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Review Article

## A Review on Study on Osteoporosis and Impact of Drugs for its Management

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### ABSTRACT

Osteoporosis is characterized by low bone mineral density (BMD) and loss of the structural and biomechanical properties that are required to maintain bone homeostasis. Physicians are knowledgeable about the association of osteoporosis with aging, postmenopausal status, and secondary causes, including chronic illnesses or lifestyle issues that promote osteoporosis. Building strong and healthy bones requires an adequate dietary intake of calcium beginning in childhood and adolescence for both sexes. Most importantly, however, a high dietary calcium intake or taking calcium supplements alone is not sufficient in treating osteoporosis and should not be viewed as an alternative to or substituted for more potent prescription medications for osteoporosis.

**Keywords:** BMD, Osteoporosis, calcium supplements, skeletal disease.

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### 1. INTRODUCTION:

Osteoporosis afflicts 10 million Americans, and another 30 million Americans have low bone mass that puts them at risk for development of osteoporosis<sup>1</sup>. The overall risk for a 50-year-old Caucasian or Asian woman to sustain an osteoporotic fracture during her lifetime is 50%. While men, African Americans, and Hispanics have lower risks, the health impact is still substantial. One-third of all hip fractures occur in men, and with our aging populace hip fracture incidence in men will eventually approach that of women<sup>1</sup>. A femoral neck or trochanteric fracture is associated with mortality of approximately 20%<sup>1</sup>

### 2. DIAGNOSIS:

Osteoporosis diagnostics can be complicated. The World Health Organization (WHO) defined osteoporosis as a systemic skeletal disease characterized by low bone mass with microarchitectural deterioration of bone tissue, thus increasing bone fragility and susceptibility to fracture<sup>1</sup>. For screening purposes, osteoporosis was defined by the WHO as a bone mineral density (BMD) at any site equal to or greater than 2.5 standard deviations below the fracture resistant mean peak bone mass of young adulthood.

### 3. SIGN AND SYMPTOMS:

Bone loss that leads to osteoporosis develops slowly. There are often no symptoms or outward signs, and a person may

not know they have it until they experience a fracture<sup>3</sup> after a minor incident, such as a fall, or even a cough or sneeze.

- Commonly affected areas are the hip, a wrist, or spinal vertebrae<sup>4</sup>.
- Breaks in the spine can lead to changes in posture, a stoop, and curvature of the spine.

### 4. CAUSES OF BONE FRAGILE:

Your skeleton is an active vital organ. It keeps you healthy through a constant process of repair, renewal, and mineral release. This process is called remodeling. The bone remodeling cycle consists of two distinct stages:

- (1) bone resorption<sup>5</sup> (breakdown and removal) and
- (2) bone formation (generation of new bone).

During resorption, cells on the bone's surface called osteoclasts dissolve bone tissue, releasing it into the bloodstream and leaving behind tiny pits, or cavities. Then, during formation, cells called osteoblasts fill these cavities with new bone tissue. In normal bone, resorption and formation are in lock step, with one matching the other.<sup>6</sup>

### 5. MEDICATION TO PREVENT FRAGILITY FRACTURES:

Here are many medications available to treat osteoporosis and reduce the risk of fracture. They fall into two basic

categories: antiresorptives and anabolics. Antiresorptive drugs include bisphosphonates (alendronate, ibandronate, risedronate, zoledronic acid), denosumab, calcitonin, estrogen/estrogen-progestin, an estrogen agonist/antagonist (raloxifene), and a tissue specific estrogen complex (estrogen/bazedoxifene).<sup>7-15</sup>

## 6. MEDICATION FOR TREATMENT OF OSTEOPOROSIS:

Bisphosphonates are the most common medications prescribed for osteoporosis treatment. These include:

- Alendronate (Fosamax)<sup>9</sup>
- Risedronate (Actonel)<sup>10</sup>
- Ibandronate (Boniva)<sup>12</sup>
- Zoledronic acid (Reclast)<sup>14</sup>

Hormones, such as estrogen, can play a role in osteoporosis prevention and treatment. However, there has been some concern about potential side effects tied to the use of hormone therapy.<sup>16</sup> Current recommendations say to use the lowest dose of hormones for the shortest period of time.

## 7. DRUG THERAPY:

Drugs that can help prevent and treat osteoporosis include:

- Bisphosphonates: These are antiresorptive drugs that slow bone loss and reduce fracture risk.<sup>9</sup>
- Estrogen agonists or antagonists, also known as selective estrogen-receptor modulators, SERMS), for example, raloxifene (Evista): These can reduce the risk of spine fractures in women after menopause.<sup>17</sup>
- Calcitonin (Calcimar, Miacalcin): This helps prevent spinal fracture in postmenopausal women, and it can help manage pain if a fracture occurs.<sup>11</sup>
- Parathyroid hormone, for example, teriparatide (Forteo): This is approved for people with a high risk of fracture, as it stimulates bone formation.
- Monoclonal antibodies<sup>18</sup> (denosumab, romosozumab): These are immune therapies given to some postmenopausal women with osteoporosis. Romosuzumab carries a black box warning due to possible adverse effects.

### 7.1 Specific Drug Treatments Include:

- Bisphosphonates<sup>9</sup>
- Denosumab
- Parathyroid Hormone
- Hormone Therapy
- SERMs<sup>19</sup> (Selective Estrogen Receptor Modulators)

## 8. CONSEQUENCES OF OSTEOPOROSIS:

Osteoporotic bone fractures are responsible for considerable pain, decreased quality of life, lost workdays, and disability. Up to 30% of patients suffering a hip fracture will require long-term nursing-home care. Elderly patients can develop pneumonia and blood clots in the leg veins that can travel to the lungs (pulmonary embolism) due to prolonged bed rest after the hip fracture. Osteoporosis has even been linked with an increased risk of death.<sup>20</sup> Some 20% of women with a hip fracture will die in the subsequent year as an indirect result of the fracture. In addition, once a person has

experienced a spine fracture due to osteoporosis, he or she is at very high risk of suffering another such fracture in the near future (next few years).<sup>21</sup> About 20% of postmenopausal women who experience a vertebral fracture will suffer a new vertebral fracture of bone in the following year.

## 9. CONCLUSION:

The goal of treatment of osteoporosis is the prevention of bone fractures by reducing bone loss or, preferably, by increasing bone density and strength. Although early detection and timely treatment of osteoporosis can substantially decrease the risk of future fractures, none of the available treatments for osteoporosis are complete cures. In other words, it is difficult to completely rebuild bone that has been weakened by osteoporosis. Therefore, prevention of osteoporosis is as important as treatment.

## 10. REFERENCES:

1. Woolf AD, Pfleger B. Burden of major musculoskeletal conditions. *Bull World Health Organ.* 2003;81(9):646-656.
2. Jones, Daniel (2003) [1917], Peter Roach; James Hartmann; Jane Setter (eds.), *English Pronouncing Dictionary*, Cambridge: Cambridge University Press, ISBN 978-3-12-539683-8
3. "Handout on Health: Osteoporosis". August 2014. Archived from the original on 18 May 2015. Retrieved 16 May 2015.
4. WHO Scientific Group on the Prevention and Management of Osteoporosis (2000 : Geneva, Switzerland) (2003). *Prevention and management of osteoporosis : report of a WHO scientific group* (PDF). pp. 7, 31. ISBN 978-9241209212. Archived (PDF) from the original on 16 July 2007.
5. Wells GA, Cranney A, Peterson J, Boucher M, Shea B, Robinson V, Coyle D, Tugwell P (January 2008). "Alendronate for the primary and secondary prevention of osteoporotic fractures in postmenopausal women". *The Cochrane Database of Systematic Reviews* (1): CD001155.
6. Wells G, Cranney A, Peterson J, Boucher M, Shea B, Robinson V, Coyle D, Tugwell P (January 2008). "Risedronate for the primary and secondary prevention of osteoporotic fractures in postmenopausal women". *The Cochrane Database of Systematic Reviews* (1): CD004523.
7. "Chronic rheumatic conditions". World Health Organization. Archived from the original on 27 April 2015. Retrieved 18 May 2015.
8. Golob AL, Laya MB (May 2015). "Osteoporosis: screening, prevention, and management". *The Medical Clinics of North America.* 99 (3): 587-606. .
9. Wells GA, Cranney A, Peterson J, Boucher M, Shea B, Robinson V, Coyle D, Tugwell P (January 2008). "Etidronate for the primary and secondary prevention of osteoporotic fractures in postmenopausal women". *The Cochrane Database of Systematic Reviews* (1): CD003376.
10. Nelson, H. D; Haney, E. M; Chou, R; Dana, T; Fu, R; Bougatsos, C (2010). "Screening for Osteoporosis: Systematic Review to Update the 2002 U.S. Preventive Services Task Force Recommendation [Internet]". Agency for Healthcare Research and Quality. PMID 20722176.
11. Wade SW, Strader C, Fitzpatrick LA, Anthony MS, O'Malley CD (2014). "Estimating prevalence of osteoporosis: examples from industrialized countries". *Archives of Osteoporosis.* 9 (1): 182.
12. Handa R, Ali Kalla A, Maalouf G (August 2008). "Osteoporosis in developing countries". *Best Practice & Research. Clinical Rheumatology.* 22 (4): 693-708.
13. Svedbom A, Hernlund E, Ivergård M, Compston J, Cooper C, Stenmark J, McCloskey EV, Jönsson B, Kanis JA (2013). "Osteoporosis in the European Union: a compendium of country-specific reports". *Archives of Osteoporosis.* 8 (1-2): 137.
14. Willson T, Nelson SD, Newbold J, Nelson RE, LaFleur J (2015). "The clinical epidemiology of male osteoporosis: a review of the recent literature". *Clinical Epidemiology.* 7: 65-76.

15. King TL, Brucker MC (2011). Pharmacology for women's health. Sudbury, Mass.: Jones and Bartlett Publishers. p. 1004. ISBN 9780763753290. Archived from the original on 8 September 2017.
16. Kasper, Dennis L.; Fauci, Anthony S.; Hauser, Stephen L.; Longo, Dan L.; Larry Jameson, J.; Loscalzo, Joseph (6 February 2018). Harrison's principles of internal medicine. Jameson, J. Larry,, Kasper, Dennis L.,, Fauci, Anthony S., 1940-, Hauser, Stephen L.,, Longo, Dan L. (Dan Louis), 1949-, Loscalzo, Joseph (Twentieth ed.)
17. Kim DH, Vaccaro AR (2006). "Osteoporotic compression fractures of the spine; current options and considerations for treatment". The Spine Journal. 6 (5): 479–87.
18. Susan Ott."Fracture Risk Calculator". Archived from the original on 14 October 2009. Retrieved 3 November 2009.
19. WHO (1994). "Assessment of fracture risk and its application to screening for postmenopausal osteoporosis. Report of a WHO Study Group". World Health Organization Technical Report Series. 843: 1–129. PMID 7941614.
20. Ganz DA, Bao Y, Shekelle PG, Rubenstein LZ (2007). "Will my patient fall?". JAMA. 297(1): 7786.
21. Waugh EJ, Lam MA, Hawker GA, McGowan J, Papaioannou A, Cheung AM, Hodsman AB, Leslie WD, Siminoski K, Jamal SA (January 2009). "Risk factors for low bone mass in healthy 40–60 year old women: a systematic review of the literature". Osteoporosis International. 20 (1): 1–21.

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