

Study of Assessment of Bone Mineral Density in Women above 45 Years of Age at a Tertiary Care Teaching Centre

Prakrati^{1*}, Deepak Sharma¹, Jaskaran Singh², Geeta², Savita Yadav³, J.K.Khatri⁴

^{1*}Senior Demonstrator, Department of Anatomy, RUHS College of Medical sciences, Jaipur, Rajasthan, India.

²Assistant Professor, ³Senior Demonstrator, ⁴Ex.Senior Professor & Head, Department of Anatomy, S.P. Medical College, Bikaner, Rajasthan, India.

ABSTRACT

Background: Osteoporosis is highly prevalent, with an estimated 30 million women diagnosed to have osteoporosis. A bone mineral density (BMD) test measures how much calcium and other types of minerals are present in a section of bone. The absolute amount of bone as measured by bone mineral density (BMD) testing generally correlates with bone strength and its ability to bear weight. By measuring BMD, it is possible to predict fracture risk in the same manner that measuring blood pressure can help predict the risk of stroke. Hence, we planned this study to evaluate BMD in women above 45 years of age in North-West Rajasthan.

Materials & Methods: The present study was conducted in S.P. Medical College and Associated Group of Hospitals, Bikaner (Rajasthan). A total of 350 women were included in the study, out of which 100 women were controls (30 year of age) and 250 women aged 45 years and above were considered as cases. An informed consent was obtained from all control and cases. BMD was determined by Pronosco X-posure system of hand radiograph to diagnose the osteoporosis and osteopenia in cases. BMD values were measured in terms of T-score and Z-score.

Results: In the patients of age group of 45 to 50 years, 25 had

osteopenia while only had osteoporosis. In the patients of age group of 56 to 60 years, 15 had osteopenia while only 4 patients had osteoporosis. Significant correlation was observed while comparing the diagnosis of the patients with the age groups.

Conclusion: Significant correlation exists while comparing the age group of the patients with the diagnosis for BMD.

Key words: Menopause, Osteoporosis, Osteopenia.

*Correspondence to:

Mrs. Prakrati
Senior Demonstrator
RUHS College of Medical sciences,
Jaipur, Rajasthan, India.

Article History:

Received: 21-08-2016, Revised: 06-09-2016, Accepted: 27-09-2016

Access this article online

Website: www.ijmrp.com	Quick Response code 
DOI: 10.21276/ijmrp.2016.2.5.053	

INTRODUCTION

One of the most rapidly emerging global health problems in women above 45 years of age is osteoporosis. In India, osteoporosis is highly prevalent, with an estimated 30 million women diagnosed to have osteoporosis.¹ Bone is the hardest form of connective tissue. It is composed of approximately 25% water, 30% organic fibrous tissue, and 45% inorganic salts (primarily calcium and phosphates). The skeleton has two major functions: mechanical and metabolic. Most of the mineral content of the body is stored in bone and contributes to its second function that of metabolic homeostasis. It is this second function that is primarily the focus of this study although it is acknowledged that changes in the metabolic processes within the bone are of less concern to patients that the physical, mechanical manifestations of osteoporosis; such as pain and fragility fractures.^{2,3}

A bone mineral density (BMD) test measures how much calcium and other types of minerals are present in a section of bone. The

absolute amount of bone as measured by bone mineral density (BMD) testing generally correlates with bone strength and its ability to bear weight.^{4,5} By measuring BMD, it is possible to predict fracture risk in the same manner that measuring blood pressure can help predict the risk of stroke. From birth to age 25 or 30, the body builds more new bone than it breaks down. By age 30, bones become the strongest they will ever be. This phase of bone development is called peak bone mass. The level of bone mass achieved at the peak is determined largely by genetics, but also by nutrition, exercise and menstrual function.^{6,7}

After about age 30, our body breaks down old bone faster than it builds new bone. This process speeds up dramatically as menopause approaches and for several years after. In the first five to seven years following menopause, bone lose up to 20 percent of bone mass.⁸ In contrast to postmenopausal bone loss, which is associated with excessive osteoclast activity, the bone loss that

accompanies aging is associated with a progressive decline in the supply of osteoblasts in proportion to the demand. This demand is ultimately determined by the frequency with which new multicellular units are created and new cycles of remodeling are initiated.^{9,10} Hence, we planned this study to evaluate bone mineral density (BMD) in women above 45 years of age in North-West Rajasthan.

MATERIALS & METHODS

The present study was conducted in S.P. Medical College and Associated Group of Hospitals, Bikaner (Rajasthan). A total of 350 women were included in the study, out of which 100 women were controls (30 year of age) and 250 women aged 45 years and above were considered as cases. An informed consent was obtained from all control and cases. Bone mineral density by A.P. view radiograph of hand was done in total 350 cases, with an average age of (57.41 ± 9.73).

Inclusion criteria

- The patients who suffered from bone and joint pain and reported to hospital for investigation.
- Aged from 45 years and above.

Exclusion criteria

- History of fracture and previously diagnosed osteoporosis.
- History of diseases such as diabetes, hypertension and cardiovascular diseases.
- With medication which may affect the BMD values.

A standardized Performa was filled which included age, family history, year since menopause, smoking, alcohol and socioeconomic status by a general questionnaire. After that all subjects were referred for bone mineral density test. BMD was determined by Pronosco X-posure system of hand radiograph to diagnose the osteoporosis and osteopenia in cases. BMD values were measured in terms of T-score and Z-score.

T-score: It is the difference between the individual patients bone mineral density and the mean results obtained in young adult population expressed in units of young population standard deviation.

Z-score: It is the difference between the individual patient's results and the mean results obtained in an age matched population expressed in units of the age matched population standard deviation.

According to WHO osteoporosis definition, based on T-score results the categories are:¹¹

Normal bone mass: T-score greater than -1.0

Osteopenia: T-score between -1 and -2.5

Osteoporosis: T-score less than -2.5

Severe osteoporosis: T-score less than -2.5 with at least one osteoporotic fracture

The statistical analysis was done using SPSS (Statistical Package for Social Sciences) Version 15.0 Statistical Analysis Software. Chi-square test was used for the assessment of level of significance.

Table 1: Cross tabulation and relation of age groups with diagnosis categories.

			Diagnosis			Total	Chi-square value	p-value
			Normal	Osteopenia	Osteoporosis			
Age group	45 to 50	Count	39	25	6	70	60.365	0.000
		% of Total	15.6	10.0	2.4	28.0		
	51 to 55	Count	15	21	3	39		
		% of Total	6.0	8.4	1.2	15.6		
	56 to 60	Count	21	15	4	40		
		% of Total	8.4	6.0	1.6	16.0		
	61 to 65	Count	8	13	9	30		
		% of Total	3.2	5.2	3.6	12.0		
	66 to 70	Count	3	20	17	40		
		% of Total	1.2	8.0	6.8	16.0		
	71 and above	Count	2	15	14	31		
		% of Total	.8	6.0	5.6	12.4		
Total		Count	88	109	53	250		
		% of Total	35.2	43.6	21.2	100.0		

RESULTS

Table 1 and Graph 1 shows the cross tabulation and relation of age groups with diagnosis categories. In the patients of age group of 45 to 50 years, 25 had osteopenia while only had osteoporosis. In the patients of age group of 56 to 60 years, 15 had osteopenia while only 4 patients had osteoporosis.

Significant correlation was observed while comparing the diagnosis of the patients with the age groups. Table 2 shows the correlation of age, T score, BMI and BMD of patients groups.

Graph 2 shows scatter plot showing correlation between T score and year since menopause. When comparing the mean age of the patients with mean T score, significant results were obtained.

Significant correlation as obtained while comparing the mean BMD with mean age of the patients. While comparing the mean BMI with the average year since menopause, significant correlation was obtained. Graph 2 shows the scatter plot showing correlation between T score and year since menopause.

Graph 1: Relationship of age groups with diagnosis categories

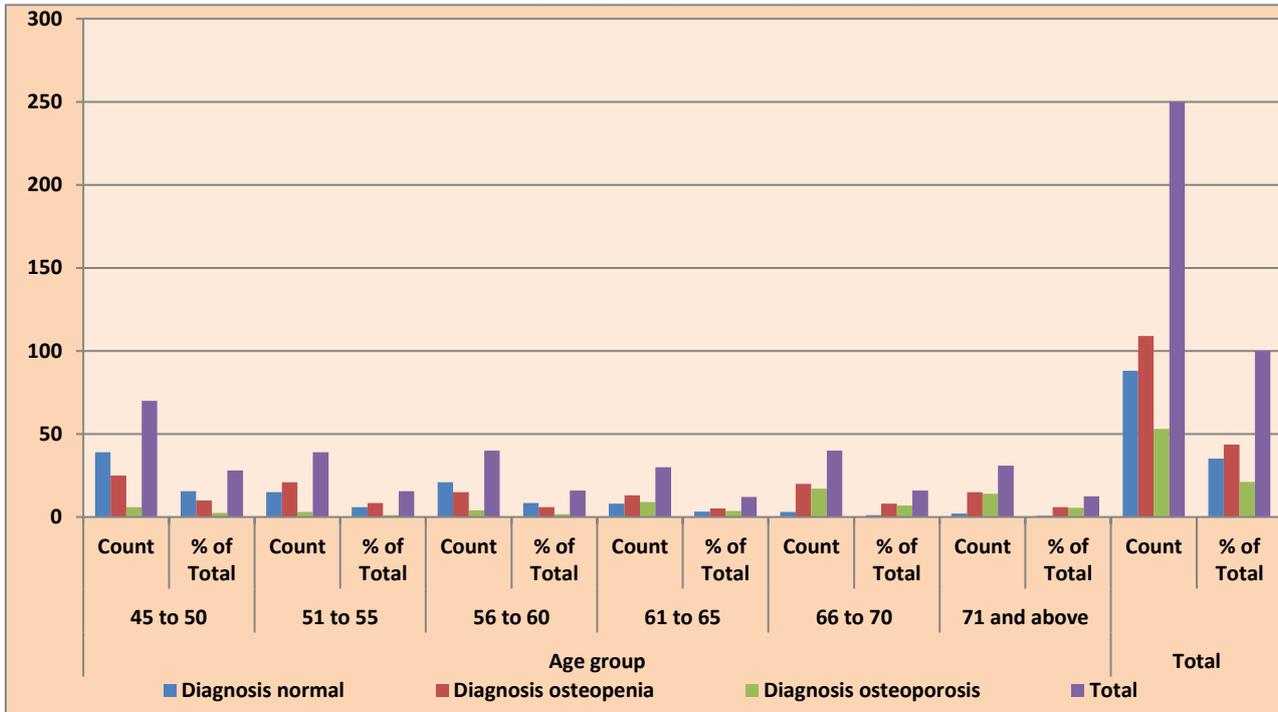
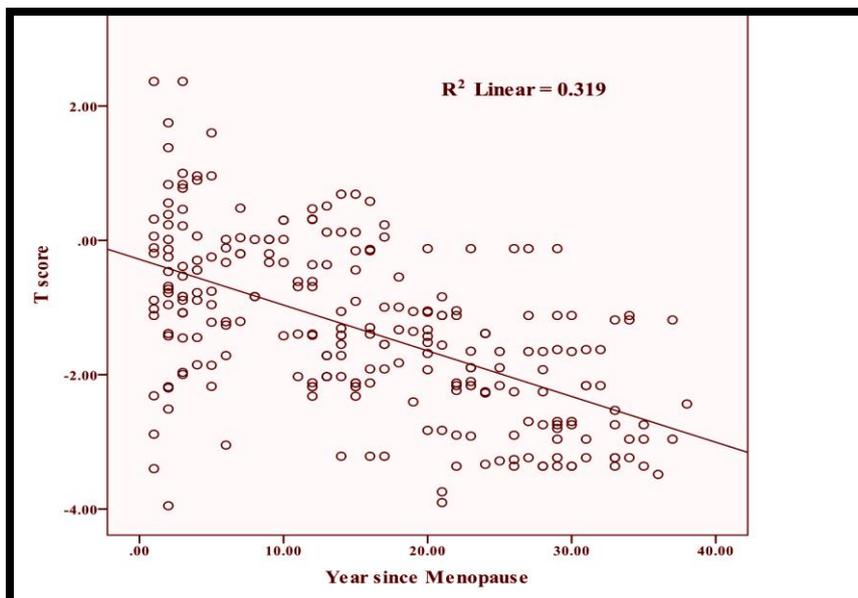


Table 2: Correlation of age, T score, BMI and BMD of patients groups.

		Age	T score	BMD gcm2	BMI	Year since Menopause
Age	Pearson Correlation	1	-.460**	-.462**	.219**	.951**
	Sig. (2-tailed)		.000	.000	.000	.000
T score	Pearson Correlation	-.460**	1	.944**	-.146*	-.565**
	Sig. (2-tailed)	.000		.000	.021	.000
BMD gcm2	Pearson Correlation	-.462**	.944**	1	-.146*	-.557**
	Sig. (2-tailed)	.000	.000		.021	.000
BMI	Pearson Correlation	.219**	-.146*	-.146*	1	.218**
	Sig. (2-tailed)	.000	.021	.021		.001
Year since Menopause	Pearson Correlation	.951**	-.565**	-.557**	.218**	1
	Sig. (2-tailed)	.000	.000	.000	.001	

** . Correlation is significant at the 0.01 level. * . Correlation is significant at the 0.05 level.

Graph 2: Scatter plot showing correlation between T score and year since menopause



DISCUSSION

Osteoporosis affects over 20 million Americans and leads to approximately 1.5 million fractures each year, making it one of the leading public health problems in the United States.¹² The most important risk factor for bone loss in midlife women is the menopause. Women lose about 50% of their trabecular bone and 30% of their cortical bone during the course of their lifetime, about half of which is lost during the first 10 yr after the menopause. Approximately 40% of all postmenopausal women eventually experience fractures. In 2001, the National Osteoporosis Foundation estimated that the annual cost of health care and lost productivity related to osteoporosis was \$17 billion.¹³

In the present study, we observed that 70 patients out of total of 250 (28 percent) belonged to the age group of 45 to 50 years. We also observed that osteopenia was present in 43.6 percent of the patients. A significant correlation was observed between the distribution of patients in various age groups and diagnosis of the patients (p -value < 0.05). Our results were in correlation with the results of Gandhi AB et al who in their study observed that 34 percent of the patients in their study were affected by osteopenia.¹⁴

The influence of obesity on BMD is believed to be mediated by mechanical loading of BMI on bone formation. Nonetheless, lean body mass has been considered as an independent predictor of femoral neck and lumbar spine BMD, whereas the fat component of obesity did not exert any protective effect on bone mass. In a study of obese or overweight African-American women, lean body mass was an independent predictor of total body, hip, spine, and total radius BMD, whereas fat mass in overweight and obese women with metabolic syndrome was a negative predictor of BMD.¹⁵⁻¹⁹

Our results showed a linear increase of osteopenia and osteoporosis with advancing age. In addition to the effects on bone mass, the aging also increases the risk of fractures, regardless of the bone mass, and an increase of 20 years of age means a fourfold risk of fractures. Published studies show that low body weight (less than 58 kg) is associated with an increased risk of osteoporosis and fractures. Weight losses after the age of 50 in women increases the risk of hip fracture, while weight gains decreases this risk.²⁰

Gur et al. reported that BMD decreased with an increase in the 'period of menopause'. These results could be inferred from the relationship between estrogen and BMD. Notelwitz reported that osteoblasts indirectly regulate the resorption activity of osteoclasts under the influence of estrogen. Simm et al. reported that estrogen produces growth factors and contributes to bone formation. Based on these findings, hormone replacement therapy has been used in the treatment of osteoporosis. The results of study of Kim et al showed that participants taking female hormones for more than 1 year had a greater number of teeth present than those not taking female hormones. Again, this result can be inferred from the decreased estrogen secretion occurring in the latter group. According to Hunziker et al., a decrease in estrogen secretion may lead to BMD loss in the mandible in patients with periodontitis. Tezal et al. reported that systemic bone resorption affected periodontal tissues and that a common pathway for destruction exists between systemic and periodontal bone.²¹⁻²⁵

The gynecologist plays an important role in establishing a biological zero in each perimenopausal patient and in controlling the rate of bone loss during postmenopausal period. They also plays an important role in early detection, and in giving advice regarding diet, exercise, calcium supplementation and drugs.¹⁰ Carranza-Lira S et al analyzed how the number of pregnancies and total breast-feeding time influence bone mineral density. 50 healthy women aged 35 to 40 years were studied. Weight, height, number of term pregnancies, time since last pregnancy and breast-feeding time were analyzed. They were divided in three groups as follows: I: nulligravidas; II: with term pregnancies, but no breast-feeding; III: with term pregnancies, with breast-feeding. Bone absorptiometry was done on all, at lumbar column and femur. The WHO criteria were used to define osteopenia and osteoporosis. Comparison among the groups was done with Student's t-test for independent samples, and simple regression analysis was done for number of gestations, total breast-feeding time, time since last birth, and T-score at lumbar column and femur. The women were divided as follows: 15 in group I, 15 in group II, and 20 in group III. The average age was 37.7 +/- 1.5 years. No differences were found among the three groups in analyzed variables, when comparing bone mineral density, T- and Z-scores in L1-L4 average, femoral neck, trochanter, and Ward's triangle. There was no correlation among the number of gestations, total breast-feeding time, time since last birth, or T-scores at lumbar column and femur. The number of pregnancies and breast-feeding time do not affect bone mineral density.²⁶

CONCLUSION

From the above results, the authors conclude that significant correlation exists while comparing the age group of the patients with the diagnosis for BMD. However, future studies are recommended.

REFERENCES

1. Bono CM, Einhorn TA. Overview of osteoporosis: pathophysiology and determinants of bone strength. *Eur Spine J.* Oct 2003; 12 Suppl 2:S90-6.
2. Hayes WC, Piazza SJ, Zysset PK. Biomechanics of fracture risk prediction of the hip and spine by quantitative computed tomography. *RadiolClin North Am.* 1991; 29:1-18.
3. WHO Study Group. Assessment of fracture risk and its application to screening for Postmenopausal osteoporosis. WHO Technical Report Series, World Health Organisation, 1994; Geneva.
4. Genant HK, Engelke K, Fuerst T, Gluer CC, Grampp S, Harris ST, Jergas M, Lang T, Lu Y, Majumdar S, Mathur A, Takada M. Noninvasive assessment of bone mineral and structure: State of the art. *J Bone Miner Res.* 1996; 11:707-730.
5. Mirsky EC, Einhorn T. Bone densitometry in orthopaedic practice: *J Bone Joint Surg (Am).* 1998; 80:1687-1698.
6. Kanis JA, McCloskey EV, De Bakker M, Pande K. Clinical assessment of Bone Mass, Quality and Architecture. *Osteoporos Int.* 1999; Suppl.2:S24-S28.
7. Pande KC. Prevalence of low bone mass in healthy Indian population. *J Ind Med Assoc.* 2002; 1000:598-600.
8. Genant HK. Radiology of Osteoporosis and other Metabolic Bone Diseases. In Favus MJ, editor. *Osteoporosis and Metabolic*

- Bone Diseases. Philadelphia: Lippincot-Raven, Publishers, 1996;152-163.
9. Unni J, Garg R, Pawar R. Bone mineral density in women above 40 years. *J Mid-life Health*. 2010;1:19-22.
 10. Acharya S, Fuchs SC, Donato G, Bastos CA, Spritzer PM. Physical, psychological, and menopause-related symptoms and minor psychiatric disorders in a community-based sample of Brazilian premenopausal, perimenopausal, and postmenopausal women. *Menopause*. 2000; 19:355–60.
 11. Hyldstrup L, Stratakis C, Hill L, Reynolds J, Galliven E, Chrousos G et al. Bone mineral density in women with depression. *N Engl J Med*. 2001; 335:1176–81.
 12. Lee JS, Scholes D, Brunner RL, Robbins J, Reed SD, Newton KM et al. Depressive symptoms, bone loss, and fractures in postmenopausal women. *J Gen Intern Med*. 2001; 23:567–74.
 13. Jagielska G, Primma S, Coyle M, Gourgiotis L, Csako G. Depression and osteoporosis: A research synthesis with meta-analysis. *Horm Metab Res*. 2002; 42:467–82.
 14. Gandhi A, Shukla A. Evaluation of BMD of women above 40 years of age. *J Obstet Gynaecol India* 2005; 55:265-7.
 15. Schneider M, Weller A, Vaisman N, Kreitler S. The relationship of depression, anxiety and stress with low bone mineral density in post-menopausal women. *Arch Osteoporos*. 2002; 7:247–55.
 16. Iketani T, Kiriike N, Nakanishi S, Nakasuji T. Effects of weight gain and resumption of menses on reduced bone density in patients with anorexia nervosa. *Biol Psychiatry*. 1995 Apr 15; 37(8):521-7.
 17. Pearce G, Bradney M, Hendrich E, Delmas P.D, Harding A, Seeman E, et al. (1996) Exercise before puberty may confer residual benefits in bone density in adulthood: studies in active prepubertal and retired gymnasts. *Journal of Bone Mineral Research* 13, 500-507.
 18. Michelson D, Stratakis C, Hill L, Reynolds J, Galliven E, Chrousos G, Gold P. Bone mineral density in women with depression. *N Engl J Med*. 1996 Oct 17; 335(16):1176-81.
 19. Duan Y, Turner CH, Kim BT, Seeman E. Sexual dimorphism in vertebral fragility is more the result of gender differences in age-related bone gain than bone loss. *J Bone Miner Res*. 1997; 16:2267–2275.
 20. Collins FS, Brooks LD. A DNA Polymorphism Discovery Resource for Research on Human Genetic Variation. *Genome Res*. 1998; 8: 1229-1231.
 21. Osei-Hyiaman D, DePetrillo M, Pacher P, Liu J, Radaeva S, Batkai S, et al. Endocannabinoid activation at hepatic CB1 receptors stimulates fatty acid synthesis and contributes to diet-induced obesity. *J Clin Invest*. 1998; 115:1298–1305.
 22. Ghannam NN, Gass M, Hall JE, Lobo R, Maki P, Rebar RW, et al. Executive summary of the stages of reproductive aging workshop + 10: Addressing the unfinished agenda of staging reproductive aging. *J Clin Endocrinol Metab*. 1999; 97:1159–68.
 23. Garnero P, Bener A, Al-Ali HM, Hammoudeh M, Liu LQ, Verjee M. Bone mineral density in midlife women: The study of women's health in Qatar. *Climacteric*. 2000; 18:316–22.
 24. Brembeck P, Winkvist A, Ohlsson C, Lorentzon M, Augustin H. Determinants of microstructural, dimensional and bone mineral changes postpartum in Swedish women. *Br J Nutr*. 2016 Nov; 23:1-9.
 25. Chaki O, Falah A. A measurement-specific quality-of-life satisfaction during premenopause, perimenopause and postmenopause in Arabian Qatari women. *J Midlife Health*. 2000; 5:126–34.
 26. Carranza-Lira S, Fournier A, Mesrine S, Boutron-Ruault MC, Clavel-Chapelon F. Risk factors for onset of menopausal symptoms: Results from a large cohort study. *Maturitas*. 2002; 60:108–21.

Source of Support: Nil.

Conflict of Interest: None Declared.

Copyright: © the author(s) and publisher. IJMRP is an official publication of Ibn Sina Academy of Medieval Medicine & Sciences, registered in 2001 under Indian Trusts Act, 1882. This is an open access article distributed under the terms of the Creative Commons Attribution Non-commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

Cite this article as: Prakrati, Deepak Sharma, Jaskaran Singh, Geeta, Savita Yadav, J.K.Khatri. Study of Assessment of Bone Mineral Density in Women above 45 Years of Age at a Tertiary Care Teaching Centre. *Int J Med Res Prof*. 2016; 2(5):246-50.