Secondary Causes of Osteoporosis

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ABSTRACT

Osteoporosis (OP) is characterized by low bone mass, microarchitectural disruption and skeletal fragility, resulting in decreased bone strength and an increased risk of fracture. In fact, fractures are the clinically relevant signs of OP, an otherwise silent disease. In inflammatory rheumatic diseases such as rheumatoid arthritis (RA), psoriatic arthritis (PsA) and ankylosing spondylitis (AS) the systemic burden caused by inflammation, glucocorticoid treatment, immobilization and reduced physical activity due to painful joints and muscle weakness are associated with decreased bone mass and deterioration of bone quality leading to increased risk of falls and fragility fractures.

Keywords
Microarchitectural disruption, Deterioration, Fragility fractures

Introduction

Osteoporosis (OP) is characterized by low bone mass, microarchitectural disruption and skeletal fragility, resulting in decreased bone strength and an increased risk of fracture. In fact, fractures are the clinically relevant signs of OP, an otherwise silent disease.

Despite being the most relevant determinant of bone strength, Bone Mineral Density (BMD) is not the only factor contributing to the increased risk of fractures with aging. Other important factors include age itself, bone turnover rate, bone geometry and microarchitecture.

OP is estimated to affect 200 million women worldwide, 75 million of which in Europe, USA and Japan [1,2]. In the year 2000, there were an estimated 9.0 million osteoporotic fractures worldwide, of which 1.6 million were at the hip, 1.7 million at the forearm and 1.4 million were clinical vertebral fractures. The greatest number of osteoporotic fractures occurred in Europe (34.8%) [3]. According to a large national health survey on rheumatic diseases (Epireuma.pt), the estimated prevalence of OP in the adult Portuguese population is 10.2% (17.0% in women and 2.6% in men) [4].

Based on the WHO diagnostic criteria (T-score less than or equal to -2.5 SD) approximately 22 million women and 5.5 million men aged 50 to 84 are estimated to have OP in the European Union (2010 figures). Due to changes in population demography the number of men and women with OP in the EU is estimated to rise to 33.9 million in 2025, corresponding to a 23% increase [5]. The remaining lifetime risk of fracture of the spine, hip, distal forearm and proximal humerus for women at 50 years of age is 46.4% [6]. Secondary causes of OP might be more frequent than once thought, highlighting the need for a careful diagnostic workup. According to the 2018 update of the Portuguese recommendations for the prevention, diagnosis and management of OP[7], secondary OP (SOP) should be suspected in the presence of: a) conditions known to induce osteoporosis (Table 1); b) in the presence of fragility fractures occurring before the age of 70 for men or before menopause for women; c) low Z scores (≤ -2.0) in dual-energy X-Ray absorptiometry (DXA) studies.

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In this review article, some of the causes of SOP will be discussed in more detail, as well as their screening, diagnosis and treatment.

In an observational study [8] taking place in an outpatient rheumatology department, causes of osteoporosis in 81 osteoporotic men were evaluated and secondary osteoporosis was diagnosed in 63 (78%). In men with vertebral crush fractures, some investigators have suggested that 55% have a secondary cause and 20% of these cases are due to hypogonadism. In a series of 214 women with vertebral crush fractures [9], 30.4% were found to have an underlying cause of osteoporosis or early menopause (36.4%) before the age of 45 years. Other estimates of osteoporosis in women suggested that approximately 20% of women who appear to have postmenopausal osteoporosis have an identifiable secondary cause, whereas the incidence of men with a secondary cause has been estimated to be as high as 64%. A cross-sectional study [10] with 173 postmenopausal women with primary osteoporosis showed that after lab tests including complete blood count, renal and hepatic function, chemistry profile, 24-h urinary calcium, 25(OH)vitamin D and PTH, 55 (32%) of them had undiagnosed disturbances of bone and mineral metabolism. The most frequent diagnosis was vitamin D deficiency (20.2%), hypercalciuria (9.8%), malabsorption (8.1%), hyperparathyroidism (6.9%) and exogenous hyperthyroidism (2.3%).

The treatment, prognosis and monitoring of osteomalacia is different from osteoporosis, thus it is an important differential diagnosis to take into account during the evaluation of SOP. A significant proportion of women with fragility fractures have osteomalacia and/or osteoporosis. In a retrospective study [11], secondary causes for reduced BMD were evaluated in 196 postmenopausal and 41 premenopausal women. Sixteen percent of these patients had 25-hydroxyvitamin D levels lower than 15 ng/ml. By using the World Health Organization (WHO) definition of osteopenia based on T score value (–1.0 to –2.5), 11% of osteopenic patients had 25-hydroxyvitamin D levels lower than 15 ng/ml. On the other hand, osteomalacia is present in 4% to 47% of men with femoral fractures, with most studies [12,13] reporting a rate of close to 20%. As shown in Table 2, several clinical conditions are associated with increased fracture risk not only by directly decreasing bone density or deteriorating bone quality, but also by increasing fall risk. In addition, pharmacological
treatment of some diseases might lead to decreased bone quality and increased fracture risk due to drug induced SOP.

**Drug-Induced Osteoporosis**

Several drugs have been associated with increased fracture risk (Table 3) and glucocorticoids are the most common cause of drug-induced osteoporosis [14]. Bone loss is due to suppression of osteoblast function, inhibition of intestinal calcium absorption leading to secondary hyperparathyroidism and increased osteoclast-mediated bone resorption. Glucocorticoid excess results in diffuse bone loss and may affect trabecular bone more than cortical bone.

Bone loss is also promoted by direct stimulation of renal excretion of calcium by glucocorticoids. Bone Mineral Density is reduced in 40% to 60% of patients with an endogenous glucocorticoid excess, and pathologic fractures have been observed in 16% to 67%. The risk of hip fractures is doubled in glucocorticoid-treated patients [15].

Studies evaluating short-term exposures to glucocorticoid therapy have indicated that glucocorticoid-induced bone loss appears greater in the first 6 to 12 months of therapy [16].

Inhaled glucocorticoid therapy was associated with a dose-related decrease in BMD at the total hip and trochanter (0.00044 g/cm² per puff per year of treatment) [17]. This finding, along with a retrospective study [10] revealing an association between doses of 2.5 mg daily prednisone and bone loss, suggest a low threshold at which glucocorticoids cause skeletal harm. Fracture risk increases with dose and duration of glucocorticoid use [18].

Serum and urine biochemical indices in patients with glucocorticoid-induced osteopenia are generally normal, but urinary markers of bone resorption may be increased. Serum PTH levels may be normal or mildly elevated (secondary hyperparathyroidism) and serum alkaline phosphatase activity and osteocalcin levels decline steadily after the initiation of glucocorticoid therapy, reflecting inhibition of osteoblast activity. Urinary calcium excretion may be increased during the first several months to years of glucocorticoid use because of the direct calciuric effect of glucocorticoids on the kidney.

The first principle in the treatment of patients with glucocorticoid-induced osteoporosis is to use the lowest effective dose of glucocorticoid. General health measures that are applicable to patients with osteoporosis should be encouraged, such as weight-bearing exercise and good nutritional status.

According to the 2017 update of the American College of Rheumatology recommendations for the prevention and treatment of glucocorticoid-induced OP, patient-taking glucocorticoids should optimize calcium (1,000-1,200 mg/day) and vitamin D intake (600–800 IU/day) [18].

For post-menopausal women and men >40 years starting glucocorticoid therapy with an anticipated duration of more than 3 months, risk stratification using FRAX and the dose of glucocorticoids are key for an adequate therapeutic decision. For low risk patients, no further treatment is recommended and patients should be monitored yearly for clinical features and with DXA every 2-3 years. For moderate to high-risk patients, oral bisphosphonates should be started and are still the first-line of treatment [18].

Bone disease associated with anticonvulsant therapy is a form of osteomalacia. In this condition, high-turnover osteoporosis is often present. Phenobarbital, diphenylhydantoin,
and carbamazepine, 3 commonly used anticonvulsants, increase the metabolism and clearance of vitamin D. Thus, Rickets has been observed in children taking anticonvulsant medication. In some reports [19,20], rates were as high as 20% to 65%, with patients being at particularly increased risk of fracture during seizures.

Table 3. Causes of drug-induced osteoporosis.

<table>
<thead>
<tr>
<th>Class</th>
<th>Generic name</th>
<th>Indications</th>
<th>Bone loss</th>
<th>Fall risk</th>
<th>Fracture risk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glucocorticoids</td>
<td>Prednisolone</td>
<td>Immune and inflammatory disorders</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Dexamethasone</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Proton Pump Inhibitors</td>
<td>Omeprazole, Esomeprazole</td>
<td>Peptic Ulcer disease</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Anticonvulsants</td>
<td>Valproic acid</td>
<td>Chronic Seizures</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Calcineurin inhibitors</td>
<td>Cyclosporine A</td>
<td>Allogenic Organ Transplantation</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Chemotherapeutic Drugs</td>
<td>Methotrexate, cyclophosphamide</td>
<td>Miscellaneous</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Unfractioned Heparin</td>
<td></td>
<td>Thromboembolic disorders</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Thiazolidinediones</td>
<td>Rosiglitazone, Pioglitazone</td>
<td>Diabetes mellitus Type 2</td>
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<td>+</td>
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</tr>
<tr>
<td>GnRH agonists</td>
<td>Goserelin, Boserelin</td>
<td>Prostate Cancer</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Anti-retroviral drugs</td>
<td>Tenofovir</td>
<td>HIV/AIDS</td>
<td>+</td>
<td>+</td>
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</tr>
<tr>
<td>Aromatase Inhibitors</td>
<td>Anastrazole, Letrozole</td>
<td>HER-positive Breast Cancer</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Progestrone*</td>
<td>Depot-hydroxyprogesterone</td>
<td>Contraception</td>
<td>+</td>
<td>+</td>
<td></td>
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<tr>
<td>Thyroid Hormone</td>
<td>Levothyroxine</td>
<td>Hypothyroidism</td>
<td>+</td>
<td>+</td>
<td></td>
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<tr>
<td>Antidepressants</td>
<td>SSRI</td>
<td>Chronic Depression</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Blood Pressure Medication</td>
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<td>Arterial Hypertension</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td>Diuretics</td>
<td>Furosemide</td>
<td>Fluid retention</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Hydrochlorothiazide</td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Alfa-adrenergic blockers</td>
<td>Tamsulosin</td>
<td>Benign Prostatic Hyperplasia</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
</tbody>
</table>


Table 4. Clinical, imaging and laboratory work-up to evaluate secondary causes of Osteoporosis.

Clinical work-up
- History and Physical examination To identify risk factors for fracture, the underlying disease and potential drugs.

Imaging work-up
- Dual-Energy X-Ray Absorptiometry (DXA) To quantify bone mineral density
- Spinal X-Rays To detect prevalent vertebral fractures To exclude osteolytic lesions or tumors

Laboratory work-up
- Complete blood count Anemia as in Multiple Myeloma, Celiac Disease Leukocytosis in Leukemia
- Renal and liver function tests Renal and liver failure, alcohol abuse
- Serum calcium and phosphate Primary Hyperparathyroidism, Multiple Myeloma
- Serum C-reactive Protein Chronic Infection / Inflammation
- Serum bone specific or total ALP Paget’s Disease, Osteomalacia
- Intact parathyroid hormone Primary Hyperparathyroidism
- 25-Hydroxyvitamin D Vitamin D deficiency, Osteomalacia
- Serum levels of basal TSH Thyroid dysfunction
- Fasting blood glucose Diabetes Mellitus
- Serum protein electrophoresis MGUS, Multiple Myeloma
- 24 hour urinary calcium excretion Hypercalciuria
- Anti-tissue transglutaminase levels Celiac Disease
- Anti HIV antibodies HIV/ AIDS
- Morning fasting serum cortisol levels* Cushing’s Syndrome
- Serum tryptase levels, Urinary histamin excretion Systemic Mastocytosis
- COL 1A genetic testing Osteogenesis Imperfecta

*After dexamethasone suppression; ABB: ALP: Alkaline Phosphatase; COL 1A: Collagen Type I Alpha 1; COPD: Chronic Obstructive Pulmonary Disease; HIV: Human Immunodeficiency Virus; MGUS:Monoclonal Gammopathy Of Undetermined Significance; TSH: Thyroid Stimulating Hormone
In the outpatient setting, abnormalities on bone biopsy specimens, such as increased osteoid, are observed in 10% to 40% of patients receiving long-term anticonvulsant therapy. However, if the patient is well nourished and exposed to adequate amounts of sunlight, clinically significant bone disease is less likely to occur.

Many retrospective studies have showed that long-term use of proton-pump-inhibitors (PPI) is associated with an increased risk of fragility fractures [21], but the underlying mechanisms have not been clarified yet. Chronic acid suppression caused by long-term proton pump inhibitor therapy may play a crucial role in decreased absorption of calcium and vitamin B12 and, therefore, indirectly affecting the bones resulting in decreased BMD. The available data suggest that proton pump inhibitors should be used with caution in patients with increased risk of osteoporosis.

**Inflammatory Rheumatic Diseases**

In inflammatory rheumatic diseases such as rheumatoid arthritis (RA), psoriatic arthritis (PsA) and ankylosing spondylitis (AS) the systemic burden caused by inflammation, glucocorticoid treatment, immobilization and reduced physical activity due to painful joints and muscle weakness are associated with decreased bone mass and deterioration of bone quality leading to increased risk of falls and fragility fractures [22]. The reason for this effect on bone is thought to come from the local and systemic action of pro-inflammatory cytokines, such as an increased expression of receptor activator of nuclear factor κB (RANKL), macrophage-colony stimulating factor (M-CSF) and the presence of Tumor Necrosis Factor (TNF α), IL-1, IL-6, IL-7, and IL-17. Other factors probably contributing to the increased risk of fracture in this population include older age, lack of physical exercise, long-term use of corticosteroids and high disability index [23,24].

The frequency of occurrence of osteoporosis in patients with inflammatory rheumatic diseases, including RA, is about 50% and the course of osteoporosis is closely connected with the activity of the underlying disease [25].

With regards to RA, local and systemic phenomena of bone loss occur, and they seem to be intimately related. Early in the course of the disease reduced BMD within joints occurs, called periarticular osteoporosis, which is associated with the destructive articular processes of bone erosion and joint space narrowing. In the course of RA, especially with longstanding active disease, secondary osteoporosis occurs. Currently, it is thought that the mechanisms causing periarticular and secondary osteoporosis are at least partly the same [26].

Unlike postmenopausal osteoporosis, osteoporosis in RA is characterised by a marked loss of cortical bone (hip and the radius), while the axial bone is relatively preserved [24], except when high cumulative doses of glucocorticoids are used.

In contrast with RA, AS is associated with both osteopenic changes and bone forming phenomena. The reported prevalence of osteoporosis in AS patients varies largely and this variation reflects the difficulties in assessing BMD in AS due to new bone formation. There are also inflammation-induced structural changes in the spine predisposing to vertebral wedge and fracture. In a prospective cohort study [27] involving 504 AS patients there was a significantly higher prevalence of OP (9.7% vs 0%) and osteopenia among AS patients (57.5% vs. 34.9%) comparing with healthy controls. The BMD was significantly lower in the patients with higher elevated erythrocyte sedimentation rate (ESR) or C-reactive protein (CRP). Risk factors for lumbar spine bone loss were Juvenile onset, morning stiffness lasting over 30 minutes and elevated ESR levels, which correlate with the inflammatory burden of the disease.

A case-control study to assess the prevalence of fragility fractures in patients with PsA [28] found that disease duration, age and femoral neck BMD were associated with increased rate of non-vertebral fractures and a large population-based cohort study [29] including 9788 PsA patients, 158,323 psoriasis patients and 821,834 matched controls reported that PsA and psoriasis patients had a 7–26% increased incidence of fracture when compared to the general population.

**Endocrine Dysfunction**

Primary hyperparathyroidism is a common disorder, with an incidence of 1 in 500 to 1 in 1000, and is usually asymptomatic [30]. Classically, primary hyperparathyroidism is associated with osteitis fibrosa cystica characterized by subperiosteal resorption of the distal phalanges, tapering of the distal clavicles, a “salt and pepper” appearance of the skull, brown tumors, and bone cysts. With the widespread
availability of screening blood tests for asymptomatic hypercalcemia, the incidence of this manifestation has decreased dramatically over the years. Osteopenia and osteoporosis are also recognized as bone diseases associated with excess parathyroid hormone (PTH). The increase in circulating levels of PTH in primary hyperparathyroidism is associated with increased bone turnover, both in osteoclast-mediated bone resorption and osteoblast activity, leading to the loss of cortical and trabecular bone. In mild hyperparathyroidism, however, BMD may be increased in areas that are primarily trabecular, whereas bone is lost in the cortical areas [31]. This anabolic effect is the basis for the treatment with PTH analogues. Nevertheless, bone resorption is enhanced by sustained elevations in the levels of PTH.

In a longitudinal cohort of patients with primary hyperparathyroidism [32], bone density has been measured at 3 sites to evaluate cortical bone, trabecular bone, and a mixture of both. At the distal radius, an area rich in cortical bone, BMD was less than 80% of age- and sex-matched controls. In contrast, at the lumbar spine, BMD was relatively well preserved. The values for the hip region, which is made of mixed trabecular and cortical bone, were midway between the data obtained for the spine and the distal radius. This finding is consistent with the observation that PTH mobilizes calcium from cortical sites before it has a negative impact on trabecular skeleton. Quantitative histomorphometric analysis of bone biopsy specimens are consistent with the loss of cortical bone and preservation of cancellous bone [31]. Cortical thinning is noted on biopsy specimens, and PTH levels correlate with cortical porosity [32].

After surgical cure of primary hyperparathyroidism, BMD increases in the forearm and lumbar spine. In a longitudinal cohort [33] of patients with primary hyperparathyroidism followed up for 10 years, parathyroidectomy resulted in normalization of biochemical values and increased BMD. The increase in bone density was prompt and sustained, but a trend toward further increase after 1 year was significant only for femoral neck values. In a subset of patients who did not undergo surgery, there was no progression of bone disease if they were asymptomatic, but one quarter had some progression with bone loss.

Regarding the thyroid gland, both thyroid hormone insufficiency and excess can lead to alterations in bone mass. Thyroid hormone increases the creation of new bone remodeling units with an enhancement of remodeling activity. Thyroid hormones directly stimulate production of osteocalcin, alkaline phosphatase, and insulin-like growth factors (IGF).

In patients with thyrotoxicosis, increased serum levels of osteocalcin and alkaline phosphatase may be seen. Despite the increase in osteoblast activity, there are also thyroid hormone–induced increases in bone resorption.

In the thyrotoxic patient, the bone remodeling cycle is shortened because of a decrease in the length of the bone formation and, overall, there is failure to replace resorbed bone completely, leading to bone loss. In patients with thyrotoxicosis, BMD is reduced [34,35].

Several studies indicated that individuals with a history of thyrotoxicosis have an increased risk of fracture and may sustain fracture at an earlier age compared with patients who have never had an increase in thyroid hormone levels. After effective treatment of the thyrotoxic patient, the decrease in BMD may be reversible. Normalization of the results of thyroid function tests results in increased BMD comparing with pre-treatment values [36].

In a meta-analysis [37], BMD was assessed in women receiving thyrotropin-suppressive doses of thyroxin. The study concluded that there was a 1% increase in annual bone loss in postmenopausal women. A large prospective study, the Study of Osteoporotic Fractures [38], examined the relationship between thyroid disease and fractures. In this study, postmenopausal women with a history of hyperthyroidism had an 80% increased risk of subsequent hip fracture. Thyroid hormone use itself was associated with a 60% increase in fracture risk.

### Eating Disorders

Anorexia nervosa and bulimia are associated with significant morbidity and mortality and are chronic in nature. They affect 5% to 10% of women and the onset may be at any time from adolescence through the fourth decade of life. Anorexia nervosa has been associated with osteoporosis. There are several metabolic disorders associated with anorexia nervosa that may adversely affect bone metabolism. These include estrogen deficiency, endogenous cortisol excess, reduced IGF-1 levels, protein-energy malnutrition, and secondary
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hyperparathyroidism due to low dietary calcium intake or vitamin D deficiency.

It has been estimated that 50% of anorexic patients have BMD values at the lumbar spine that are more than 2 SDs below those of age-matched, healthy controls [39].

Total alkaline phosphatase activity may be elevated, but liver enzyme levels are also elevated. Osteocalcin has been noted to be very low in women with anorexia nervosa and may be due to the excess endogenous cortisol levels. Markers of bone resorption, such as pyridinoline and N-telopeptide excretion are usually increased.

Celiac Disease

Regarding bone involvement in celiac disease, it can present as osteoporosis, osteomalacia, or both. Untreated adults usually present with reduced bone mineral at the time of diagnosis, whereas children may present with growth retardation.

Chronic inflammatory intestinal diseases, including celiac disease, can affect bone and mineral metabolism due to changes in both systemic and local regulatory factors. The pathogenetic processes are still controversial, but two main mechanisms seem to be involved: intestinal malabsorption and the presence of chronic inflammation [40].

A strict and lifelong gluten free diet can help recover normal bone density when a diagnosis of Celiac disease is made in children and adolescents [41,42].

In the case of adult celiac disease with associated bone disease, a gluten free diet rarely normalizes BMD in adulthood [40, 43].

A cross-sectional study [44] as shown that despite long-term strict adherence to gluten free diet (GFD), 74% of patients displayed low BMD. Among these, 24% had osteoporosis and 76% osteopenia. Therefore, nutritional supplementation should be considered for all patients with celiac disease.

Patients may present with normal serum biochemical analysis results or with reduced serum and urine calcium levels and elevated alkaline phosphatase levels. With GFD, biochemical abnormalities and BMD measurements may improve [44].

Alcohol Consumption

Alcohol abuse is a major cause of liver disease, and cirrhosis contributes to the severity of bone disease. On the other hand, alcohol directly inhibits bone cell activity, resulting in reduced bone formation and increased bone resorption.

Spinal osteopenia may be observed in up to 50% of ambulatory patients with alcoholism, and fractures of the ribs or vertebrae occur in 30% of this population [45]. Osteoporosis is usually the predominant disease, although osteomalacia may occur. Sex differences occur in that chronic alcohol abuse has a detrimental effect on the male skeleton, whereas a neutral or beneficial effect with light-to-moderate alcohol consumption is seen on the female skeleton. Trabecular bone is more affected than cortical bone when evaluating BMD.

Serum concentrations of calcium, phosphorus, and magnesium tend to be in the low normal range. Poor nutrition contributes to the picture, with a reduction in serum albumin levels. Serum PTH levels may be elevated or high normal and may be due to short-term administration of alcohol or the low calcium and magnesium levels. 25-hydroxyvitamin D levels are usually low, and 1,25-dihydroxyvitamin D levels have been reported as low, normal, or high.

Cessation of alcohol consumption can reverse or at least stop progression of disease. If 25-hydroxyvitamin D levels are low, cholecalciferol therapy should be considered. There is not enough available data on the effect of alcohol cessation and the reversibility of osteoporosis.

Conclusion

An appropriate diagnostic workup should be performed for patients presenting with fragility fractures or those presenting with risk factors for secondary osteoporosis. As shown in Table 4, this should include a careful clinical history and physical examination, imaging work-up with DXA to quantify BMD and spinal X-rays to detect prevalent vertebral fractures and exclude osteolytic lesions or tumours. In terms of laboratory work-up, a complete blood count should be performed to exclude anaemia (as in multiple myeloma, celiac disease or leukaemia), renal and liver function tests to exclude renal and liver impairment as well as alcohol abuse, serum calcium and phosphate as well as intact parathyroid hormone to exclude hyperparathyroidism, ESR and CRP to exclude chronic infection/inflammation,
25-hydroxyvitamin D to exclude vitamin D deficiency and osteomalacia, serum bone specific or total alkaline phosphatase for Paget’s disease of the bone, serum protein electrophoresis for monoclonal gammopathy of undetermined significance (MGUS)/multiple myeloma and a 24 hour urinary calcium excretion to check for hypercalciuria.

References


6. Rodrigues AM 2018


36. Faber J, Gallowe AM. Changes in bone mass during prolonged subclinical

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