Assessment and treatment of osteoporosis are recommended following hip fracture. Osteoporosis treatment assumes an adequate calcium intake and a normal vitamin D plasma level. The authors conducted a study in three phases. Phase I: circulating 25-hydroxyvitamin D levels were retrospectively recorded from in the case records of 381 consecutive patients with 387 hip fractures, between March 2010 and September 2011. Only 27 patients had sufficient (> 75 nmol/L) circulating vitamin D, and of these 22 were taking vitamin D supplements. The remainder, 354 patients, had abnormally low vitamin D levels, with a mean value of 26.4 nmol/L. These findings confirmed literature data, and gave rise to the prospective Phase II (October 2011): 14 consecutive patients with a hip fracture received rapid substitution therapy with 50,000 IU cholecalciferol (vitamin D3) daily for 3 days. Patients with corrected calcium level (calcium level based on the serum albumin level) > 2.60 mmol/L were excluded from phase II (and phase III), in order to avoid hypercalcemia. Substitution resulted in an increase in vitamin D plasma levels from +/-29.6 nmol/L to +/-81.4 nmol/L (p < 0.0001), after +/-14 days. However, vitamin D level remained below the desired threshold of 75 nmol/L in 29%. Therefore it was decided to increase the treatment period from 3 days to 7 days in the next 54 patients with a hip fracture in a prospective phase III (October 2011-January 2012). This time rapid substitution resulted in an increase from +/-31.4 nmol/L to +/-131.1 nmol/L (p < 0.0001), after +/-16 days, and 100% of treated patients achieved plasma levels above the desired threshold of 75 nmol/L. Conclusion: virtually all patients with a hip fracture have low vitamin D plasma levels; substitution with 50,000 IU oral cholecalciferol daily for 7 days increases vitamin D plasma levels rapidly, safely and consistently. Keywords: cholecalciferol; ergocalciferol; uploading; rapid substitution; hip fractures; osteoporosis; fragility fracture; calcium; D3.

INTRODUCTION

Hip fractures are a major cause of morbidity and mortality in the elderly, and are associated with chronic pain, reduced mobility, disability, and an increasing degree of dependence (24). These low-energy injuries are costly to treat and have a significant physical and social impact.
An estimated worldwide incidence of hip fractures in the year 2000 was 1.6 million (23). This number is projected to rise up to 2.6 million by 2025, and to more than 4.5 million by 2050 (17). Most hip fractures in the elderly can be related to the development of osteoporosis, a progressive, systemic skeletal disorder characterised by low bone mass and micro-architectural deterioration of bone tissue, with a consequent increase in bone fragility and susceptibility to fracture (28).

It has been shown that identification and management of patients at risk can significantly reduce the risk of a further fracture through modification of lifestyle, adequate treatment of osteoporosis, and a falls prevention programme (13, 27). Managing hip fractures appropriately has been rewarded in the UK by the introduction of Best Practice Tariff, in which the fee paid to NHS hospitals for treating patients with a proximal femoral fracture is increased by approximately 20% if a number of quality targets, including assessing and initiating appropriate osteoporosis treatment are met.

In the last decades, the development of effective therapies (e.g. bisphosphonates, RANK ligand inhibitors, selective estrogen receptor modulators (SERMs) and strontium ranelate) has extended the spectrum of osteoporosis treatment. Most current osteoporosis guidelines assume that patients who receive treatment for secondary prevention of osteoporotic fragility fractures have an adequate calcium intake and are vitamin D replete. Adequate levels of calcium and vitamin D are needed to ensure optimum effects of the treatments for osteoporosis and are recommended in the data sheets of all the commonly used drugs.

Vitamin D status is best evaluated by measuring the circulating 25-hydroxyvitamin D concentration in the serum. Although controversy surrounds the definition of low vitamin D status, there is general acceptance that the optimal circulating 25-hydroxyvitamin D level should be 75 nmol/L or above (3). A threshold for optimal 25-hydroxyvitamin D and hip Bone Mineral Density (BMD) has been established from 13,432 individuals of NHANES III (Third National Health and Nutrition Examination Survey), including both younger (20-49 years) and older (≥ 50 years) individuals with different ethnic racial background (7). This study shows that high serum 25-hydroxyvitamin D levels are associated with higher BMD throughout the reference range of 22.5-94 nmol/L in all subgroups. In younger whites and younger Mexican Americans, higher 25-hydroxyvitamin D was associated with higher BMD even beyond 100 nmol/L. A meta-analysis (9) of 12 double-blind RCTs for non-vertebral fractures (n = 42,279) and eight RCTs for hip fractures (n = 40,886) found that the efficacy of vitamin D in preventing fractures is dose dependent and increases significantly with a higher achieved level of 25-hydroxyvitamin D in the treatment group, starting at 75 nmol/L (6, 9). However, serum 25-hydroxyvitamin D levels above 220 nmol/L have been associated with hypercalcaemia and other signs of toxicity (15, 34).

Bone remodelling is a balance between bone formation by osteoblasts and bone resorption by osteoclasts and mononuclear cells. It involves several hormones (Fig. 1). Indeed, low calcium levels increase parathyroid hormone (PTH) production by the parathyroid glands. This stimulates calcium release from the bones and increases the reabsorption of calcium in the distal renal tube cells. PTH also stimulates the renal production of biologically active vitamin D, which in turn increases calcium absorption from the gut. All these actions increase the calcium level in plasma. High calcium levels on the other hand stimulate the thyroid gland to produce and release calcitonin, which inhibits bone resorption by blocking PTH receptors on the osteoclasts, decreases calcium reabsorption in the kidney, and decreases calcium absorption from the gut. Thus reducing plasma calcium levels.

Vitamin D deficiency may be characterized biochemically by the presence of secondary hyperparathyroidism, which can also contribute to the bone loss in osteopenic patients. Secondary hyperparathyroidism is a physiological response to hypocalcaemia associated with vitamin D deficiency, and treatment with vitamin D will normalise the elevated PTH levels without significantly elevating the serum calcium level. It is important to distinguish secondary hyperparathyroidism from primary hyperparathyroidism due to a parathyroid adenoma, hyperplasia, or malignancy, in which excessive
PTH secretion leads to bone resorption and high calcium levels in the plasma. In primary hyperparathyroidism substitution with vitamin D will not normalise PTH levels, and a potentially dangerous hypercalcaemia may ensue.

In the general population, particularly in the elderly, vitamin D levels are commonly reduced as a result of low dietary intake, decreased sun exposure, decreased intrinsic vitamin D production, and decreased vitamin D receptor activity. There is some evidence that vitamin D deficiency is also becoming widespread in younger patients (33). Other authors have also measured vitamin D levels in patients with a hip fracture, and demonstrated that levels are consistently low. Three UK audits (Belfast, Glasgow and London) identified 694 patients with a hip fracture. Mean levels of 25-hydroxyvitamin D ranged from 24.7 nmol/L to 36.1 nmol/L. Of these patients 91-99% had a level below 80 nmol/L (11). However this does not seem to have been taken on board by trauma surgeons and no guidelines are currently available on how to safely and effectively optimise vitamin D levels in patients with a hip fracture.

In this study, we noted the levels of circulating 25-hydroxyvitamin D in a consecutive series of 381 patients sustaining a hip fracture (phase I, retrospective), and evaluated the effect of rapid substitution therapy with high dose oral vitamin D3 (cholecalciferol) (phase II: 3 days, and phase III: 7 days, both prospective). From these data, guidelines were developed for the management of low vitamin D levels in patients with a hip fracture.

**MATERIALS AND METHODS**

Phase I (retrospective): circulating 25-hydroxyvitamin D and serum calcium levels were recorded from the files of 381 consecutive patients admitted with 387 hip fractures between March 2010 and September 2011. All patients were included.

Vitamin D analysis was performed on serum using a direct competitive chemiluminescence immunoassay (Liaison®, DiaSorin). Levels above 75 nmol/L were defined as sufficient, between 25-75 nmol/L as insufficient, and below 25 nmol/L as deficient. Calcium and corrected calcium (= calcium level based on the serum albumin level) analyses were also performed, using spectrophotometry (Cobas® 6000, Roche Diagnostics Limited, UK).

Phase II and phase III (both prospective) focused on rapid substitution therapy, respectively during 3 and 7 days. Inclusion criteria: all patients presenting with a hip fracture, with corrected calcium below 2.60 mmol/L (to avoid hypercalcaemia), who agreed with a written informed consent. Exclusion criteria: patients with corrected calcium above 2.60 mmol/L, because treatment
could potentially lead to dangerously high calcium levels; they were assessed biochemically for primary hyperparathyroidism. Prior intake of a low dose of vitamin D and a calcium supplement was not an exclusion criterion. Patients with cognitive impairment resulting in non-compliance with therapy, and patients with a significantly reduced life expectancy (ASA grade 5) or significant renal impairment (GFR < 30 mL/min/1.73m²) were also excluded. Clinical side effects of hypercalcaemia and hypervitaminosis D, such as nausea, vomiting, constipation, thirst and polyuria were recorded.

Phase II (prospective): rapid substitution therapy with 50,000 IU oral vitamin D3 (cholecalciferol) (SunVit-D³®) daily for 3 days was started in 14 consecutive patients with a hip fracture (October 2011).

Phase III (prospective): rapid substitution therapy with 50,000 IU oral vitamin D3 (cholecalciferol) (SunVit-D³®) daily for 7 days was started in 54 consecutive patients with a hip fracture, between October 2011 and January 2012. Initially 60 patients had been considered, but 6 were excluded (2 non-compliant, 2 absent repeat sample, 2 initial corrected calcium above 2.60 mmol/L).

Both patient groups (phase II and phase III) had similar demographic characteristics, and did not differ in terms of age-related comorbidities, drug intake or dwelling status. Repeat vitamin D and calcium measurements were performed between 7 and 42 days after the start of substitution therapy. After completion of the study all patients were recommended to take a low dose vitamin D and calcium supplement (Adcal D³®) for life: calcium carbonate (1500 mg) and vitamin D3 (400 IU).

RESULTS

Phase I: over an 18 month period 381 patients were admitted with 387 hip fractures (95 men, 286 women) (mean age 83 years, range 34-97 years). Serum 25-hydroxyvitamin D analysis was performed in all patients (Fig. 2). Only 27 patients had a sufficient (> 75 nmol/L) circulating vitamin D level (mean 91.2 nmol/L, SD 20.0 nmol/L, range 75.6-171 nmol/L), and of these 22 were taking vitamin D supplements. The remainder, 354 patients, had abnormally low vitamin D levels, with a mean value of 26.4 nmol/L (SD = 17.9 nmol/L, range < 10-74.4 nmol/L): 155 patients (44%) were vitamin D insufficient with a level between 25 and 75 nmol/L, 199 patients (56%) were vitamin D deficient with a level below 25 nmol/L. Of these patients, 43 had a low vitamin D level despite taking low dose vitamin D (400 or 800 IU) and calcium supplements with a mean level of 57.8 nmol/L (SD = 10.3 nmol/L, range 32.5-74.4 nmol/L).

Fig. 2. — Phase I: vitamin D levels (nmol/L) in 381 patients (387 hips). Only 27 patients had a level above 75 nmol/L, 22 of whom were taking supplements. The remainder, 354 patients, had abnormally low vitamin D levels (below 75 nmol/L), with a mean value of 26.4 nmol/L: 155 patients (44%) were vitamin D insufficient with a level between 25 and 75 nmol/L, 199 patients (56%) were vitamin D deficient with a level below 25 nmol/L. Of these 354 patients, 43 had a low vitamin D level (+/- 57.8 nmol/L) despite taking low dose vitamin D and calcium supplements (400 or 800 IU).
Table I. Vitamin D, calcium and corrected calcium before and after substitution with 50,000 IU oral cholecalciferol (vitamin D3) for 3 days (phase II : group 1) and 7 days (phase III : group 2)

<table>
<thead>
<tr>
<th>Group 1</th>
<th>Before substitution</th>
<th>After substitution</th>
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<tr>
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<tr>
<td>Vitamin D (nmol/L)</td>
<td>29.6 (SD = 15.8, range &lt;10-56.8)</td>
<td>81.4 (SD = 17.0, range 47.4-108.0, p &lt; 0.0001)</td>
</tr>
<tr>
<td>Calcium (mmol/L)</td>
<td>2.12 (SD = 0.13, range 1.91-2.39)</td>
<td>2.30 (SD = 0.12, range 2.12-2.54, p &lt; 0.0001)</td>
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<tr>
<td>Corrected calcium (mmol/L)</td>
<td>2.43 (SD = 0.04, range 2.33-2.47)</td>
<td>2.55 (SD = 0.10, range 2.38-2.73, p &lt; 0.0001)</td>
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<tr>
<th>Group 2</th>
<th>Before substitution</th>
<th>After substitution</th>
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<tbody>
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<tr>
<td>Vitamin D (nmol/L)</td>
<td>31.4 (SD = 24.2, range &lt;10-113.0)</td>
<td>131.1 (SD = 30.7, range 85.6-243.0, p &lt; 0.0001)</td>
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<tr>
<td>Calcium (mmol/L)</td>
<td>2.19 (SD = 0.13, range 1.94-2.50)</td>
<td>2.33 (SD = 0.17, range 1.84-2.78, p &lt; 0.0001)</td>
</tr>
<tr>
<td>Corrected calcium (mmol/L)</td>
<td>2.47 (SD = 0.09, range 2.21-2.60)</td>
<td>2.54 (SD = 0.12, range 2.18-2.81, p &lt; 0.0002)</td>
</tr>
</tbody>
</table>

Group 1 (phase II ; n = 14) : substitution with 50,000 IU oral vitamin D3 (cholecalciferol) daily for 3 days. Significant increase in circulating 25-hydroxyvitamin D levels and normalizing of calcium levels 14 days (range, 7-23 days) after start of substitution; 29% of patients do not reach desired threshold of 75 nmol/L.

Group 2 (phase III ; n = 54) : substitution with 50,000 IU oral vitamin D3 (cholecalciferol) daily for 7 days. Significant increase in circulating 25-hydroxyvitamin D levels and normalizing of calcium levels 16 days (range, 7-42 days) after start of substitution. All patients achieve the desired vitamin D threshold of 75 nmol/L.

DISCUSSION

Vitamin D is responsible for the absorption of calcium from the intestine, and may have other bone-protective effects independent of those on calcium metabolism (14,32). In addition, vitamin D plays an important role in muscular function: higher levels of vitamin D are associated with increased muscular strength and balance, reducing the risk of falls and subsequent fractures (5,12). In several double blind randomised controlled trials, vitamin D supplementation increased muscle strength and balance, and reduced the risk of falling (4,8,10,29,30). A study by Glerup et al (16) suggests that vitamin D deficiency may cause muscular impairment even before adverse effects on bone occur.
Hip fracture correlated with vitamin D deficiency

This study clearly confirms that the vast majority of patients with hip fracture are vitamin D deficient and require supplements. The current target for 25-hydroxyvitamin D serum concentration in secondary fracture prevention is 75 nmol/L. The fact that the majority of the patients taking low dose supplements were also found to be vitamin D deficient raises the question as to whether the current dose recommended for supplementation is adequate. This question has also been raised by others (2,19). According to studies in younger adults, intakes of as high as 4,000-10,000 IU are safe (18,35). Furthermore, other studies indicate that individuals with a low starting level may need a high dose of vitamin D to achieve desirable vitamin D levels (18,20).

Vitamin D3 preferable to vitamin D2

The aim of the current study was to examine the effect of rapid vitamin D substitution with vitamin D3 (cholecalciferol). Pappasmonnou et al (26) studied in 2011 the effect of a loading dose of vitamin D2 (ergocalciferol) and found no long-term advantage over daily supplementation (26). However, it has been shown that, although both vitamin D2 and D3 produce similar rises in serum 25-hydroxyvitamin D concentration over 3 days, levels decline much more rapidly in subjects treated with D2 (1). This may possibly reflect lower affinity of vitamin D2 for vitamin D binding protein in the circulation, leading to more rapid clearance. As such, use of supplements containing vitamin D3, rather than vitamin D2, is generally recommended (22).

The current study demonstrates that, in patients with abnormal corrected calcium levels (< 2.60 mmol/L), rapid high dose substitution with 50,000 IU of vitamin D3 can normalise circulating serum vitamin D levels quickly and safely. This is important because all drugs used in the management of osteoporosis require normal vitamin D levels before they are started. However, the authors suspect that the majority of patients started on such treatment do not have their vitamin D levels checked systematically and are almost all deficient. This was certainly the case in the authors’ institution before they undertook this study. This means that rapid substitution with high doses vitamin D3 makes sense after hip fractures. Interestingly, also calcium levels increase from an abnormal low value to within the normal range.

Vitamin D protects against falls and non-vertebral fractures

In addition, the fact that fall prevention and non-vertebral fracture prevention increased significantly with higher achieved 25-hydroxyvitamin D levels in two meta-analyses (8) of double-blind RCTs, is in itself a good reason to normalise vitamin D as soon as possible after admission with a fracture. Fall prevention was effective with 25-hydroxyvitamin D levels of 60 nmol/L up to 95 nmol/L, while 75-112 nmol/L were required for non-vertebral fracture prevention (9,25). Whether higher levels of vitamin D give even more protection against falls and fracture is not known, but has been postulated (6).

Routine high dose substitution after hip fractures is cost saving

Sixty-five out of 381 fracture patients (Fig. 2) took low dose vitamin D and calcium supplements, but only 22 (34%) of these had a normal vitamin D level. The remaining 316 patients did not take supplements, and only 5 (2%) of these had a normal vitamin D level. The authors feel that this allows to assume that all patients with a hip fracture, on supplements or not, have low vitamin D levels and should be treated accordingly. This would obviate the need for a routine vitamin D analysis, which saves approximately 7.5€ per patient, and would allow even earlier initiation of a high dose rapid substitution. At unit level this is a small saving but if it became standard practice for all hip fracture patients the saving would be considerable. The cost of treating one patient with 50,000 IU oral vitamin D a day for seven days is only 2€. In this study patients were included, regardless of their prior vitamin D intake, and rapid substitution was found to be effective and safe in all.

Acta Orthopaedica Belgica, Vol. 79 - 5 - 2013
Limitations

Phase II and III: patients were not randomised, but were treated sequentially. Both phase II and phase III groups were relatively small (14 and 54 patients), but the dramatic rise in vitamin D levels ($p < 0.0001$) makes the likelihood of the effects being fortuitous improbable. More research is needed to establish if the effect of a short course of high dose oral vitamin D3 substitution is maintained if followed by long term low dose therapy or if an occasional booster is required.

Of more concern is the possibility that a patient with primary hyperparathyroidism might get high dose vitamin D supplementation causing severe hypercalcaemia and the complications thereof. For this reason only patients with uncompromised renal function (GFR > 30 mL/min/1.73 m$^3$) and a calcaemia < 2.60 mmol/L received such a large dose. Patients who do not fit these criteria should be treated with much more caution. An algorithm for the management of all patients is shown (Fig. 3). The authors’ current practice for patients who present with a corrected calcium of 2.60-2.80 mmol/L is to measure the PTH level and renal function, and start a weekly dose of 50,000 units vitamin D3 and regular calcium monitoring. In the majority of cases initial high PTH levels are due to secondary hyperparathyroidism and at six weeks their repeat PTH level and calcium levels will have returned to normal. They can then receive additional vitamin D supplementation as required before remaining on a lifelong low dose vitamin D3 and calcium regime. Patients who present with corrected calcium > 2.80 mmol/L and patients with corrected calcium between 2.60-2.80 mmol/L, whose PTH does not fall after a week of rapid vitamin D substitution, are referred to an endocrinologist for further investigation.

CONCLUSION

The benefits of maintaining adequate vitamin D levels in hip fracture patients have been established and the very low cost of the treatment makes it highly cost effective. The very low levels of vitamin D in this patient group took us by surprise, and further studies to evaluate levels in younger populations are clearly indicated. There have been recent reports of the re-emergence of vitamin D deficient related rickets in children in the UK (31), and it has been postulated that the national obsession with preventing sunburn because of the risk of skin cancer in later life, coupled with less time spent outdoors by

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**Fig. 3.** — Algorithm for the management of low vitamin D levels in hip fractures (see full text).

CC = corrected calcium (mmol/L); EGFR = estimated glomerular filtration rate (mL/min/1.73m$^2$); PTH = parathyroid hormone.
the majority of the population, has had the unintended consequence of creating an entire population which is vitamin D deficient (21). The authors believe that the management of many other types of osteoporotic fractures faces similar challenges with regard to vitamin D supplementation, although this needs to be explored. Investigating this should be a major public health priority, particularly because treatment would be easy and inexpensive.

Acknowledgement

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