

OSTEOPOROSIS, OSTEOPENIA IN MEN OF MARDAN REGION K.P.K PAKISTAN

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ABSTRACT

Objectives: Finding out how common Osteo-penia and Osteo-porosis are among men in the Mardan region of K.P.K. Pakistan is the goal.

Methods: A total of 1726 male participants between the ages of 40 and 70 were randomly selected for this study. sufferers under the age of 40 and above the age of 70, long-term steroid users, sufferers of rheumatoid arthritis, and bedridden patients all met the exclusion criteria.

Results: With a total of 1726 people, 61 (3.5%) had Osteo-porosis, and 858 (49.7%) had Osteo-penia, with a mean age of 52.70(7.71) years and a range of 40-70 years. Men (n=874) between the ages of 50 and 59 showed a significant frequency of Osteo-penia (446; 52%) and Osteo-porosis (30; 49.2%). men (n=[528]) in the 40-49 age group had 250 (16.4%) osteoporotic and 250 (29.1%) osteopenic individuals, whereas men (n=[324]) in the 60-70 age group had 21 (34.4%) Osteo-porosis and 162 (18.9%) Osteo-penia individuals.

Conclusions: Osteo-porosis was discovered to be a widespread occurrence, affecting men between the ages of 50 and 59.

Keywords: Bone mineral density, prevalence, and Osteo-porosis

INTRODUCTION

Osteo-porosis literally means “porous bone.” A condition characterised by bone tissue thinness and loss, ultimately leading to bone density reduction and an increased susceptibility to fractures.¹ There is a widespread myth that women are disproportionately affected by Osteo-porosis. A research article estimates that 6 percent of men over 50 have Osteo-porosis, and that percentage rises with age. An estimated 2 million American males developed Osteo-porosis in 2002, while another 12 million had Osteo-penia.² With an estimated 11.3 million cases in 2020 and 12.9 million cases in 2050, the prevalence of Osteo-porosis

in Pakistan is predicted to rise in the coming years.³ One Australian study found that Osteo-porosis causes fractures in one out of every three men over the age of 60.4 Men’s bone mineral density used to decrease by about 1% annually as they got older.⁵

40 percent of men and postmenopausal women suffer from type I Osteo-porosis. The second kind of Osteo-porosis, often called senile Osteo-porosis, generally affects adults over the age of 75. There were twice as many female casualties as male ones.

Men are more likely to develop Osteo-porosis due to chronic steroid usage, hormonal imbalance, smoking, inactivity, heredity, hypogonadism, and vitamin D insufficiency than women. Because oestrogen and testosterone impact bone tissue through aromatization to oestrogen, mutations of aromatase enzymes or oestrogen receptors have been associated to severe Osteo-porosis in males.^{6,7,8}

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Male Osteo-porosis has a number of proposed mechanisms, one of which is a drop in sex steroid hormone production and/or sensitivity. Bone loss in both sexes is thought to result from hypogonadism. Although males do not go through a process analogous to menopause, at the ages of 50 and 60, both oestrogen and androgen levels, and more especially their bioavailable fractions, decline, leading to complex abnormalities in reproductive physiology.⁹⁻¹² Prolonged use of anabolic steroids, smoking cigarettes, hormonal disruptions, insufficient physical activity, hypogonadism, and a lack of vitamin D are all known to induce Osteo-porosis in men. Aromatization of bone tissue is influenced by both oestrogen and progesterone.¹³⁻¹⁵

Back pain, short stature, and fragility fractures are the most typical early symptoms of Osteo-porosis in men. There is substantial debate over what constitutes an acceptable bone mineral density cutoff in men. While more information is gathered, it is acceptable to use gender-specific criteria (a T score that is 2.5 standard deviations below the young male advertence mean) to diagnose Osteo-porosis in men.^{16,17}

To diagnose Osteo-porosis, bone mineral density may be assessed by ultrasonography (calcaneal), DXA scan (dual X-ray absorption), SPA (single photon absorption), and QCT (quantitative computerised tomography). Strong correlations between the values of bone mineral density determined by a DXA scan and ultrasonography have been found in prior studies.^{18,19} Compared to DXA, ultrasound measurements of bone mineral density are less intrusive, more inexpensive, and more accessible. Ultrasound is used to evaluate the heel bone mineral density, which is then converted to T-scores by comparing the results to those of the general adult population. The World Health Organisation defines Osteo-penia as a T-score of 2.5 or above.²⁰⁻²¹

Our study seeks to ascertain the prevalence of Osteo-penia and Osteo-porosis in men in the Mardan region of K.P.K. Pakistan.

MATERIAL & METHODS

Ethical Review Committee blessing allowed us to begin the project. From May through October of 2014, researchers at the Mardan Medical Complex Hospital gathered data in a prospective cross-sectional study. All patients who came via the hospital's outpatient clinic (OPD) throughout the study's enrolment period were considered participants. After participants were briefed

on the study's goals, informed consent was collected from them. Non-probabilistic convenience sampling was used to collect data. Males aged 40-70 made up the bulk of the study's 1726 participants. People who were chronic steroid users, had rheumatoid arthritis, or were bedridden were also not allowed to participate. Participants' ages were used to create three groups: those aged 40-49, 50-59, and 60-70. Participants were briefed beforehand and given the opportunity to consent to the experiment. Bone mineral density was assessed using the SONOSOT 3000 (Software Version: 3.03.06) bone densitometer by Calcaneal, Inc. After bone mineral density measurements were automatically converted, the T-score suggested by the World Health Organisation was used to assess bone fragility. SPSS (20.0) was used to generate descriptive statistics.

RESULTS

Mean age was 52.70(7.71) years, and the age range of the 1726 participants was 40–70 years old. Of these, 60 (03.05% were osteoporotic, and [858] (48%) were osteopenic. (Figure 1 of Table 1)

Out of a total of [528] people, 10 (16.4%) were diagnosed with Osteo-porosis between the ages of 40 and 49; out of a total of 874, 30 (49.5%) were diagnosed between the ages of 50 and 59; and out of a total of 324, 21 (34.4%) were diagnosed between the ages of 60 and 70. Osteo-penia was found in 162 (18.9%) of those aged 60-70, 446 (52.1%) of those aged 50-59, and 250 (29.1%) of those aged 40-49. (Table 2)

DISCUSSION

Metabolic Osteo-porosis reduces bone density and microarchitectural architecture, causing fragility fractures.²² Osteo-porosis will certainly become the most common age-related bone disease as the global elderly population grows. Hip fractures are expected to increase sixfold by 2050 due to this age change.²³

Meier et al.²⁴, Jones et al.²⁵, and Hannan et al. discover a statistically significant relationship between age and Osteo-porosis prevalence in multivariate and univariate analysis.²⁶ Age significantly affected bone mineral density, and men's bone mass dynamically reduced with age, especially in older men.

Our study found 30.9% of men aged 50–59 had Osteo-porosis, and 19.8% of those aged 60–70 had. A study in Lahore, Pakistan, found 20.6% of males over 45 had Osteo-porosis and 10.7% of men under

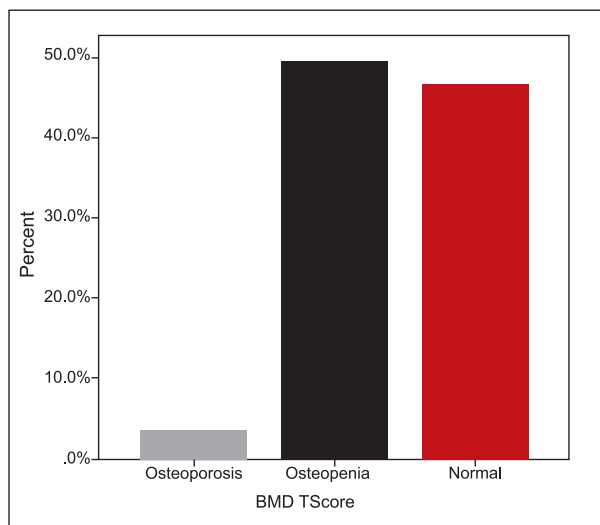


Figure 1: Bone mineral density of total population

Table 1: BMD of total population n=1726

BMD T-Score	Male
Osteoporosis	61-(03.5%)
Osteopenia	858 -(48.7%)
Normal	806 -(45.8%)
Total	1727- (39.2%)

Table 2: T-score, sex, and age ranges for BMD

Age range	BMD T-Score	Male
40-49 years	Osteoporosis	11(15.4%)
	Osteopenia	251(28.1%)
	Normal	267(23.2%)
	Total	528
50-59 years	Osteoporosis	31(48.2%)
	Osteopenia	445(51%)
	Normal	397(48.3%)
	Total	875
60-70 years	Osteoporosis	21(36%)
	Osteopenia	163(19%)
	Normal	141(18%)
	Total	324

45.27 In Saudi Arabia, 23% of men over 50 have Osteoporosis, according to El-Desouki²⁸. According to Garg N et al.²⁹, 60 (66.8%) of 170 males aged 50 and older in Muzaffarnagar district had Osteopenia and 10 (11.1%) had Osteoporosis. Older Brazilian men had 6.4%-16.1% Osteoporosis and 33.5%-57.4% Osteopenia.³⁰ Westgaard et al.³¹ found 17.7% of Danish men over 50 had Osteoporosis. A British

study found Osteoporosis in 6% of males over 50.32 Canada had 4.8% femoral neck and 2.9% lumbar spine Osteoporosis.³³

Our 40-49-year-old subjects also had osteopenic and osteoporotic bone density. This suggests that bone mineral density loss in men begins sooner than previously assumed, emphasising the need for early prevention. Calcium intake and absorption are worse in the elderly. Circumlocutory oestrogens help women absorb calcium from their guts and kidneys.

Calcium and vitamin D deficiency affects many Pakistanis. A Karachi hospital reported that 92% of its “Out Door Patients” had vitamin D deficiency, with a 5:1 female-to-male ratio.³³ Another study indicated that the average Pakistani adult ingested 400–600 milligrammes of calcium per day, significantly less than the recommended 1,000–1,200.34 Pakistanis ingested much less calcium than African-Caribbean, Pakistani, and European adults in another study.³⁵

Early diagnosis and treatment of Osteoporosis reduces fracture risk. Due to the difficulty of restoring bone mass after Osteoporosis, prevention is as important as treatment. Giving up smoking and drinking, exercising more, and eating a balanced diet rich in calcium and vitamin D can improve health. Calcium and vitamin D-rich foods strengthen bones.

There were limitations to our investigation. It was a hospital study, which may have overestimated Osteoporosis prevalence. Bone mineral density was measured using quantitative calcaneus ultrasonography. Quantitative ultrasonography of the calcaneus produces bone mineral density indices that match DXA scans, however DXA is the gold standard and should be used whenever possible. The high cost of DXA and our lack of funds prevented us from proceeding. However, quantitative ultrasonography is reliable and has been used in Osteoporosis studies. Large-scale DXA scan population studies are needed to estimate local Osteoporosis prevalence.

CONCLUSION

It's possible that the low BMD is an ethnic variation. In the Mardan region of K.P.K. Pakistan, males in their 50s and 60s are disproportionately affected by Osteopenia and Osteoporosis. Adequate calcium intake and physical activity are particularly important in reducing the risk of fracture in this population.

REFERENCES

1. Alldredge BK, Koda-Kimble MII. Applied Therapeutics: the clinical use of drugs. Philadelphia: Wolters Kluwer Health/Lippincott Williams and Wilkins. 2009; pg. 101- 3.
2. Orwoll ES. Osteoporosis in men. *Endocrinol Metab Clin North Am.* 1998; 27:349-367.
3. Mithal A, Dhingra V, Lau EB. The Asian audit: epidemiology, costs and burden of Osteoporosis in Asia. China: International Osteoporosis Foundation(IOF) Publication; 2009.
4. Osteoporosis and men, leaflet from Osteoporosis Australia. *Med J* 1997; 167:51-5.
5. Hannan MT, Felson DT, Dawson-Hughes B, Tuck KL. Risk fractures for longitudinal bone loss in elderly men and women: The Framingham Osteoporosis study. *J Bone Miner Res* 2000; 15:710-720.
6. Ebeling PR. Osteoporosis in men: new insights into aetiology, pathogenesis, prevention and management. *Drugs Aging* 1998; 13:421-34.
7. Smith EP, Boyd J, Frank GR. Estrogen resistance caused by a mutation in the estrogen-receptor gene in a man. *N Engl J Med* 1994; 331:1056-61.
8. Morishima A, Grumbach MM, Simpson ER, Fisher C, Qin K. Aromatase deficiency in male and female siblings caused by a novel mutation and the physiological role of estrogens. *J Clin Endocrinol Metab* 1995; 80:3689-98.
9. Riggs BL, Khosla S, Melton LJ III. Sex steroids and the construction and conservation of the adult skeleton. *Endocr Rev* 2002; 23(3):279-302.
10. Harman SM, Metter JF, Tobin JD. Longitudinal effects of aging on serum total and free testosterone levels in healthy men. *J Clin Endocrinol Metab* 2001; 86(2):724-31.
11. Khosla S, Melton LJ III, Atkinson EJ. Relationship of serum sex steroid levels to longitudinal changes in bone density in young versus elderly men. *J Clin Endocrinol Metab* 2001; 86(8):3555-61.
12. Gennari L, Merlotti D, Martini G. Longitudinal association between sex hormone levels, bone loss, and bone turnover in elderly men. *J Clin Endocrinol Metab* 2003; 88(11):5327-33.
13. Lamberts SWJ, Van den Beld AW, Van der Lely A. The endocrinology of aging. *Science* 1997; 278:419-24.
14. Gray A, Feldman HA, McKinlay JB. Age, disease and changing sex hormone levels in middle-aged men: results of the Massachusetts male aging study. *J Clin Endocrinol Metab* 1991; 73:1016-25.
15. Amin S, Zhang Y, Sawin CT. Association of hypogonadism and estradiol levels with bone mineral density in elderly men from the Framingham study. *Ann Intern Med* 2000; 133(12):951-63.
16. Orwoll E. Assessing bone density in men. *J Bone Miner Res* 2000; 15:1867-70.
17. Lewiecki EM, Watts NB, McClung MR. International Society for Clinical Densitometry. Official positions of the international society for clinical densitometry. *J Clin Endocrinol Metab* 2004; 89:3651-5.
18. Falcini F, Bindi G, Ermini M, Galluzzi F. Comparison of quantitative calcaneal ultrasound and dual energy X-ray absorptiometry in the evaluation of osteoporotic risk in children with chronic rheumatic diseases. *Calcif Tissue Int* 2000; 67: 19-23.
19. Massie A, Reid DM, Porter RW. Screening for Osteoporosis: comparison between dual energy X-ray absorptiometry and broadband ultrasound attenuation in 1000 perimenopausal women. *Osteoporos Int* 1993; 3:107-10.
20. World Health Organization. WHO assessment of fracture risk and its application to screening for postmenopausal Osteoporosis: report of a WHO study group. Geneva: World Health organization, 1994.
21. Kanis JA, McCloskey EV, Johansson H A. A reference standard for the description of Osteoporosis, *Bone* 2008; 42:3:467-475.
22. Consensus Development Conference V, 1993. Diagnosis, Prophylaxis and treatment of Osteoporosis. *Am J Med* 1994; 90: 646-50.
23. Cooper C, Campian G, Melton IJ. Hip fractures in the elderly: a worldwide projection. *Osteoporosis Int* 1992; 2: 285-9.
24. Meier DE, Orwoll ES, Jones JM. Marked disparity between trabecular and cortical bone loss with age in healthy men. *Ann Intern Med* 1984; 101:605-12.
25. Jones G, Nguyen T, Sambrook P. Progressive loss of bone in the femoral neck in elderly people: longitudinal findings from the Dubbo Osteoporosis epidemiology study. *BMJ* 1994; 309:691-5.
26. Hannan MT, Felson DT, Anderson JJ. Bone mineral density in elderly men and women: results from the Framingham Osteoporosis Study. *J Bone Miner Res* 1992; 7:547-53.
27. Nagi D, Butt Z, Farooq F. Frequency of Osteoporosis in ambulatory setting in Lahore using calcaneal ultrasound. *J Pak Med Assoc* 2013; 63:8:965-968.
28. El-Desouki MI, Sulimani RA. High prevalence of Osteoporosis in Saudi men. *Saudi Med J* 2007; 28(5):774-777.
28. Garg N, Kumar A, Goel P. Prevalence of Osteoporosis in rural population of Muzaffarnagar district. *JACM* 2012; 13(3):185-8.

30. Rodrigues Camargo MB, Cendorogolo MS, Ramos LR. Bone mineral density and Osteoporosis among a preminiantly Caucasian elderly population in the city of San Paulo, Brazil. *Osteoporosis International* 2005; 16:1451-1460.
31. Westgaard, Rejumark L, Mosekilde L. Osteoporosis is markedely underdiagnosed: a nationwide study from Denmark. *Osteoporosis Int* 2005; 16:134-141.
32. Hoh G, Khan KT, Reid DM. Prevalence of osteoporotic bone mineral density of the hip in Britain differs substantially from U.S. *Br J Radio* 2002; 75:736-742.
33. Tenhouse A, Joseph L, Krieger N. Estimation of the prevalence of low bone mineral density in Canadian women and men. *Osteoporos Int* 2000; 11:897-904.
34. Zuberi LM, Habib A, Haque N, Jabbar A. Vitamin D deficiency in ambulatory patients. *J Pak Med Assos* 2008; 58: 482-4.
35. Iqbal R, Khan AH. Possible causes of vitamin D deficiency (VDD) in Pakistani population residing in Pakistan. *J Pak Med Assos* 2010; 60: 1-2.
36. Vyas A, Greenhalgh A, Cade J, Sanghera B, Riste L. Nutrient intakes of an adult Pakistani, European and African-Caribbean community in inner city Britain. *J Hum Nutr Diet* 2003; 16: 327-37.