotype (p=0.369) and BMI (p=0.697) influence.

Conclusion: prepubertal TS patients present normal BD when corrected for HA, without influence of karyotype and BMI.

Key words Turner syndrome, Bone density, Densitometry
Introduction

Turner Syndrome (TS) is characterized by the presence of an intact X chromosome and the complete or partial loss of the second sex chromosome (X or Y) and is associated with skeletal abnormalities that include short stature, in addition to other clinical manifestations such as characteristic physical features, ovarian failure, early hearing loss, congenital cardiovascular changes, and a higher frequency of other diseases such as hypothyroidism and celiac disease. It has a prevalence of 1 in every 2,000 female live births. 45,X monosomy is present in around 40% to 50% of cases, while the other cases present mosaicism with one or more additional cell lines (X or Y) and/or structural alterations (X or Y). 1,3

The etiology and mechanism of bone fragility in TS have not yet been fully elucidated and may be associated to intrinsic bone defects induced by chromosomal alteration that are exacerbated by hormonal factors. 2-4 Patients with TS have a 25% increase in the risk of fractures. 5,6 Normal bone architecture is influenced by the continuous process of bone modeling and remodeling, the regulation of chondrocytes is complex and involves endocrine factors (growth hormone, estrogen, thyroid hormone), and local/paracrine factors. 6

Bone densitometry (BD) using dual-energy X-ray absorptiometry (DXA) has been used since the 1980s to assess bone mineral content, but it only analyzes the two-dimensional bone measurement (g/cm²) and not its actual volume (g/cm³). This is the most used method worldwide, justified by its easiness, speed, reproducibility, precision, safety, low cost, and availability to predict fracture risk. 7

In children, the interpretation of the BD must be performed using the Z-score parameter (it evaluates the mean of the normal population of the same sex and age). In patients with short stature, it is necessary to correct the study values for bone age or, as more recently recommended, for height age. 8-10

The diagnosis of low bone mass is made when a Z-score of less than two standard deviations (< -2.0) for age is evidenced and, for the diagnosis of osteoporosis, it is necessary that the child has fractured one or more vertebrae in the absence of local disease or high-energy trauma or having a significant history of previous fractures associated with low bone density. 11

Several studies have shown that patients with TS have low BD when evaluated via studies with DXA. However, the studies have limitations because they use a two-dimensional method and patients with short stature, such as in TS, may receive an underestimated result with decreased BD, which can be corrected for height or for volumetric BD (g/cm³). 5,6,12

Bone mineral density (BMD) is believed to be influenced by modifiable genetic and environmental factors such as adiposity, hormone levels, sun exposure, blood pressure, physical activity, and diet. 13

Therefore, this study aimed to evaluate BD in patients with TS with correction for age and height and to verify the interference of karyotype and BMI.

Methods

This is a descriptive and analytical cross-sectional study and retrospective analysis of medical records of patients diagnosed with TS, followed at the Pediatric Endocrinology Outpatient Clinic at the Teaching Hospital of the University of Campinas (HC-Unicamp – Portuguese acronym). This outpatient clinic at HC-Unicamp monitors patients with Turner Syndrome with biannual consultations for treatment and guidance, seeking to improve their quality of life with modifiable factors such as diet and physical activity.

Inclusion criteria were: patients diagnosed with TS, in the prepubertal stage (aiming to homogenize the sample, avoiding hormonal interference and the characteristic events of puberty in bone mass); with BD evaluation performed at the HC-Unicamp, using the Hologic Discovery Wi bone densitometer (S/N83901).

The exclusion criteria were the use of medications aimed to improve growth (such as growth hormone) and the presence of chronic disease or use of medications that could interfere with bone metabolism.

The diagnosis of TS was confirmed through the karyotype with evaluation of at least 20 metaphysis. The variables analyzed were: karyotype, length of follow-up at the service, age at BD; height and BMI obtained in the medical appointment closest to the performance of the BD and the BD result in Z-score. To evaluate the results without the bias of short stature, the BD Z-score was corrected using, instead of chronological age (CA), the height age (HA), that is, using as age that corresponding to the 50 percentile of the measured height. 14

The BMI in Z-score was used to assess adiposity. SPSS software, version 16.0 was used for data statistical analysis. The mean, standard deviation, minimum, maximum, median, and interquartile range of quantitative variables were determined. The dependent variables were: BD Z-score for CA and
HA; to compare the results of the Z-score for both BD-CA and BD-HA, the Wilcoxon test was used. To compare the distribution of BDs in relation to karyotype and BMI, the Mann-Whitney test was used and the probabilities were calculated using the bootstrap technique, Monte Carlo method, from 10,000 samples. A 5% significance level was used for results interpretation.

Results

In this study, 37 patients aged 8–13 years were selected. Among them, 13 were excluded for not having undergone any BD at the HC-Unicamp, eight for having used growth hormone, two patients for having spontaneous puberty at the time of the BD assessment, and one patient with an associated diagnosis of non-classic Congenital Adrenal Hyperplasia and uses steroids. Thus, 13 patients were included in the study, four of whom had had more than one BD assessment. Therefore, exams performed between the ages of 10 and 13 were selected in order to homogenize the study. Patients included in the study had follow-up at the service for an average period of eight years.

Regarding the chromosomal constitution, six individuals (46.1%) had 45,X karyotype and seven (53.8%) had mosaicism with a 46,XX normal lineage and/or with structural alterations of the X chromosome. None had Y chromosome intact or not. The CA at the time of BD was significantly higher ($p<0.001$; Wilcoxon test) compared to HA $(12.5 \pm 1.05 \times 9.05 \pm 1.20)$ (Table 1).

The mean Z-score of the BD-CA was significantly lower ($p<0.001$; Wilcoxon test) than that of the BD-HA $(−1.62 \pm 1.32 \times 1.39 \pm 1.18)$ (Table 1) (Figure 1).

The mean of the Z-score of the BD-CA did not differ significantly ($p=0.525$; Mann-Whitney test) in relation to 45,X and other karyotypes $(−1.48 \pm 1.25 \times -1.74 \pm 1.46)$ (Table 2), the same occurring for BD-HA $(p=0.369$ – Mann-Whitney test) $(0.70 \pm 0.51 \times 0.13 \pm 1.54)$ (Table 2).

The mean of the Z-score of the BD-CA did not differ significantly ($p=0.734$; Mann-Whitney test) in relation to the median (group above vs. group below) of the BMI $Z$ score $(−1.48 \pm 0.76 \times -1.74 \pm 1.72)$ (Table 3), the same occurring for BD-HA $(p=0.697$; Mann-Whitney test) $(0.63 \pm 0.46 \times 0.19 \pm 1.57)$ (Table 3).

Discussion

The results found in this study indicate that, although patients with TS present compromised Z-score of BD for CA when correction for HA is performed, the results are within the expected range.

The association between TS and low bone mineral density/osteoporosis is still controversial in the literature. The low bone mass found in this group when evaluated using BD for CA seems to be associated with short stature, as described in other studies,6,15 since, when the result was corrected using HA, no patient was found with low BD. The same has been described in other studies that performed the correction of BD for HA.16,17 However, other authors have already reported that the BD alteration remained even after correction for HA and suggested that the risk of osteoporosis is greater in patients with TS due to early ovarian failure and possibly due to bone alterations resulting from haploinsufficiency of the gene $SHOX$.18

In our study, the two groups of karyotypes (45,X and karyotypes with mosaicism) were compared and no differences were found regarding the Z-score for height, BMI and BD results, which is in agreement with the literature.16,19,20 Short stature in TS can be largely attributed to the haploinsufficiency of the

| Table 1 |

| Distribution of chronological age (CA) and height age (HA) and Bone Densitometry (BD) by CA and HA of 13 prepubertal patients with Turner Syndrome. |

| Age (years) | BD (Z-score) |
|------------|--------------|
| CA         | HA           |
| Average    | 12.54        | 9.05         |
| Minimum    | 10.00        | 7.40         |
| Maximum    | 13.90        | 11.00        |
| Median     | -1.50        | 0.70         |
| $p$        | 0.001 (teste de Wilcoxon) | <0.001 (teste de Wilcoxon) |
Figure 1

Distribution of BD-CA and BD-HA in 13 prepubertal patients with Turner Syndrome.

Table 2

Z-score of bone densitometry for chronological age (CA) and height age (HA) in relation to karyotype.

| Karyotype  | Distribution of bone densitometry in relation to karyotype |
|------------|----------------------------------------------------------|
| 45,X       |                                                          |
| N          | Z_BMD CA*       | Z_BMD HA**       |
| 6          | -1.48          | 0.70             |
| Average    | -2.90          | -0.10            |
| Minimum    | -1.40          | 0.80             |
| Median     | 0.70           | 1.40             |
| Maximum    | 1.25           | 0.51             |
| Standard deviation | 1.25      | 1.54             |
| Mosaicism  |                                                          |
| N          | Z_BMD CA*       | Z_BMD HA**       |
| 7          | -1.74          | 0.13             |
| Average    | -3.90          | -2.00            |
| Minimum    | -1.80          | 0.30             |
| Median     | 0.20           | 2.40             |
| Maximum    | 1.46           | 1.54             |

*p=0.525 (Mann-Whitney test); **p=0.369 (Mann-Whitney test).
SHOX gene, which has been correlated with short stature and bone phenotype in TS.\(^3,16,20,21\)

There are studies that describe an increased risk of fractures associated with the syndrome, however with an inconclusive relationship with BD,\(^5,12,21\) since they present normal BD Z-score values after correction for HA, although clinically they present an increased risk for fractures.\(^16\)

Although some studies have already demonstrated a positive relationship between weight and BMD,\(^22,23,24\) in our study with BMI, no difference was observed between groups with higher or lower BMI values. These findings are similar to those of Costa \textit{et al.}\(^19\) Alghadir \textit{et al.}\(^13\) estimate a variation in bone mineral density (BMD) from 60% to 80%, that is influenced not only by ethnic and genetic factors, but also by modifiable environmental factors such as adiposity, hormone levels, sun exposure, blood pressure, physical activity, and diet. Body weight is a factor that would have a positive influence on BMD, although this relationship is not linear, as even in obese patients, BMD can be compromised.\(^22-24\)

The limitations of our study were the small sample and being a retrospective study, and not having a control group, which was partially compensated by the fact that all BD variables were analyzed by Z-score.

The strengths of the study were a group of patients in a limited age group, all prepubescent, and the fact that, because they were followed by the same group with specific guidelines on nutrition and physical activity, some of the factors that could negatively affect the mass bone were minimized.

There was no confirmed evidence of low bone mass in this group of patients with TS. Lower BD values are associated with short stature bias and when corrected for HA values are within normal limits. Karyotype and BMI do not modify the BD result in prepubertal patients with TS.

### Authors’ contribution

Dallago RT, Lemos-Marini SHV and Guerra-Junior G contributed to the planning, preparation and review of the study. Data collection was performed by Dallago RT, Marmos DB and Galon MV. Morcillo AM assisted in the statistical analysis. For help with techniques and with the densitometry device, we have the collaboration of Santos AO. The manuscript was written and revised by Dallago RT and Lemos-Marini SHV. All authors approved the final version of the manuscript and are publicly responsible for the content of the article.

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**Table 3**

Distribution of the bone densitometry Z-score in relation to BMI (median cut-off point = 0.61) for CA and HA of 13 prepubertal patients with Turner Syndrome.

| Median Z_BMI | Distribution of BD in relation to BMI using as a cutoff point the median=0.61 for CA and HA |
|--------------|---------------------------------------------------------------------------------------------|
| >0.61        | Z BMD CA*                                                                                   |
| N            | 6                                                                                           |
| Average      | -1.48                                                                                      |
| Minimum      | -2.50                                                                                      |
| Median       | -1.45                                                                                      |
| Maximum      | 0.720                                                                                      |
| Standard deviation | 0.76  |
| ≤0.61        | Z BMD HA**                                                                                |
| N            | 7                                                                                           |
| Average      | -1.74                                                                                      |
| Minimum      | -3.90                                                                                      |
| Median       | -1.80                                                                                      |
| Maximum      | 0.70                                                                                       |
| Standard deviation | 1.72  |

\(^*p=0.734\) (Mann-Whitney test); **\(^p=0.697\) (Mann-Whitney test); BD= bone densitometry; BMI= body mass index; CA=chronological age; HA= height age.
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