Osteoporosis and Sarcopenia 6 (2020) 168-172

Contents lists available at ScienceDirect

Osteoporosis and Sarcopenia

journal homepage: http://www.elsevier.com/locate/afos

Original article

Different reference ranges affect the prevalence of osteoporosis and osteopenia in an urban adult Malaysian population



Osteoporosis Sarcopenia

Swan Sim Yeap ^{a, b, *}, Subashini C. Thambiah ^c, Intan Nureslyna Samsudin ^c, Geeta Appannah ^d, Nurunnaim Zainuddin ^c, Safarina Mohamad-Ismuddin ^c, Nasrin Shahifar ^c, Salmiah Md-Said ^e, Siti Yazmin Zahari-Sham ^c, Subapriya Suppiah ^f, Fen Lee Hew ^{a, b}

^a Puchong Specialist Centre, Puchong, Selangor, Malaysia

^b Department of Medicine, Subang Jaya Medical Centre, Subang Jaya, Selangor, Malaysia

^c Department of Pathology, Faculty of Medicine and Health Sciences, Universiti Putra Malaysia, Selangor, Malaysia

^d Department of Nutrition and Dietetics, Faculty of Medicine and Health Sciences, Universiti Putra Malaysia, Selangor, Malaysia

^e Department of Community Health, Faculty of Medicine and Health Sciences, Universiti Putra Malaysia, Selangor, Malaysia

^f Department of Imaging, Faculty of Medicine and Health Sciences, Universiti Putra Malaysia, Selangor, Malaysia

A R T I C L E I N F O

Article history: Received 28 June 2020 Received in revised form 16 September 2020 Accepted 17 November 2020 Available online 27 November 2020

Keywords: Osteoporosis Osteopenia Reference ranges Malaysia

ABSTRACT

Objectives: To determine the prevalence of osteopenia (OPe) and osteoporosis (OP) in an urban adult population in Malaysia, and to compare the change in the prevalence when using a Caucasian compared to an Asian reference range.

Methods: A cross-sectional random sample of the population aged between 45 and 90 years from the state of Selangor, Malaysia, was invited to attend a bone health check-up. Participants with diseases known to affect bone metabolism or who were on treatment for OP were excluded. Bone mineral density was measured using dual energy X-ray absorptiometry. Based on the World Health Organization definitions, the prevalence of OPe and OP was calculated using the Asian and Caucasian T-scores.

Results: A total of 342 subjects (222 females, 120 males), with a mean age of 59.68 (standard deviation: 8.89) years, who fulfilled the study criteria were assessed. Based on the Asian reference range, there were 140 (40.9%) subjects with OPe and 48 (14.0%) with OP. On applying the Caucasian reference range, there were 152 (44.4%) subjects with OPe and 79 (23.1%) with OP, with significant increases in males, females, and Chinese ethnic groups. Overall, 75 (21.9%) of subjects had a change in their diagnostic status. T-scores were consistently lower when the Caucasian reference range was used.

Conclusions: In a healthy urban Malaysian population, the prevalence of OP is 14.0% and OPe is 40.9%. Application of a Caucasian reference range significantly increased the number of subjects with OP and may potentially lead to over-treatment.

© 2020 The Korean Society of Osteoporosis. Publishing services by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

1. Introduction

Osteoporosis (OP) is defined as a skeletal disorder characterized by compromised bone strength predisposing to an increased risk of fracture [1]. The term bone strength is used to include both bone mineral density (BMD) and bone quality. Although we cannot

E-mail address: swanyeap@gmail.com (S.S. Yeap).

Peer review under responsibility of The Korean Society of Osteoporosis.

directly measure bone quality, BMD can be readily measured using dual-energy X-ray absorptiometry (DXA). The purpose of such assessments would be to identify those at risk of an osteoporotic fracture, and/or to determine in whom to start treatment. The World Health Organisation (WHO) defined OP as a BMD ≥ 2.5 SD below the mean for healthy young women (ie, T-score ≤ -2.5) at any site (spine, hip or 1/3 radius), with the mean based on American Caucasian female data. This level would identify 30% of all postmenopausal women as having OP, of which more than half would have sustained a previous osteoporotic fracture [2]. Thus, OP can be diagnosed following a fragility fracture, or when there is a T-score of ≤ -2.5 as measured by DXA [3].

https://doi.org/10.1016/j.afos.2020.11.005

^{*} Corresponding author. Department of Medicine, Subang Jaya Medical Centre, No. 1, Jalan SS 12/1A, 47500, Subang Jaya, Malaysia.

^{2405-5255/© 2020} The Korean Society of Osteoporosis. Publishing services by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (http:// creativecommons.org/licenses/by-nc-nd/4.0/).

Table	1
-------	---

Characteristics of study cohort.

Variable	Male ($n = 120, 35.1\%$)	Female (n = 222, 64.9%)	P- value
Age, yr	59.70 ± 9.51	59.67 ± 8.57	0.974 ^a
Race:			
Malay (78)	30 (25.0%)	48 (21.6%)	0.624 ^b
Chinese (157)	51 (42.5%)	106 (47.7%)	
Indian (107)	39 (32.5%)	68 (30.6%)	
Menopause	NA	191 (86.0%)	
LS A T-score	0.008 ± 1.431	-0.748 ± 1.315	< 0.001 ^a
LS C T-score	-0.600 ± 1.555	-1.167 ± 1.366	0.001 ^a
FN A T-score	-0.537 ± 1.269	-1.150 ± 1.200	< 0.001 ^a
FN C T-score	-1.104 ± 1.164	-1.524 ± 1.136	0.002 ^a
TH A T-score	-0.144 ± 1.134	-0.374 ± 1.145	0.086 ^a
TH C T-score	-0.732 ± 0.997	-1.107 ± 1.090	0.003 ^a
BMD Category (A)			
Normal	74 (61.7%)	80 (36.0%)	< 0.001 ^b
Osteopenia	37 (30.8%)	103 (46.4%)	
Osteoporosis	9 (7.5%)	39 (17.6%)	
BMD Category (C)			
Normal	53 (44.2%)	58 (26.1%)	0.002 ^b
Osteopenia	48 (40.0%)	104 (46.9%)	
Osteoporosis	19 (15.8%)	60 (27.0%)	

Values are presented as mean \pm standard deviation or number (%).

NA, not applicable; LS, lumbar spine; FN, femoral neck; TH, total hip; A, Asian; C, Caucasian; BMD, bone mineral density.

^a One way ANOVA.

^b Chi-square.

A group from the WHO Collaborating Center for Metabolic Bone Disease [4] and the International Osteoporosis Foundation (IOF) [5] have suggested that the femoral neck (FN) BMD measured with DXA to be used as the reference standard for diagnosing OP in both genders and all ethnic groups, using T-scores derived from the United States (US) National Health and Nutrition Examination Survey (NHANES) III database of Caucasian women aged 20-29 years. However, many studies have shown that there are ethnic differences in BMD [6,7]. Generally, Asian populations were found to have lower BMDs compared to Caucasian populations [8,9]. The BMD reference curves for Chinese women have been found to be lower compared to Japanese and Caucasian women. Chinese women had peak BMD at the lumbar spine (LS) and hip regions that were 2.7–7.9% lower than those of Japanese women and 0.5–7.2% lower than those of Caucasian women [10]. Even within the same country, the BMD of Cambodian, Laotian, and Vietnamese women in Rochester, Minnesota, US was found to be lower than those of Caucasian women living in the same area [11]. Thus, the mean BMD of a healthy young Asian woman would be lower than that of a Caucasian. As a result, if a Caucasian reference range is used, the Tscore will be lower compared to a local country-specific reference range. Several other studies from Asian countries such as Korea [12], Thailand [13], and Vietnam [14] have shown that using a

 Table 2

 The prevalence of normal, osteopenia and osteoporosis in the different ethnic groups.

Variable	Malay (n=78)	Chinese ($n = 157$)	Indian ($n = 107$)
BMD Category (A) ^a			
Normal	42 (53.8%)	49 (31.2%)	63 (58.9%)
Osteopenia	29 (37.2%)	73 (46.5%)	38 (35.5%)
Osteoporosis	7 (9.0%)	35 (22.3%)	6 (5.6%)
BMD Category (C) ^a			
Normal	34 (43.6%)	32 (20.4%)	45 (42.1%)
Osteopenia	31 (39.7%)	72 (45.9%)	49 (45.8%)
Osteoporosis	13 (16.7%)	53 (33.7%)	13 (12.1%)

Values are presented as number (%).

A, Asian; C, Caucasian.

^a Chi-square P < 0.001 when comparing the different ethnic groups.

Caucasian/USA reference range leads to a higher incidence of OP.

This is the first study to examine the prevalence of OP and osteopenia (OPe) in a multi-ethnic Asian urban population in Malaysia, and to compare the change in prevalence with the use of a Caucasian, compared to an Asian, BMD reference range to determine the T-score.

2. Methods

2.1. Study design and study location

A cross-sectional study was conducted in 3 selected residential areas in Puchong, Serdang, and Kajang, in the state of Selangor, Malaysia. Data were collected from June 2016 to August 2018.

2.2. Subject sampling

All adults aged 45 and above from randomly selected addresses from the voters' registry were invited to participate in this study. The selection of respondents used stratified random sampling with equal proportions. In addition, systematic random sampling was performed to select adequate respondents in each ethnic stratum. Research assistants distributed house-to-house brochures with details of the research project by hand. Potential subjects were screened when they called for an appointment. The inclusion criteria were those aged between 45 and 90 years and belonging to the Malay, Chinese or Indian ethnic groups. The exclusion criteria were subjects already diagnosed with OP, were taking/had taken medication for OP (including calcitriol or alfacalcidol), have a known secondary cause of OP, subjects with renal impairment (estimated glomerular filtration rate $< 60 \text{ mL/min}/1.73 \text{ m}^2$), known to have or had metabolic bone disorders, malabsorption, thyroid disease, immobilization or taking other drugs which affected bone homeostasis (eg, corticosteroids, phenytoin, methotrexate, cyclosporine, oral contraceptive pill) or subjects who had a computed tomography scan in the past 1 year.

Eligible subjects were scheduled for a face-to-face clinical assessment and BMD measurement by DXA. The study protocol

was approved by the Ethics Committee of Universiti Putra Malaysia, approval reference FPSK (FR16) P002 dated 9th May 2016. All subjects signed written informed consent.

2.3. Anthropometric parameters and BMD measurement

Anthropometric measurements, ie, height, weight, and body mass index (BMI), were performed for all subjects. The designated personnel measured the height of the subjects, who were required to stand barefoot on the base of the stadiometer. The weight of the subjects was also taken at the same time. The height was measured in centimetres (cm) and the weight in kilograms (kg) and recorded to the approximate value of one decimal point. Subsequently, the BMI was calculated as weight/height².

The DXA was performed using a HOLOGIC Discovery W densitometer (Hologic Corporation, Bedford, MA, USA) to measure the BMD of the lumbar spine (LS) that represented the mean value of L1 to L4, the left femoral neck (FN), and the left total hip (TH). The precision of the machine is $\pm 2\%$. A calibration using the phantom was done before the first scan of each clinic session. The reference populations used were the manufacturer's White (Caucasian) and Asian (Japanese) population databases.

BMD was classified into normal, OPe, and OP based on T-scores using the WHO classification [2]. A T-score greater than -1.0 was classified as normal, between -1 and -2.5 was classified as OPe, and less than-2.5 was classified as OP.

2.4. Plain radiograph assessment of the lumbar spine

The antero-posterior and lateral lumbar spine radiographs were taken with the patient in the erect position using a portable X-ray machine (Toshiba 25 kW Radiography System, Tokyo, Japan). The radiation exposure was 40–80 mAs and 70–80 kVp depending on the body habitus of the subjects. The diagnosis of a vertebral fracture was made using a semi-quantitative technique, based on the Genant classification of vertebral fractures in OP [15]. A morphometric fracture was diagnosed based on a greater than 25% reduction in the anterior, central, and/or posterior segment height of the vertebra relative to the adjacent vertebra accompanied with a reduction in the area.

2.5. Statistical analysis

Analysis was performed using IBM SPSS Statistics for Windows, Version 24.0 (IBM Corp, Armonk, NY, USA). As all data were normally distributed, mean and standard deviation (SD) were used for the continuous variables. One way ANOVA, *t*-tests and chi-square tests were used to determine association between variables. A P value of < 0.05 was considered to be statistically significant.

3. Results

BMD was measured in 386 subjects. A total of 342 subjects were studied after 44 subjects who were found to have vertebral fractures were excluded. There were 222 females (64.9%) and 120 males (35.1%). The mean age of the group was 59.68 (SD: 8.89) years. The majority of the subjects were of the Chinese ethnic group (45.9%), followed by Indian (31.3%), and Malay (22.8%). The characteristics of subjects are shown in Table 1. Blood tests performed showed normal full blood counts, renal, liver, and bone profiles for the subjects (data not shown).

In Malaysia, when analyzing BMD measured by DXA, it is recommended that race-specific reference ranges are used when available [16]. As there is no Malaysian reference range, the manufacturer's Asian reference range is normally used to calculate the T- and Z-scores in the DXA printouts. As shown in Table 1, mean Tscores were significantly lower in females compared to males at the LS and FN but not TH when the Asian reference range was used. When the White (Caucasian) range was applied to the data, the mean T-scores were significantly lower in females compared to males in all 3 sites. Mean T-scores were significantly lower at each site in both males and females when the Asian T-scores were compared to Caucasian T-scores (T test P \leq 0.001 at LS, FN and TH).

Based on the Asian reference range, there were 140 (40.9%) subjects with OPe and 48 (14.0%) with OP. After applying the Caucasian reference range, there were 152 (44.4%) subjects with OPe and 79 (23.1%) with OP. This represented an 8.6% (12/140) increase in subjects with OPe and a 64.6% (31/48) increase in subjects in OP. Overall, 75 (21.9%) subjects had their diagnostic status changed. The proportion of male and female subjects in each BMD category based on the Caucasian or Asian range is shown in Table 1. In addition, there was a significant increase in the percentage of subjects with OP in both males and females when the Caucasian reference range was applied, compared to using the Asian reference range (chi-square P = 0.044 in males and P = 0.017 in females).

Table 2 shows the prevalence of normal, osteopenic and osteoporotic BMD in the 3 ethnic groups in Malaysia using the Asian and Caucasian reference ranges. There was an increase in the prevalence of OP when the Caucasian reference range was used compared to the Asian reference range in all 3 ethnic groups, but this was only significant in Chinese (chi-square P = 0.151 in Malay, P = 0.024 in Chinese, and P = 0.093 in Indian).

4. Discussion

The WHO defined OP on the basis of a value for BMD 2.5 SD or more below the young female adult mean (T-score less than or equal to -2.5 SD) based on DXA measurements [2].

For the purpose of standardization of research studies on OP, it was decided by WHO [4] and IOF [5] that the recommended reference range is the NHANES III reference database for FN measurements in Caucasian women aged 20–29 years, for both women and men. The International Society for Clinical Densitometry (ISCD) advises to use a uniform Caucasian (non-race adjusted) female normative database for women and men of all ethnic groups [17], which is similar to the other 2 organizations. In the discussion from the paper from Kanis and colleagues [4], they mention that differences in BMD in different regions of the world only vary by approximately 1 SD. These small variations in BMD between populations therefore appear to be substantially less than variations in fracture risk. For example in China and India, hip fracture risk is lower than in Europe or the US [18], despite generally lower BMD in Asia [8–10]. They conclude that in view of the disparity between population fracture risks and BMD, it is uncertain whether reference ranges drawn from local populations would be of any added value. This is especially true if the local population BMD data is not linked to robust fracture risk data.

The mean BMD of the young healthy reference group used for the calculation of the T-score is related to the achievement of peak bone mass in the population. Studies in Asian populations have shown that BMD at peak bone mass can be lower [19,20] or occur at different ages, including later than that achieved in Caucasians [21]. This would be another reason that the T-scores would be different depending on the reference range used.

In this study, we found that the prevalence of OP significantly increased in both males and females when a Caucasian reference range was applied; the prevalence of OP increased from 7.5% to 15.8% in males and from 17.6% to 27.0% in females when the Asian and Caucasian reference range were applied, respectively. This was associated with a consistent reduction in the mean T-scores at all

measured sites. This is similar to several other Asian studies, which have shown that using a Caucasian database in Asian patients will lead to an increase in the diagnosis of OP. In a study from Thailand looking at women aged between 40 and 80 years, the number of women with OP increased when the USA reference range from the DXA machine was used compared to a local Thai population database [13]. This increase was seen in all age groups and also both at the LS and FN. For example, the prevalence of OP at the FN in 60–64 year olds increased from 20.1% using a Thai reference range to 52.6% using the USA reference range. A study from Vietnam that examined BMD in men and women between the ages of 18-89, found that in those over the age of 50, the proportion of subjects with OP increased when the DXA machine reference range was used compared to a Vietnamese reference range [14]. The number of subjects with FN OP increased from 28.6% to 43.7% in women and from 10.4% to 29.6% in men when the DXA machine reference range was used compared to the Vietnamese reference range, respectively. This is similar to our study showing increased prevalence of OP in both men and women using the DXA machine (Caucasian) reference range. However, not all studies have shown that the nonlocal reference range leads to an increased diagnosis of OP. In a study from Korea, when they compared OP prevalence based on using the NHANES database compared to the Korean reference range, they found that the prevalence of OP in men was less when the NHANES reference was used, but increased in women [12]. Another factor in this discussion is that even within Asia itself, there are differences in the BMDs of different populations. A study from Korea compared the prevalence of OP based on the Japanese reference standard given in the DXA machine and the local Korean reference ranges, and found that the prevalence varied depending on gender and site [21]. Thus, even in countries that are geographically close together, there is a difference in BMD and hence T-scores. Similarly, in a study from Sri Lanka, it was found that the Asian BMD reference values provided by the DXA machine manufacturer was lower than the BMD values found in their crosssectional randomly sampled community survey [20]. Therefore, as long as the diagnosis of OP is based on a T-score value of ≤ -2.5 , with the implication that it is a treatment threshold, the reference range used to calculate the T-score remains important. Using a reference range that increases the prevalence of OP would mean potentially many more people being treated, thus raising the spectre of over-treatment.

This study has also provided information on the prevalence of OP and OPe in the community in a multi-ethnic Malaysian population. There have only been a handful of previous studies. In a similar study looking at an urban population, the prevalence of OP was 8.4% in males and 16.1% in females [22], comparable to our results of 7.5% in males and 17.6% in females. However, the investigators used a Singapore reference range to calculate the Tscores. Our study also shows that the Chinese have the highest proportion of OP, 22.3% using the Asian database, out of the 3 ethnic groups. This is slightly higher than the prevalence of 15.8% found in a study from Malaysia that used a Singapore reference range [23]. The higher prevalence of OP in the Chinese in Malaysia was also shown in a study published in 2005 that did not specify which reference range that was used to calculate the T-score. In that study, the prevalence of FN OP was highest in the Chinese, 24.8%, followed by Malay, 14.8%, and Indian, 9.1% [24], a pattern similar to our study. This lower BMD in the Chinese may be a factor in the increased risk of hip fractures found in the Malaysian Chinese population [25].

There are a few limitations to this study. Firstly, our population may not be fully representative of the Malaysian population in terms of the ethnic breakdown of the subjects. We had 20.2% (76/386) Malay, 40.7% (157/386) Chinese, and 27.7% (107/386) Indian subjects. This is a higher proportion of Chinese and Indian subjects

compared to the population in the state of Selangor, where the proportion of ethnic groups were 54.1% Malays, 24.0% Chinese, and 11.2% Indians [26]. Further studies will be needed to fully examine the BMD distribution in Malays. In addition, because we specifically wanted to look at a healthy population, we excluded those who already had a diagnosis of OP, or were on treatment for OP. This may have led to an under-estimation of the prevalence of OP and OPe. However, in just looking at healthy subjects, we eliminated the possible effect of treatment on increasing BMD, which may have reduced the prevalence of OP and OPe.

5. Conclusions

In a healthy Malaysian urban population aged 45 to 90, 14.0% had OP and 40.9% had OPe. Application of a Caucasian reference range increased the number of subjects with OP to 23.1% and those with OPe to 44.4%, with 21.9% of subjects having changed BMD diagnostic category. Overall, we found that Caucasian T-scores are lower than Asian T-scores in both males and females, and this result was consistently present in the 3 ethnic groups studied. Thus the use of a White (Caucasian) reference range rather than an Asian population specific reference range would lead to a higher rate of diagnosing OP, which can potentially lead to over-treatment.

CRediT author statement

Swan Sim Yeap: Conceptualization, Investigation, Formal analysis, Writing - original draft, Writing - review & editing. Subashini C. Thambiah: Conceptualization, Investigation, Formal analysis, Writing - review & editing, Supervision, Funding acquisition. Intan Nureslyna Samsudin: Conceptualization, Investigation, Writing review & editing, Funding acquisition. Geeta Appannah: Conceptualization, Investigation, Writing - review & editing. Nurunnaim Zainuddin: Investigation. Safarina Mohamad-Ismuddin: Investigation. Nasrin Shahifar: Investigation. Salmiah Md-Said: Conceptualization, Methodology, Investigation, Writing - review & editing. Siti Yazmin Zahari-Sham: Conceptualization, Investigation, Writing - review & editing. Subapriya Suppiah: Formal analysis, Writing - review & editing. Fen Lee Hew: Conceptualization, Investigation, Formal analysis, Writing - review & editing.

Conflicts of interest

The authors declare no competing interests.

Acknowledgments

We would like to thank HK Lim and AK Nurhafizdza for performing the imaging studies. This project was funded by a grant from the Ministry of Education, Malaysia (FRGS/1/2015/SKK03/ UPM/02/1). **ORCID** Swan Sim Yeap: 0000-0002-0474-3667. Subashini C. Thambiah: 0000-0001-6939-9185. Intan Nureslyna Samsudin: 0000-0002-7224-8457. Geeta Appannah: 0000-0003-4636-6529. Nurunnaim Zainuddin: 0000-0002-6076-4112. Safarina Mohamad-Ismuddin: 0000-0002-7040-8939. Nasrin Shahifar: 0000-0002-3298-9170. Salmiah Md-Said: 0000-0001-5865-2499. Siti Yazmin Zahari-Sham: 0000-0002-3765-5700. Subapriya Suppiah: 0000-0002-2495-6408. Fen Lee Hew: 0000-0001-6135-4257.

References

- NIH Consensus Development Panel on osteoporosis prevention, diagnosis, and therapy. Osteoporosis prevention, diagnosis, and therapy. J Am Med Assoc 2001;285. 785–95.
- [2] World Health Organization. Assessment of fracture risk and its application to screening for postmenopausal osteoporosis. Geneva: WHO; 1994. WHO

Technical Report Series, No.843.

- [3] 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.
- [4] Kanis JA, McCloskey EV, Johansson H, Oden A, Melton LJ, Khaltaev N. A reference standard for the description of osteoporosis. Bone 2008;42. 467–75.
- [5] Kanis JA, Adachi JD, Cooper C, Clark P, Cummings SR, Diaz-Curiel M, et al. Standardising the descriptive epidemiology of osteoporosis: recommendations from the epidemiology and quality of life working group of IOF. Osteoporos Int 2013;24. 2763–64.
- [6] Leslie WD. Ethnic differences in bone mass clinical implications. J Clin Endocrinol Metab 2012;97. 4329–40.
- [7] Zengin A, Prentice A, Ward KA. Ethnic differences in bone health. Front Endocrinol 2015;6:1–6.
- [8] Nam HS, Kweon SS, Choi JS, Zmuda JM, Leung PC, Lui LY, et al. Racial/ethnic differences in bone mineral density among older women. J Bone Miner Metabol 2013;31. 190–98.
- [9] Russell-Aulet M, Wang J, Thornton JC, Colt EWD, Pierson RN. Bone mineral density and mass in a cross-sectional study of white and Asian women. J Bone Miner Res 1993;8. 575–82.
- [10] Wu XP, Liao EY, Huang G, Dai RC, Zhang H. A comparison study of the reference curves of bone mineral density at different skeletal sites in native Chinese, Japanese, and American Caucasian women. Calcif Tissue Int 2003;73. 122–32.
- [11] Marquez MA, Melton LJ, Muhs JM, Crowson CS, Tosomeen A, O'Connor MK, et al. Bone density in an immigrant population from Southeast Asia. Osteoporos Int 2001;12. 595–04.
- [12] Lee S, Choi MG, Yu J, Ryu OH, Yoo HJ, Ihm SH, et al. The effects of the Korean reference value on the prevalence of osteoporosis and the prediction of fracture risk. BMC Muscoskel Disord 2015;16:1–13.
- [13] Limpaphayom KK, Taechakraichana N, Jaisamrarn U, Bunyavejchevin S, Chaikittisilpa S, Poshyachinda M, et al. Prevalence of osteopenia and osteoporosis in Thai women. Menopause 2001;8:65–9.
- [14] Ho-Pham LT, Nguyen Ud T, Pham HN, Nguyen ND, Nguyen TV. Reference ranges for bone mineral density and prevalence of osteoporosis in Vietnamese men and women. BMC Muscoskel Disord 2011;10:182.
- [15] Genant HK, Wu CY, Van Kuijk C, Nevitt MC. Vertebral fracture assessment

using a semiquantitative technique. J Bone Miner Res 1993;8. 1137-48.

- [16] Academy of medicine Malaysia/CPGs/Clinical practice guidelines on the management of osteoporosis. second ed, 2015. http://www.acadmed.org.my/ index.cfm?&menuid=67. [Accessed 27 May 2020].
- [17] Shuhart CR, Yeap SS, Anderson PA, Jankowski LG, Lewiecki EM, Morse LR, et al. Executive summary of the 2019 ISCD position development conference on monitoring treatment, DXA cross-calibration and least significant change, spinal cord injury, peri-prosthetic and orthopedic bone health, transgender medicine, and pediatrics. J Clin Densitom 2019;22. 453–71.
- [18] Kanis JA, Odén A, McCloskey EV, Johansson H, Wahl DA, Cooper C. A systematic review of hip fracture incidence and probability of fracture worldwide. Osteoporos Int 2012;23. 2239–56.
- [19] Begum RA, Ali L, Takahashi O, Fukui T, Rahman M. Bone mineral density: reference values and correlates for Bangladeshi women aged 16–65 years. J Orthop Sci 2015;20. 522–28.
- [20] Rathnayake H, Lekamwasam S, Wickramatilake C, Lenora J. Trabecular bone score and bone mineral density reference data for women aged 20–70 years and the effect of local reference data on the prevalence of postmenopausal osteoporosis: a cross-sectional study from Sri Lanka. Arch Osteoporos 2019;14:91.
- [21] Jung KJ, Chung CY, Park MS, Kwon SS, Moon SY, Lee IH, et al. Different reference BMDs affect the prevalence of osteoporosis. J Bone Miner Metabol 2016;34. 347–53.
- [22] Chan CY, Subramaniam S, Mohamed N, Ima-Nirwana S, Muhammad N, Fairus A, et al. Determinants of bone health status in a multi-ethnic population in Klang Valley, Malaysia. Int J Environ Res Publ Health 2020;17:1–16.
- [23] Chan CY, Subramaniam S, Chin KY, Ima-Nirwana S, Muhammad N, Fairus A, et al. Knowledge, beliefs, dietary, and lifestyle practices related to bone health among middle-aged and elderly Chinese in Klang Valley, Malaysia. Int J Environ Res Publ Health 2019;16:4115.
- [24] Lim PS, Ong FB, Adeeb N, Seri SS, Noor-Aini MY, Shamsuddin K, et al. Bone health in urban midlife Malaysian women: risk factors and prevention. Osteoporos Int 2005;16. 2069–79.
- [25] Lee JK, Khir ASM. The incidence of hip fracture in Malaysians above 50 years of age: variation in different ethnic groups. APLAR J Rheumatol 2007;10. 300–5.
- [26] Population Quick Info/population by ethnic group. Selangor, http://pqi.stats. gov.my/result.php?token=1354f1d60aff658d8824da1a93ee5310; 2019. accessed 24 June 2020.