ORIGINAL ARTICLE

Oral vitamin D replacement is effective in chronic liver disease

Fréquence du déficit en vitamine D et effet de la supplémentation orale au cours des maladies chroniques du foie

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Summary

Background & aims. — End-stage chronic liver disease is associated with vitamin D deficiency but the prevalence across a broad-spectrum of liver disease is unknown. This study prospectively examines prevalence of vitamin D deficiency and response to replacement in chronic liver disease.

Methods. — One hundred and fifty-eight outpatients with chronic liver disease were enrolled. Serum 25-hydroxyvitamin D (25\(^{\text{OH}}\)D) levels were classified as: severely deficient less than 25 nmol/l, deficient 25—54 nmol/l or replete greater than 54 nmol/l. Sixty-five of 158 (41%) had cirrhosis.

Results. — 25\(^{\text{OH}}\)D was suboptimal in 101/158 (64%), including severe deficiency in 24 patients (15%). Vitamin D deficiency occurred in liver disease of all aetiologies, including patients with only mild liver disease. 25\(^{\text{OH}}\)D increased by 60.0% (19.11 ± 13.20 nmol/l) in patients with deficiency after vitamin D replacement and decreased by 25.2% (-18.33 ± 12.02 nmol/l) in non-treated initially replete patients over a median of 4 months.

Conclusions. — Vitamin D deficiency improves with oral vitamin D supplementation and levels fall without supplementation. Chronic liver disease patients are at very high risk of vitamin D deficiency regardless of etiology or severity.

Vitamin D deficiency is common in chronic disease, but the prevalence across the broad spectrum of chronic liver disease (CLD) is not well documented. The importance of vitamin D deficiency is highlighted by the increasingly recognized role vitamin D has in immunoregulation [1], cellular growth and differentiation, antioxidation, and regulation of fibroblast proliferation and collagen synthesis [2].

Previous studies have focussed on vitamin D deficiency in cholestatic liver disease, especially primary biliary cirrhosis [3], and in patients with advanced cirrhosis [4]. Although
Oral vitamin D replacement is effective in chronic liver disease. Vitamin D deficiency is well accepted as a consequence of chronic liver disease (CLD), no prospective studies have determined the response to standard vitamin D replacement therapy. We sought to examine prospectively the prevalence of vitamin D deficiency in chronic liver disease of different etiologies and stages, evaluate if serum 25-hydroxyvitamin D (25(OH)D) levels correlate with severity of liver disease and determine the response to standard oral replacement therapy.

Patients and methods

Patients attending the outpatient liver clinic between November 2006 and July 2007, during a period of the year when cumulative sunlight exposure is higher, were enrolled prospectively (n = 158) with etiologies detailed in Table 1. Sixty-five of 158 (41%) had cirrhosis and 82 (52%) were men. The majority of the patients were of Anglo-Saxon, European or Asian descent. The Melbourne Health Human Research Ethics Committee approved the study.

Laboratory analysis, treatment and statistical analysis

25(OH)D were measured by radioimmunoassay (Diasorin, Stillwater, MN, USA). Consistent with our laboratory’s ranges and the literature, 25(OH)D was classified as: severely deficient (< 25 nmol/l), mildly deficient (25—54 nmol/l) or replete (> 54 nmol/l) [5].

All subjects had measurements of serum albumin, bilirubin, aminotransferases, alkaline phosphatase and gamma-glutamyl transferase. Cirrhosis was diagnosed on liver biopsy or a combination of appropriate radiological findings and clinical diagnosis.

Patients with suboptimal 25(OH)D were treated with standard oral replacement doses of ergocalciferol (vitamin D2 1000IU) or cholecalciferol (vitamin D3 50 mcg) supplements daily.

One-way analysis of variance and regression analyses were performed, and scatter plots were generated using Minitab 15 (Minitab Inc. [2007] Pennsylvania). Validity checks of statistical assumptions were satisfactory.

Results

Suboptimal 25(OH)D levels were present in 101 patients (64%) and 24 (15%) had severe vitamin D deficiency (Table 1). There was no significant difference in 25(OH)D between males and females. Patients with cirrhosis were more likely to be deficient (75%) (P = 0.028) and were over-represented in the severely deficient category (63%). Vitamin D deficiency was seen in liver disease across all etiologies including in subgroups with a low incidence of cirrhosis. Surprisingly, no patients with cholestatic liver disease were severely deficient of 25(OH)D, with 40% recording replete levels.

Patients with initially replete 25(OH)D without supplementation demonstrated an average 25.2% decrease (−18.33 ± 12.02 nmol/l, 72.72 vs. 54.39 nmol/l) over a median period of 4 months. Oral vitamin D supplementation resulted in a 41.0% increase in mean 25(OH)D in patients with mild deficiency (16.38 ± 12.74 nmol/l, 39.92 vs. 56.30 nmol/l) and a 141.1% increase in patients with severe deficiency (22.23 ± 14.12 nmol/l, 15.75 vs. 37.98 nmol/l). Overall, for patients with vitamin D deficiency there was a 60% increase in mean 25(OH)D (19.11 ± 13.20 nmol/l, 31.86 vs. 50.97 nmol/l) over a median period of 4 months.

25(OH)D levels correlated with serum albumin (r = 0.230, P = 0.006) and inversely with bilirubin (r = −0.240, P = 0.004) as surrogate markers of liver disease severity and with the presence of cirrhosis (r = −0.175, P = 0.028).

Discussion

Although previous investigators have noted vitamin D deficiency as a complication of CLD, this is the first study to...
Vitamin D levels can be improved by low dose oral supplementation and may have a negative impact on the natural history of chronic liver disease patients regardless of severity, how- ever, the degree of deficiency correlates with liver disease severity. This predisposes patients to hepatic osteodystrophy and may have a negative impact on the natural history. Vitamin D levels can be improved by low dose oral supplementation and serum vitamin D levels continue to fall in patients without supplementation. Further research to determine optimal replacement doses in chronic liver disease patients are required.

Conflict of interest statement

No potential conflict of interest relevant to this article was reported.

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References

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