

Peripheral BMD T-Scores in the Diagnosis of Osteoporosis

Wanna Trivitayaratana MD, MSc*, Pichit Trivitayaratana MSc*

*Department of Radiological Technology, Faculty of Medical Technology, Mahidol University

The purposes of this study were to provide the reference value of peripheral bone mineral density from young healthy adults (20-40 years) in both sexes and to examine the prevalence of osteoporosis and fracture risk in normal young adults by T-scores following the World Health Organization (WHO) definition. Non-dominant sites of 4 peripheral skeletal of 1,128 young healthy women and 225 men were examined with the following techniques: (1) dual energy X-ray absorptiometry (DXA) at supradistal and distal 1/10 of forearm (2) digital X-ray radiogrammetry (DXR) of metacarpal and distal forearm and (3) quantitative ultrasound of the os calcis (stiffness index). The results showed that young adult mean (YAM) of all peripheral sites were similar in both sexes ($p > 0.05$ for all). Peak bone mass of all peripheral sites between Thai males and females were not different. $YAM \pm SD$ of women were used as Thai reference values and could be appropriately applied to men for individual calculation of T-scores. YAM at corresponding area of dominant and non-dominant sites of the above measurements ($n = 421, 183$ and 467 for forearm DXA, ultrasound heel and DXR, respectively) were not significantly different. It indicated that the BMD values of estimating bone mass would be correct whether the dominant or non-dominant site was measured. Applying the WHO definition of normal, osteopenia and osteoporosis to the T-scores level among 4 measurements in normal young adults for screening of peri-pheral osteoporosis, the prevalence of osteopenia and osteoporosis were 12.56-17.74% and 0.62-1.04%, respectively. There was a moderate fracture risk of 1.24-2.39% and marked risk of $< 0.5\%$

Keywords: Reference value, Peripheral BMD, Osteoporosis

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DXA is acceptably used as the standard method for diagnosis and monitoring of osteoporosis and for assessment of fracture risk. However, axial DXA is not suitable and available for mass screening because it is usually confined to specialized centers⁽¹⁾. Peripheral sites of BMD measurements may be useful to pre-select postmenopausal women in whom axial DXA is indicated to confirm/exclude osteoporosis at the spine or hip⁽²⁻⁸⁾. The non-uniform skeletal involvement in osteoporosis argues for multi-site evaluation^(9,10). Moreover, an inappropriate reference range for peak BMD may result in identification of an incorrect proportion of subjects with osteopenia and osteoporosis. Individual populations should use their own reference range T-scores to avoid misdiagnoses of osteopenia and osteoporosis⁽¹¹⁾. BMD at different sites of healthy young adult Thai are needed, unfortunately, the values have not been reported. So the authors revealed the reference database of 4 different

peripheral sites measured by DXA (Panasonic DXA-70), Ultrasound (Archillis ultrasound, Lunar) and combined radio-grammetric analysis with texture analysis (Pronosco X-posure system) or DXR in young adults (20-40 years) for T-scores in the diagnosis of osteoporosis in Thai people. YAM was calculated in both sexes because there was a different relationship between bone density and fractures in both sexes⁽¹²⁾.

Material and Method

The study included 1,128 women and 225 men, aged 20-40 years (mean aged 28.284 ± 4.160) who were healthy and randomly selected from the screening project of osteoporosis of the department of Radiological Technology, Faculty of Medical Technology, Mahidol University from 1998 to 2001. Exclusion criteria were traumatic fractures, surgical menopause, disease or conditions known to affect bone, e.g. endocrine disease, rheumatoid arthritis, hyperparathyroidism, chronic disease, medication (current or past treatment with corticosteroids, fluoride, calcitonin, bisphosphonates). Conventional radiographs of the lateral lumbar spine were evaluated for degenera-

Correspondence to : Trivitayaratana W, Department of Radiological Technology, Faculty of Medical Technology, Mahidol University, Bangkok 10700, Thailand. Phone: 0-6791-5950, E-mail wanna_kae@hotmail.com

tive changes. Fractured vertebral bodies, if present, were excluded from the analysis. All subjects were examined at the non-dominant site with a DXA scanner of forearm, stiffness at calcaneus and DXR of hand and forearm. 114 women and 27 men could not be scanned at the non-dominant forearm because of the problem of a foreign body. 192 women and 39 men radiographs of hand including forearm for DXR measurement were also excluded because of their exposure problems. Patients were excluded if they had evidence of any secondary cause of osteoporosis and could not be scanned by any peripheral measurements. For comparison of non-dominant and dominant sites of peripheral BMD, DXA forearm, ultrasound heel and DXR measurements were also performed at both sites of 421, 183 and 467 subjects, respectively.

Forearm bone mineral density values by DXA were given in grams per centimeter squared, and the individual subjects results were expressed as T-scores. Distal forearms were scanned and BMD at 4 different regions of interest (supradistal, distal 1/10, distal 1/6 and distal 1/3) were computed. Only two regions of forearm BMD were reported because BMD at supradistal and distal 1/10 of radius had been proved a strong correlation with BMD of lumbar spine, hip, femoral neck and Ward's triangle⁽¹³⁾. During the same session, the subjects were examined at the calcaneus with a quantitative sonography device, the Achilles, according to the manufacturer's recommended standard procedures. The sonography derived stiffness of the calcaneus was automatically determined by the scanner software according to the following formula: stiffness = $(0.67 \times \text{broadband ultrasound attenuation}) + (0.28 \times \text{speed of sound}) - 420$ ⁽¹⁴⁾. Stiffness is the default parameter used by the manufacturer for demographic comparison of the patient's data, and, therefore, a T-score is given.

The recent development of computed radiogrammetric analysis of a plain radiograph of the hand and forearm was a valuable tool for quantitating

bone mass^(15,16). Cortical bone loss was monitored by using the DXR method.

The diagnostic bone mass threshold for defining osteoporosis in individuals without fracture was recently clarified by the WHO⁽¹⁷⁾ to be T-scores less than -2.5 ⁽¹⁸⁾. WHO also defined people whose BMD was normal and osteopenia if T-scores > -1 and $-2.5 < \text{T-scores} < -1$, respectively. T-scores of DXA were used to define moderate fracture risk by T-scores < -2 and marked risk by T-scores < -3 ⁽²⁾.

Statistics

YAM \pm SD and range for reference BMD values were calculated. For comparing the means of peripheral BMDs between females and males, Student's t test was used. BMD at dominant and non-dominant of peripheral sites were compared using paired t test. Prevalence of percentage of normal, osteopenia, osteoporosis and fracture risks in healthy Thais was computed. All descriptive and inferential statistics were performed by SPSS/PC software.

Results

1,353 healthy young adults, 1,128 females and 225 males were studied aged ranging from 20-40 years. YAM \pm SD of DXA BMD at supradistal and distal 1/10 of forearm, ultrasound heel and digital radiogrammetry of hand and forearm were calculated in both sexes as shown in Table 1. The Student's t test was used to compare YAM of each measurements in females and males. The results showed that YAM of all peripheral sites were similar in both sexes ($p > 0.05$ for all). From paired t-test in Table 2, YAM at the corresponding area of non-dominant and dominant sites of the 4 measurements were not significantly different ($p > 0.05$ for all). T-scores of individual BMD were computed using the reference YAM in Table 1. Table 3 shows the number and percentage of 3 levels of T-scores among 4 different measurements in normal young adults. There were

Table 1. YAM \pm SD and range of peripheral BMD in females and males

Measurement technology	sites	n	Females YAM \pm SD	range	n	Males YAM \pm SD	range	p
DXA	Supradistal of forearm	1,014	0.4590 \pm 0.032	0.322-0.663	198	0.5589 \pm 0.031	0.446-0.683	0.186
DXA	Distal 1/10 of forearm	1,014	0.5405 \pm 0.025	0.415-0.710	198	0.6191 \pm 0.027	0.558-0.713	0.121
Ultrasound	Calcaneus	1,128	92.4043 \pm 7.258	30-127	225	101.3600 \pm 9.114	61-139	0.124
DXR	Radius, ulna and three middle metacarpals	936	0.5538 \pm 0.030	0.482-0.642	186	0.5871 \pm 0.022	0.521-0.662	0.143

Table 2. Comparison of non-dominant and dominant sites of peripheral BMD measurements

Measurement techniques	$\bar{x} \pm SD$		p	
	n	Non-dominant		Dominant
DXA forearm (supradistal)	421	0.519±0.059	0.524±0.059	0.119
DXA forearm (distal 1/10)	421	0.471±0.074	0.483±0.067	0.132
Ultrasound calcaneus	183	91.445±14.365	91.192±14.449	0.821
DXR	467	0.560±0.032	0.568±0.032	0.415

Table 3. T-scores level among 4 different measurements in normal young Thai adults

Level of T-scores	Supradistal		Distal 1/10		Ultrasound calcaneus		DXR	
	n	%	n	%	n	%	n	%
> -1	1,025	84.57	986	81.35	1,169	86.40	960	85.56
-1 < and > -2.5	176	14.52	215	17.74	170	12.56	155	13.82
< -2.5	11	0.91	11	0.91	14	1.04	7	0.62
Total	1,014		1,014		1,128		936	

Table 4. Fracture risk level among 4 different measurements in normal young adults

Fracture risk	SDs below YAM	Supradistal		Distal 1/10		Ultrasound calcaneus		DXR	
		n	%	n	%	n	%	n	%
moderate	2	24	1.98	29	2.39	23	1.69	14	1.24
marked	3	2	0.17	2	0.17	4	0.30	0	0.00

>80% normal bone mass and <1.1% osteoporosis. <3% moderate fracture and <0.5% marked fracture risk are also shown in Table 4.

Discussion

Female and male's YAM were not significantly different as shown in Table 1. Peak bone mass of all peripheral sites in Thai males were not different from Thai females. Although axial BMD in European men were much more than women⁽¹²⁾. The current WHO definition of osteoporosis in postmenopausal women can be appropriately applied to men. The authors suggested that YAM \pm SD of women were used as Thai reference values and could be appropriately applied to men for individual calculation of T-scores.

The standard peripheral site for screening of bone mass is the non-dominant site. Sometimes it was not practical for routine measurement because a foreign body was hard to remove. The present study showed that BMD at corresponding sites of non-dominant and dominant peripheral bone were similar. It indicated that the BMD value of estimating bone mass was correct whether the non-dominant or dominant site was measured or not.

T-scores level among 4 measurements in normal young adults in Table 3 shows 81.35-86.40% normal, 12.56-17.74% osteopenia and 0.62-1.04%

osteoporosis. In comparison to the screening test for osteoporosis in 2,789 women aged 51-75 years, 34% of patients were osteoporotic, 42% were osteopenic and 24% had normal bone density results performed by spinal DXA⁽¹⁹⁾. Fracture risk level in normal young Thais were 1.24-2.39% moderate and <0.5% marked fracture risk. Although the occurrence of osteoporosis and marked fracture risk were low in normal young adults, maintenance of peak bone mass since young adult life by promoting bone formation and reducing bone resorption should be regarded for the prevention of osteoporosis fracture.

There is a need for low cost screening methods to detect low bone mass in postmenopausal women. Peripheral measurement techniques are attractive because the equipment cost is substantially lower, radiation exposure is small, the devices require less space and sometimes they are portable⁽²⁰⁾. Computed radiogrammetry of appendicular bone densitometric method^(16,21), forearm DXA^(8,22,23), stiffness index of the os calcis^(24,25) confirmed their predictive ability for fragility fractures.

The BMD values are expressed in terms of the number of standard deviations above or below the young normal value (commonly referred to as the T-scores). If the normative populations of the various systems are consistent, the standard deviation scores

should also be consistent. For this reason, the WHO recently established diagnostic criteria for osteoporosis base on T-scores but not BMD⁽²⁶⁾. The patients were categorized into three diagnoses: normal, osteopenia and osteoporosis⁽²⁷⁾. The classification of osteoporosis varies according to skeletal site⁽¹⁰⁾, which is affected by multi-factors (aging, calcium intake, estrogen, life style, geographic area, etc.). The reliable prevalence of accurate diagnosis of osteoporosis and fracture risk for Thai people require Thai reference database of T-scores. In the present research, young healthy adults mean and standard deviation of bone density of 4 different peripheral measurements were established for the calculation of T-scores. They were useful for diagnosis and assessment of the fracture risk of osteoporosis in a Thai population. Because of the fracture risk increase with decreasing BMD, these important findings may lead to public health planning in promoting and preventing a health policy.

References

1. Lippuner K, Fuchs G, Ruetsche AG. How well do radiographic absorptiometry and quantitative ultrasound predict osteoporosis at spine or hip? A cost effectiveness analysis. *J Clin Densitom* 2000; 3: 241-9.
2. Swezey RL, Draper D, Swezey AM. Bone densitometry: comparison of dual energy x-ray absorptiometry to radiographic absorptiometry. *J Rheumatol* 1996; 23: 1734-8.
3. Benitez CL, Schneider DL, Barrett-Connor E, Sartoris DJ. Hand ultrasound for osteoporosis screening in postmenopausal women. *Osteoporos Int* 2000; 11: 203-10.
4. Takada M, Engelke K, Hagiwara S. Assessment of osteoporosis: comparison of radiographic absorptiometry of the phalanges and dual X-ray absorptiometry of the radius and lumbar spine. *Radiology* 1997; 202 : 759-63.
5. Mulder JE, Michaeli D, Flaster E, Siris E. Comparison of bone mineral density of the phalanges, lumbar spine, hip and forearm for the assessment of osteoporosis in postmenopausal women. *J Clin Densitom* 2000; 3: 373-81.
6. Ayers M, Prince M, Ahmadi S, Baran DT. Reconciling quantitative ultrasound of the calcaneus with x-ray base measurements of the central skeleton. *J Bone Miner Res* 2000; 15: 1850-5.
7. Frost ML, Blake GM, Fogelman I. Contact quantitative ultrasound: an evaluation of precision, fracture discrimination, age-related bone loss and applicability of the WHO criteria. *Osteoporos Int* 1999; 10: 441-9.
8. Jones T, Davie MW. Bone mineral density at distal forearm can identify patients with osteoporosis at spine and femoral neck. *Br J Rheumatol* 1998; 37: 539-43.
9. Weiss M, Ben-shlomo AB, Hagag P, Rapoport M. Reference database for bone speed of sound measurement by a novel quantitative multi-site ultrasound device. *Osteoporos Int* 2000; 11: 688-96.
10. Sahata O, Pearson D, Cawte SW. Site-specific variation in the classification of osteoporosis, and the diagnostic reclassification using the lowest individual lumbar vertebra T-scores compared with L₁-L₄ mean, in early postmenopausal women. *Osteoporos Int* 2000; 11: 852-7.
11. Gurlek A, Bayraktar M, Ariyarek M. Inappropriate reference range for peak bone mineral density in dual-energy x-ray absorptiometry: implication for the interpretation of T-scores. *Osteoporos Int* 2000; 11: 809-13.
12. Selley PL, Davies M, Adams JE. Do men and women fracture bones at similar bone densities? *Osteoporos Int* 2000; 11: 153-7.
13. Trivitayaratana W, Trivitayaratana P, Kongkiatikul S. Prediction of bone mineral density of lumbar spine, hip, femoral neck and Ward's triangle by forearm bone mineral density. *J Med Assoc Thai* 2001; 84: 390-6.
14. Hans D, Schott AM, Chapuy MC. Ultrasound measurements on the os calcis in a prospective multi center study. *Calcif Tissue Int* 1994; 55: 94-9.
15. Rico H, Revilla M, Villa LF. Comparison between metacarpal bone measurements by computerized radiogrammetry and total body DEXA in normal and osteoporotic women. *Clin Rheumatol* 1994; 13: 593-7.
16. Jorgensen JT, Anderson PB, Rosholm A, Bjarnason NM. Digital X-ray radiogrammetry: a new appendicular bone densitometric method with high precision. *Clin Physiol* 2000; 20: 330-5.
17. Kanis JA. Assessment of fracture risk and its application to screening for postmenopausal osteoporosis: synopsis of a WHO report. *Osteoporos Int* 1994; 4: 368-81.
18. Miller PD, Bonnick SL, Rosen CJ. Consensus of an international panel on the clinical utility of bone mass measurements in the detection of low bone mass in the adult population. *Calcif Tissue Int* 1996; 58: 207-14.

19. Saadi H, Litaker D, Mills W. Practice variation in the diagnosis and treatment of osteoporosis: a case for more effective physician education in primary care. *J Women Health Gend Based Med* 1996; 8: 767-71.
20. Gluer CC, Jergas M, Hans D. Peripheral measurement techniques for the assessment of osteoporosis. *Semin Nucl Med* 1997; 27: 229-47.
21. Adami S, Zamborlan N, Gatti D. Computed radiographic absorptiometry and morphometry in the assessment of postmenopausal bone loss. *Osteoporos Int* 1996; 6: 8-13.
22. Augat P, Fuerst T, Genant HK. Quantitative bone mineral assessment at the forearm: a review. *Osteoporos Int* 1998; 8: 299-310.
23. Duppe H, Gardsell P, Nilsson B, Johnell O. A single bone density measurement can predict fracture over 25 years. *Calcif Tissue Int* 1997; 60: 171-4.
24. Cunningham JL, Fordham JN, Hewitt TA, Speed CA. Ultrasound velocity and attenuation at different skeletal sites compared with bone mineral density measured using dual energy x-ray absorptiometry. *Br J Radiol* 1996; 69: 25-32.
25. Grampp S, Henk CB, Fuerst TP. Diagnostic agreement of quantitative sonography of the calcaneus with dual x-ray absorptiometry of the spine and femur. *AJR* 1999; 173: 329-34.
26. Faulkner KG, Roberts LA, McClung MR. Discrepancies in normative data between Lunar and Hologic DXA systems. *Osteoporos Int* 1996; 6: 432-6.
27. Woodson G. Dual x-ray absorptiometry T-scores concordance and discordance between the hip and spine measurement sites. *J Clin Densitom* 2000; 3: 319-24.

ความหนาแน่นของกระดูกกระยางค์สำหรับคำนวณ T-scores ในการวินิจฉัยโรคกระดูกพรุน

วรรณา ตริวิทย์รัตน์, พิชิต ตริวิทย์รัตน์

วัตถุประสงค์ของงานวิจัยนี้คือ หาค่าอ้างอิงของความหนาแน่นของกระดูกกระยางค์ จากผู้ใหญ่วัยหนุ่มสาว (20-40 ปี) ที่มีสุขภาพแข็งแรง ทั้งเพศชายและหญิง และเพื่อหาอัตราอุบัติการณ์เกิดโรคกระดูกพรุนและความเสี่ยงต่อกระดูกหักจากค่า T-scores ที่นิยามโดยองค์การอนามัยโลก ทำการวัดกระดูกกระยางค์ 4 ตำแหน่งเฉพาะข้างที่ไม่ถนัดของผู้หญิง 1,128 คนและผู้ชาย 225 คน โดยใช้วิธีวัดดังนี้ (1) การดูคลื่นเอกซเรย์ (2) ค่าพลังงานที่กระดูกปลายแขนส่วนปลายสุดและส่วนปลาย 1/10 (3) รูปร่างของกระดูกฝ่ามือและปลายแขน และ (3) คลื่นเสียงความถี่สูงที่สันเท้า ผลการวิจัยพบว่าค่าเฉลี่ยของความหนาแน่นของกระดูกกระยางค์ทุกแห่งในผู้หญิงและผู้ชาย ไม่พบความแตกต่าง ($P > 0.05$ ทุกตำแหน่ง) นั่นคือมวลกระดูกสูงสุดของกระดูกกระยางค์ในผู้หญิงและผู้ชายไม่ต่างกัน จึงสามารถใช้ค่าเฉลี่ยมวลกระดูกในผู้หญิงเป็นค่าอ้างอิงของคนไทย และสามารถประยุกต์ใช้กับการคำนวณ T-scores ในผู้ชายได้ ค่าเฉลี่ยความหนาแน่นของกระดูกกระยางค์ข้างที่ไม่ถนัดและถนัดทั้ง 4 ตำแหน่ง ($n = 421, 183$ และ 467 สำหรับการวัดการดูคลื่นเอกซเรย์ 2 ค่าพลังงานที่ปลายแขน, คลื่นเสียงความถี่สูงที่สันเท้าและวัดรูปร่างของกระดูกตามลำดับ) ไม่แตกต่างกัน แสดงว่าค่าความหนาแน่นของกระดูกที่ได้จะถูกตั้งไม่ว่าจะวัดข้างที่ไม่ถนัดหรือถนัด จากการใช้นิยามขององค์การอนามัยโลก พบว่าในวัยหนุ่มสาวไทย มีอุบัติการณ์เกิดกระดูกบางร้อยละ 12.56-17.74 เกิดโรคกระดูกพรุน ร้อยละ 0.62-1.04 ความเสี่ยงต่อการเกิดกระดูกหักระดับปานกลาง พบร้อยละ 1.24-2.39 และระดับสูงพบร้อยละ < 0.5
