Review Article

Bisphophonates in CKD Patients with Low Bone Mineral Density

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Patients with chronic kidney disease-mineral and bone disorder (CKD-MBD) have a high risk of bone fracture because of low bone mineral density and poor bone quality. Osteoporosis also features low bone mass, disarranged microarchitecture, and skeletal fragility, and differentiating between osteoporosis and CKD-MBD in low bone mineral density is a challenge and usually achieved by bone biopsy. Bisphophonates can be safe and beneficial for patients with a glomerular filtration rate of 30 mL/min or higher, but prescribing bisphosphonates in advanced CKD requires caution because of the increased possibility of low bone turnover disorders such as osteomalacia, mixed uremic osteodystrophy, and adynamic bone, even aggravating hyperparathyroidism. Therefore, bone biopsy in advanced CKD is an important consideration before prescribing bisphosphonates. Treatment also may induce hypocalcemia in CKD patients with secondary hyperparathyroidism, but vitamin D supplementation may ameliorate this effect. Bisphosphonate treatment can improve both bone mineral density and vascular calcification, but the latter becomes more unlikely in patients with stage 3-4 CKD with vascular calcification but no decreased bone mineral density. Using bisphosphonates requires considerable caution in advanced CKD, and the lack of adequate clinical investigation necessitates more studies regarding its effects on these patients.

1. Introduction

At the onset of chronic kidney disease (CKD), the systemic mineral metabolism and bone composition start to change. This alteration is known as CKD-MBD. The greater the decrease in renal function, the worse the progression of CKD-MBD. CKD-MBD involves serum calcium, serum phosphate, parathyroid hormone (PTH), and vitamin D metabolism derangement, and its main endpoints are altered bone turnover, bone mineralization, bone volume, bone linear growth, bone strength, and vascular/other soft tissue calcification [1].

Those who have CKD-MBD may have another bone problem, osteoporosis, which is the most common metabolic bone disease resulting in fragility fractures. With increasing lifespans, the population with osteoporosis is growing [2, 3], and, in the general population, about 85% of women with osteoporosis have some deterioration of renal function [4]; both are probably attributable to greater age [5]. However, osteoporosis also involves low bone mass, a disarranged microarchitecture, and skeletal fragility, and thus this condition contributes to the risk of fracture, especially in the spine, hip, wrist, humerus, and pelvis [6].

Because fragility fractures, reduced glomerular filtration rate (GFR), and low bone mineral density (BMD) are common in the older population, it can be a challenge to differentiate the cause of a fragility fracture and/or low BMD in elderly CKD (particularly with estimated eGFR <30 mL/minute) patients as either osteoporosis or another bone and mineral disorder related to CKD (e.g., hyperparathyroidism, adynamic bone disease, and osteomalacia) [7] (Table 1). Patients with an age-associated reduction in
Table 1: Bone diseases may associate with fragility fractures (low trauma fractures).

| Name of bone diseases                          | General metabolic bone diseases | Chronic kidney disease-mineral bone disease (CKD-MBD) |
|-----------------------------------------------|---------------------------------|-----------------------------------------------------|
| Adynamic bone disease (including aluminum bone disease) | Yes (AL-amyloidosis.)         | Yes (β₂-microglobulin)                              |
| Amyloid bone disease                          | Yes                             |                                                     |
| Mixed uremic osteoporosis                     | Yes (primary hyperparathyroidism) | Yes (secondary hyperparathyroidism)                 |
| Osteitis fibrosa cystica (severe)             | Yes                             |                                                     |
| Osteogenesis imperfecta                      | Yes                             |                                                     |
| Osteomalacia                                  | Yes                             |                                                     |
| Vitamin D-related                            | Yes                             |                                                     |
| Nonvitamin D-related                         | Yes                             |                                                     |
| Chronic metabolic acidosis                    | Yes                             |                                                     |
| Phosphate depletion                           | Yes                             |                                                     |
| Osteoporosis                                  | Yes                             |                                                     |
| Primary osteoporosis                          | Yes                             |                                                     |
| Secondary osteoporosis (all causes)           | Yes                             |                                                     |
| Chronic liver disease                         | Yes                             |                                                     |
| Hypogonadism or premature menopause          | Yes                             |                                                     |
| Inflammatory bowel disease                    | Yes                             |                                                     |
| Malabsorption                                | Yes                             |                                                     |
| Steroid-induced osteoporosis                  | Yes                             |                                                     |
| Pathologic fractures (malignancies)           | Yes                             |                                                     |
| Paget's disease                               | Yes                             |                                                     |

eGFR to 30 mL/min benefit from oral or intravenous bisphosphonates for osteoporosis [8, 9], but the use of bisphosphonates in CKD requires some caution. Primary management of CKD with low mineral density is essential for understanding the pathogenesis of these two bone disorders and designing a rational approach to treatment and prevention of complications.

2. Definition of Osteoporosis

Osteoporosis is a gradual systemic skeletal disease that usually does not cause any symptoms, so many patients do not know that they have it until a bone fracture unexpectedly occurs. In general, osteoporosis is affected by age, sex, body size, ethnicity, family history, sex hormones, and lifestyle.

The structures of the bone consist of the bone matrix, bone cells, and mineral salts. Bone quality is influenced by bone mass, bone density, bone geometry (shape and size), microarchitectural features (cortical or trabecular connections), and molecular elements (collagen type and linkages, bone mineral composition, and crystal orientation). Bone cells include osteoblasts, which form the bone matrix and produce increased bone mass, and osteoclasts, which resorb the bone matrix by secreting acids and digestive enzymes and triggering osteocyte apoptosis and decreased bone mass. In healthy adult bone, the new bone formation will be followed by bone resorption to keep the bone mass in a dynamic balance. However, in osteoporosis, the balance shifts toward bone resorption even though bone mineral and protein are normal, leading to thinning of bone mass and increased fragility [10, 11].

According to 1994 World Health Organization (WHO) criteria, osteoporosis can be defined as a BMD T-score < −2.5, as measured by dual-energy X-ray absorptiometry (DXA) at the hip, spine, or forearm [12]. This measure can be applied in both men and women, as well as in younger people with medical problems along with increased risk of low-trauma fracture [13, 14]. This T-score is an important diagnostic tool for defining normal density (+1 to −1), osteopenia (−1 to −2.5), and osteoporosis (<−2.5) in clinical practice. Those with fractures who subsequently undergo BMD testing, however, are more often found to have osteopenia than osteoporosis. One reason is that many other factors not associated with low BMD contribute to fragile bone [15, 16]. Thus, the National Institutes of Health has stated that osteoporosis as “a skeletal disorder characterized by impairment in bone strength predisposing a person to an increased risk of fracture. Bone strength primarily reflects the integration of bone density and bone quality” [17]. This statement does not serve, however, as a diagnostic tool, and, in the general adult population, clinical diagnosis of osteoporosis is made based on one of two conditions: the presence of a low-trauma fracture unrelated
to BMD level, or in the absence of a preexisting fracture, a certain level of BMD defined by the standard deviation of the T-score [18].

Although the low BMD or fragility fracture is a clear guide for osteoporosis, they do not aid in differentiating cause from metabolic bone diseases that also can lead to fragility fractures. These metabolic bone diseases can be related not only to osteoporosis but also to osteomalacia and adynamic disorders, among others [19]. In addition, DXA checks only area BMD but not bone volumetric BMD, which cannot evaluate bone microarchitecture or bone turnover. To overcome these limitations, some new noninvasive three-dimensional technologies have been developed to evaluate bone microarchitecture, including high-resolution micro-CT and micro-MRI, hip structural analysis, and finite element analysis. Few data with these techniques have been related to CKD patients, however [20].

3. Bone Remodeling in CKD

3.1. Remodeling. Physiological bone remodeling is a lifelong and highly coordinated process of bone resorption and formation [21, 22]. It involves continuous removal of old bone, replacement with a newly synthesized proteinaceous matrix, and subsequent mineralization of the matrix to form new bone. Therefore, maintaining mineral homeostasis is another essential role of bone remodeling [23]. This resorption and reformation cycle is crucial for repairing bone microfractures and modifying bone structure during stress or other responses to biomechanical forces. Remodeling in the adult bone is not entirely clarified, but the main purposes are removing dead osteocytes, maintaining oxygen and nutrition supply, keeping a suitable level of matrix blood supply, and correcting fatigue damage [24].

The stages of bone remodeling begin with osteoclast resorption and move to reversal, osteoblast maturation, osteoid or unmineralized bone formation, and mineralization to the quiescent stage. In resorption, the osteoclasts gather on the bone surface, releasing acid and hydrolytic enzymes to resorb bone and digest bone minerals and fragments of collagen. The resorption process leads to circulation of free pyridinoline and deoxypyridinoline substances circulating in the blood and eliminated in the urine [25–27]. Bone formation starts from the osteoblasts synthesizing type I collagen and other proteins (e.g., osteocalcin); all of these particles aggregate extracellularly to form osteoid, and then bone is mineralized [22]. The cell membranes of osteoblasts contain alkaline phosphatase. The synthesis of type I collagen is a combination of one alpha-2 and two alpha-1 polypeptide chains to shape a coiled construction known as procollagen [21].

The bone remodeling process occurs through the action of osteoprotegerin (OPG) and the RANKL/RANK (receptor activator of nuclear factor kB; NF-kB ligand/receptor activator of nuclear factor kB; NF-kB) system, which is affected by hormones (PTH, calcitriol, estrogen, and glucocorticoids), cytokines, and interleukins. Increased PTH levels expand the destroyed surface, osteoid surface, and mineralizing surface, which enhance the amount of osteoclasts and osteoblasts [28]. Avbersek-Luznik et al. showed that PTH can enhance bone resorption through a mechanism involving increased osteoblast RANKL synthesis [29]. RANKL/RANK interaction provides essential signals to osteoclast progenitors, leading to osteoclast differentiation, an important step in osteoclastogenesis, and bone resorption. This process can be blocked by OPG, a decoy receptor for RANKL that protects against bone resorption and extensive deterioration [30].

3.2. Bone Turnover. Bone strength is determined by its mass, microarchitecture, macrogeometry, and rate of turnover. Bone turnover occurs in both cortical bone and trabecular bone, and the latter has a relatively higher turnover rate [23]. Therefore, this important step of bone turnover is orchestrated by a tightly coupled interplay of osteoclasts and osteoblasts to constantly replace dead bone with new bone [31].

In CKD patients, the effect of high levels of PTH on bone results in the high-turnover bone disease, osteitis fibrosa, with excessive osteoclastic bone resorption and bone marrow fibrosis [32]. In contrast, low-turnover bone disease is common in patients with CKD, especially in dialysis patients, and is characterized by an extremely slow rate of bone formation [19].

Osteomalacia, aluminum-induced bone disease, and adynamic bone disease are low-turnover bone diseases. Osteomalacia is characterized by defective bone mineralization and very slow bone formation rate [33]. This mineralization defect is related to reduced active vitamin D and chronic metabolic acidosis [34]. Aluminum ingestion causes another mineralization defect, owing to reducing both osteoclast resorption and the osteoblast surface, and is associated with low-turnover bone disease. In addition, chronic low-dose aluminum exposure with high intake of vitamin D in dialysis patients reduces PTH synthesis and secretion. These patients may present with adynamic bone disease rather than osteomalacia [35].

4. Diagnosis of Osteoporosis or Bone Disease in CKD

4.1. Osteoporosis versus Bone Disease in CKD. It is not uncommon that CKD patients develop impairment of bone strength and low-trauma fractures, the latter because of osteoporosis or some other metabolic bone disease associated with CKD. The challenge for clinical practice is to differentiate osteoporosis with CKD-MBD in fracturing patients. However, there are some clues that can be generalized.

First, in early CKD (stages 1 to 3), patients may present with few abnormalities of mineral metabolism and have the same osteoporosis risk factors as the general population. The Kidney Disease: Improving Global Outcomes (K/DIGO) guidelines mention that early CKD metabolic status may involve intermittent hyperphosphatemia, mild increases in PTH, or other bone turnover marker changes without altered bone strength or fragility fractures [36]. Hence, low-trauma
fractures in early CKD without abnormalities of vitamin D metabolism and PTH can be diagnosed using the WHO osteoporosis criteria, a T-score of −2.5 or lower or fragility fractures, as in the postmenopausal population. In other words, these patients without biochemical abnormalities of CKD-MBD can receive standard treatment [18].

Second, in advanced CKD (stages 4 to 5, 5D), there are no universally accepted criteria for making the diagnosis of osteoporosis [12]. The significant derangements in bone and mineral metabolism may lead to more impairment in bone strength and increased risk for low-trauma fractures. Patients with stage 5D CKD may have a higher risk for fragility fractures than postmenopausal women or elderly men do. These bone fragility conditions are often confounded on DXA with BMD increased due to calcified soft tissue in the path of the X-ray beam or altered bone composition [37]. Therefore, the K/DIGO guidelines do not recommend that patients with stage 3 to 5 CKD receive regular BMD DXA [38].

4.2. Diagnosis of Bone Disease in Advanced CKD. As previously described, there are no accredited standards for diagnosing osteoporosis in advanced CKD. For now, the best way to differentiate between osteoporosis and CKD-MBD is bone histomorphometry and/or bone biochemical markers of bone metabolism to describe the bone disease that may be identified for low-trauma fractures in advanced CKD [12]. The K/DIGO guidelines suggest bone biopsy for unexplained fractures, persistent bone pain, unexplained hypercalcemia, high PTH but low alkaline phosphatases, and before therapy with bisphosphonates [38]. To describe the various abnormalities in CKD-MBD, the recent recommendation from the K/DIGO classification of renal metabolic bone disease uses the TMV (turnover: from low to high; mineralization: from normal to abnormal; and bone volume: from low to high) but also does not provide a working diagnosis of osteoporosis [1]. Although many useful radiologic techniques can examine bone microarchitecture to offer the potential for defining turnover, mineralization, and volume noninvasively in advanced kidney disease, they cannot distinguish between renal osteodystrophy and osteoporosis [20, 39–41]. In the future, these noninvasive imaging techniques may lead to methods for connecting turnover, mineralization, and volume to bone strength and open up a totally new way to categorize bone strength [18].

5. Bisphosphonates in Advanced CKD

5.1. Bisphosphonate Pharmacokinetics. Bisphosphonates are standard drugs for osteoporosis and are like pyrophosphate compounds (Figure 1), which are crucial endogenous inhibitors of ectopic mineralization. In postmenopausal women, effective bisphosphonate treatment results in a quantifiable reduction in urinary excretion of N-telopeptide of type I collagen and improvement in bone density [42]. The bone lining cells prevent pyrophosphate from entering the structure by cleaving it with alkaline phosphatase [43]. Because a carbon atom substitutes for the oxygen in pyrophosphate, bisphosphonates are resistant to cleavage by alkaline phosphatase. Therefore, bisphosphonates can enter bone by binding to the mineralized surface [44].

The intestinal absorption of bisphosphonates is low (e.g., 1% for alendronate, ibandronate, and risedronate and 3% to 7% for etidronate [45]) and will be even less when taking medicines without a glass of water or on an empty stomach. When bisphosphonates are absorbed, about 40%–60% enters bone; the rest is excreted unchanged through glomerular filtration with active proximal tubular secretion [46]. Bisphosphonates stack up in bone, with a more than 10-year half-life [47], and are slowly released back into the circulation and taken up again or excreted.

As bone-seeking antireceptive agents [12], bisphosphonates can bind strongly to hydroxyapatite in bone. Bisphosphonates have a carbon atom with two phosphate groups, forming a P-C-P structure that is a stable analogue of inorganic pyrophosphate [48]. Their effects on hydroxyapatite crystals may influence the overall action of osteoclasts, but the most essential effect is on osteoclasts themselves. During osteoclastic bone resorption, bisphosphonates impair osteoclast cell function by inhibiting enzyme activity [47]. Their mechanism of action, "osteoclast inhibition," makes them a valuable treatment option in all bone disease associated with increased osteoclast activity. In addition, bisphosphonates can reduce the progression of soft tissue calcification, and recent studies have shown that bisphosphonates have the potential to reduce the progression of vascular calcification [49–51]. Therefore, other roles for bisphosphonates include reducing progression of vascular calcification in CKD.

5.2. Mechanisms of Action of Bisphosphonates. Bisphosphonates can be divided into two molecular groups (Table 2): nonnitrogen-containing bisphosphonates (clodronate and etidronate) and nitrogen-containing bisphosphonates (alendronate, ibandronate, pamidronate, risedronate, and zolendronate). When they enter bone and bind with osteoclasts, the latter group is more effective in improving bone strength.
The molecular structure of the nonnitrogen-containing group is very similar to that of pyrophosphate. Thus, these bisphosphonates are incorporated into nonhydrolyzable analogs of ATP, which may limit ATP-dependent enzymes in osteoclasts to block mitochondrial energy production and induce osteoclast apoptosis [52].

The nitrogen-containing bisphosphonates are internalized by endocytosis of osteoclasts and go through other specific biochemical processes to apoptosis (Figure 2). At the same time, they restrict enzymes of the mevalonate pathway and inhibit farnesyl pyrophosphate (FPP) synthase [44]. Furthermore, this inhibition of FPP blocks the prenylation of small GTPase signaling proteins (e.g., Ras, Rho, and Rac), which are essential for osteoclast function and survival [53]. Therefore, the most important key to inhibiting small GTPase binding proteins is to block the biosynthesis of isoprenoid compounds. This capacity can explain the loss of osteoclast activity and function as a result of inhibition of protein prenylation and the disruption of the function of these essential regulatory proteins.

In addition, during bone resorption, the secretion of vacuolar-type proton pumps on the ruffled border of the osteoclast membrane acidifies the space beneath osteoclasts to dissolve bone mineral. When bisphosphonates adsorb to bone mineral, the extracellular bone matrix is destroyed by the action of proteolytic enzymes, and the acidic microenvironment results in the dissolution of the hydroxyapatite bone mineral because bisphosphonates accumulate at the sites of bone resorption where the mineral is most exposed [54, 55]. Osteoclasts, as the most bisphosphonate-attractive cells in bone, take up bisphosphonate separated from bone mineral in the acidified environment beneath osteoclasts. Hence, osteoclasts are exposed to the highest concentrations of free, nonmineral-bound bisphosphonate [48].

Nitrogen-containing bisphosphonates can affect osteoclast functions: terminal differentiation, attachment, endocytosis, cell shape, and apoptosis. In addition to influencing osteoclast function, this type of bisphosphonate can block FPP production in monocytes to gather plenty of isopentenyl diphosphate (IPP). IPP is a bacterial antigen that stimulates T cells to release THF-α and INF-Υ and to increase IL-6 and CRP levels. This response is why patients receiving their first bisphosphonate treatment often complain of flu-like symptoms [56].

5.3. Bisphosphonate in CKD with Low BMD. In both the general population and patients with stages 1 to 3 CKD, proper treatment with bisphosphonates can prevent fracture [57]. A meta-analysis of postmenopausal women with CKD stages 1 to 3 in nine clinical trials concluded that it is

### Table 2: Two classes of bisphosphonates.

| Bisphosphonates | Agents | R1 side chain | R2 side chain |
|-----------------|--------|--------------|--------------|
| Nonnitrogenous  | Etidronate | OH | CH<sub>3</sub> |
|                 | Clodronate | Cl | Cl |
|                 | Tiludronate | H | ClS |
| Nitrogenous     | Pamidronate | OH | NH<sub>2</sub> |
|                 | Neridronate | OH | NH<sub>2</sub> |
|                 | Olpadronate | OH | N |
|                 | Alendronate | OH | N |
|                 | Ibandronate | OH | |
|                 | Risedronate | OH | |
|                 | Zoledronate | OH | |

The bisphosphonates are analogous to that of the naturally occurring pyrophosphates, with two-side chains (R1 and R2) attached to the carbon core. The R1 side chain determines bone-binding affinity, and the R2 side chain determines interception potency.
safe to provide bisphosphonates to low-BMD patients without secondary causes or deranged blood levels of calcium, phosphate, PTH, or alkaline phosphatase, and vitamin D abnormalities (laboratory features of CKD-MBD) to reduce fractures [58]. From several prospective studies and clinical trials, oral or intravenous bisphosphonates can offer benefits to patients with aged-related decreased eGFR, down to 30 mL/min [8, 59]. Other researchers as well suggest that it is safe to use bisphosphonates in patients with an eGFR of 30 mL/min or higher [60, 61]. However, these drugs might not be safe when the patients have preexisting renal parenchymal disease (e.g., diabetes) or use other agents that could affect renal function (e.g., nonsteroidal anti-inflammatory drugs). Therefore, when using intravenous bisphosphonates in these specific higher-risk secondary renal disease groups, care is required [18].

The value of bisphosphonate treatment in patients with advanced CKD is vague because suitable data are lacking for CKD stages 4 to 5D (eGFR less than 30 mL/min). Some patients with stages 4 to 5 CKD with fractures and BMD

Figure 2: Effects of nitrogenous bisphosphonate on mevalonate metabolism (right) and on the osteoclasts of bone and calcified vessels (left).
levels in the osteoporotic range may derive benefits from the administration of bisphosphonates [62], as may some dialysis patients with osteoporosis due to gonadal hormone deficiency such as postmenopausal osteoporosis, glucocorticoid-induced osteoporosis, or male osteoporosis [63].

Even though data are limited describing bisphosphate use in stage 5 CKD, a randomized placebo-controlled study comparing alendronate and placebo in 31 hemodialysis patients proved that hip BMD remained stable after 6 months with alendronate while BMD fell in those treated with placebo [64]. When considering prescribing bisphosphonates to CKD stage 5 patients suffering fragility fractures, renal osteodystrophy must be thoroughly ruled out [65, 66]. Because bone biopsy is not available in most clinics, in patients with CKD-MBD, who already have low bone turnover (e.g., adynamic bone disorder; bone alkaline phosphatase <20ng/mL and iPTH < 100 pg/mL) [67], the use of bisphosphonates may aggravate clinical symptoms. Therefore, in advanced CKD, a bone biopsy should be considered before providing bisphosphonates, and therapy should involve individual-specific conditions. In patients with advanced CKD, only those who have low BMD and high bone resorption might receive bisphosphonates because these drugs have not been used to prevent fractures in people with normal BMD or with low bone formation.

Current studies show that a 4-hour regular hemodialysis can remove 35% to 40% of the administered bisphosphate dose, which is close to renal bisphosphate elimination in normal renal function [68]. But these data are not sufficient evidence for other different bisphosphonates, different dialysis membranes, or the effect of bisphosphonate administration timing related to dialysis timing [69]. Lu et al. (2003) found that pamidronate therapy is associated with reduced plasma iCa levels and increased PTH secretion, resulting in aggravated secondary hyperparathyroidism in postmenopausal hemodialysis patients with SHPT with short-term therapy [70]. However, Huang et al. (2012) [71] demonstrated that, in considering pamidronate treatment in postmenopausal patients with osteoporosis receiving hemodialysis, it might be safer to add calcitriol to prevent increased PTH secretion [71]. Bisphosphate treatment may induce hypocalcemia, and in CKD patients with secondary hyperparathyroidism, vitamin D supplementation may ameliorate this hypocalcemic effect. Therefore, for patients with early CKD (stages 1–3), nutritional vitamin D (cholecalciferol or ergocalciferol) and calcium should be provided. However, active vitamin D (calcitriol or paricalcitol) could be considered when advanced CKD or dialysis patients receive bisphosphonate treatment.

When managing dialysis patients with osteoporosis, Miller has suggested cutting the bisphosphonate dose to half of the FDA-registered dose for postmenopausal osteoporosis, simply based on limited pharmacokinetic and dialysis data. Moreover, Miller advises limiting treatment to 2-3 years because the reuptake of bisphosphonates might lead to accumulation and unknown bone retention of bisphosphonates in this population [63].

5.4. Bisphosphonate and Atherosclerosis with Vascular Calcification

5.4.1. Bisphosphonates and Atherosclerosis. In soft blood vessel tissue, bisphosphonates not only have high affinity for calcium in atherosclerotic deposits but also enter the arterial wall by macrophage phagocytosis [72]. During phagocytosis, bisphosphonates change the function of macrophages to internalize atherogenic LDL cholesterol, consequently forming LDL into foam cells [73]. Hence, bisphosphonates stimulate macrophage apoptosis through inhibiting intracellular enzymes as well as inhibiting sterol biosynthesis [74]. However, some studies have found that etidronate, pamidronate, and clodronate suppress developing atherosclerosis without influencing cholesterol or lipid levels [74–77].

A study of etidronate therapy showed that four cycles (200 mg daily for 2 wk every 3 months) of etidronate treatments resulted in a remarkable decrease in intima-media thickness. But the selection criteria of participants for those receiving etidronate (on the basis of low BMD) differed from those using placebo (no BMD criteria) [78].

5.4.2. Bisphosphonates and Vascular Calcification. Multiple risk factors cause vascular calcification in CKD patients, such as vascular smooth muscle cell (VSMC) differentiation into osteoblast-like cells, high calcium and phosphorus levels in abnormal bone metabolism, inflammation, low levels of circulating and locally produced inhibitors, impaired renal excretion, and current therapies. It is a highly complicated process that includes a complex interaction of calcification inducers and inhibitors similar to osteogenesis but different from passive mineral deposition [50].

High levels of calcium and phosphate can enhance the activity of VSMC sodium phosphate (NaPi) cotransporters, leading to intracellular phosphate concentration elevation [79, 80]. This result will influence Cbfa-1, an essential controller of cellular differentiation, inducing VSMCs to transdifferentiate into the osteoblast phenotype cell [62].

In the 1970s, some results indicated that bisphosphonates would be able to inhibit vascular calcification in both humans and non-human animals [81, 82]. Bisphosphonates do inhibit the expression of TNF-α, which can induce osteogenic transdifferentiation processes and calcium deposition in atheromatous injury of rabbit aorta [77].

Several studies have shown that CKD 4-5 patients usually have adynamic bone disease in bone biopsy samples along with some degree of arterial calcification [83, 84], even with the progression of coronary artery calcification [85]. However, Toussaint et al. [86] found that 51 patients with stage 3-4 CKD treated with alendronate for 18 months in clinical trials did not show inhibition of progression of vascular calcification compared with placebo. Therefore, in CKD patients with low BMD, bisphosphonate treatment may improve both the BMD and vascular calcification, but in CKD patients with vascular calcification but no decreased BMD, bisphosphonate treatment is unlikely to do so [86]. In addition, PTH levels had a meaningful increase with alendronate versus placebo (+3.2 pmol/L [95% CI, 0.8–5.5]; \( P = 0.009 \)) at 18 months [86].
During low bone turnover, serum phosphate does not easily enter the bone, and the bone therefore cannot buffer elevation of the phosphate burden any longer. Cannata-Andia and colleagues suggested that bone inhibition and vascular calcifications create a vicious cycle [87]. Sclerostin, an inhibitor of Wnt signaling, is released from transdifferentiated VSMCs as a defensive response that aims to block the Wnt pathway to reduce the mineralization in the vascular tissue [88]. It not only slows the transformation of VSMCs into osteoblasts but also may overflow to the circulation and lessen bone formation, which in turn weakens the ability of bone to absorb phosphate. Hence, bisphosphonates would worsen bone low turnover, then result in high serum phosphate, which would further increase VSMC transformation to osteoblast-like cells [87, 89].

The mechanism by which bisphosphonates inhibit vascular calcification remains unclear. They may inhibit bone resorption, reduce serum calcium and phosphate, and limit their deposition in the vascular wall [90] or their ability to influence the activity of the VSMC NaPi cotransporter. Other case studies indicate that both etidronate [91] and pamidronate are useful [92] in treating calciphylaxis (uremic arteriolopathy), which is a rare but life-threatening complication of CKD.

5.5. Renal Complications of Bisphosphonates. Monitoring proteinuria in patients when administering bisphosphonates is recommended because several groups have mentioned that bisphosphonates damage kidney tissues [93, 94]. As an oncology medicine, bisphosphonates produce direct nephrotoxicity, particularly when used at high dosage [95]. Administering high-dose IV pamidronate can bring on collapsing focal glomerulosclerosis and resulting nephrotic proteinuria [93]. The FDA suggests adjusting the dosage of zoledronic acid according to the baseline creatinine clearance and administering the infusion over 15 minutes [63]. A fast IV of bisphosphonates has been associated with acute renal failure complication [96–98]. Treatment of pamidronate may cause nephrotic syndrome with renal impairment [94] or acute tubular necrosis with consequent acute renal impairment [99].

Markowitz et al. analyzed bone biopsies from six patients who developed renal failure after treatment with zoledronate and identified tubular atrophy, interstitial fibrosis, interstitial inflammation, and mild-to-moderate vascular disease as the main mechanisms of renal damage [98]. Lower doses of bisphosphonate only rarely induce clinically significant nephrotoxicity [100].

6. Conclusions

CKD patients often suffer from osteoporosis. To diagnosis osteoporosis in CKD patients, WHO criteria and/or low-trauma fractures can be widely used for stages 1 to 3 CKD, but there is no consensus regarding the criteria for the diagnosis of osteoporosis in stage 4, 5, or 5D CKD. Patients in stages 4, 5, and 5D CKD have a high prevalence of other metabolic bone diseases and CKD-MBD, which increases difficulties in managing these more complex situations, and the WHO criteria or fragility fractures cannot be used for the diagnosis of osteoporosis in these stages.

That bisphosphonates in advanced CKD may be associated with low bone turnover is uncertain because they decrease bone turnover in a preexisting low bone turnover state, which can influence bone in different ways and result in more or less cardiovascular disease.

Bone biopsy in advanced CKD is a priority consideration before prescribing bisphosphonates in these patients because these medicines will increase the possibility of worsening bone turnover, osteomalacia, and mixed uremic osteodystrophy, and aggravate hyperparathyroidism. In the dialysis population, bisphosphonates should be carefully considered as therapeutic agents and with very specific requirements: carefully defined osteoporosis, dose adjustment considerations, and limited exposure time.

Thus, the bisphosphonates should be used with caution in the CKD patient population and treatment tailored to each individual. Currently, the role of bisphosphonates in BMD and vascular calcification in CKD population remains unclear. It must be stated strongly that most of these compounds are substantially renally excreted and thus tend to accumulate significantly in plasma and bone, with prolonged and often excessive skeletal actions. In patients with early CKD (stages 1–3), nutritional vitamin-D (cholecalciferol or ergocalciferol) and calcium should be provided. However, in patients with advanced CKD or dialysis active vitamin-D (calcitriol or paricalcitol) could be considered. Diagnosis and treatment of low BMD in CKD, and any impact on associated tendencies to vascular calcification, remain challenging and underresearched. Because the current evidence for this population is insufficient, further and detailed research is required to better delineate how to prescribe these medicines appropriately.

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