Treatment of osteomalacia associated with primary biliary cirrhosis with parenteral vitamin D$_2$ or oral 25-hydroxyvitamin D$_3$

JULIET E. COMPSTON, L. W. L. HORTON, AND R. P. H. THOMPSON

From the Gastrointestinal Research Unit, Rayne Institute, and Department of Surgical Pathology, St Thomas’ Hospital, London

SUMMARY The histological and biochemical response of osteomalacia has been studied in four patients with primary biliary cirrhosis, who were treated with oral 25-hydroxyvitamin D$_3$, 50 µg daily, or intramuscular vitamin D$_2$, 150 000 units once weekly, for five to 12 months. All patients showed complete histological healing of osteomalacia, despite rapidly deteriorating liver function in three. Plasma 25-hydroxyvitamin D concentrations were low in all patients before treatment, but became normal during either vitamin therapy. Serum calcium and phosphate levels, and urinary calcium excretion were not always reliable in predicting the histological response to treatment. Serum alkaline phosphatase activity decreased in all patients during vitamin D therapy. We conclude that both high-dose parenteral vitamin D$_2$ and oral 25-hydroxyvitamin D$_3$ may be effective in healing osteomalacia associated with primary biliary cirrhosis. Measurement of plasma 25-hydroxyvitamin D levels during vitamin D therapy provides useful information about 25-hydroxylation of the parent vitamin and intestinal absorption of orally administered 25-hydroxyvitamin D$_3$.

Vitamin D is 25-hydroxylated in the liver to 25-hydroxyvitamin D (25-OHD) (Ponchon et al., 1969), which is further hydroxylated in the kidney (Fraser and Kodicek, 1970) to 1,25-dihydroxyvitamin D (1,25-(OH)$_2$D) (Holick et al., 1971), the major active metabolite. Plasma concentrations of 25-OHD are reduced in most patients with primary biliary cirrhosis (PBC) (Long et al., 1976; Wagonfeld et al., 1976); possible causes include malabsorption of dietary vitamin D (Thompson et al., 1966; Krawitt et al., 1977), and of 25-OHD excreted into the bile and undergoing enterohepatic circulation (Arnaud et al., 1975; Compston and Thompson, 1977), and increased urinary loss of vitamin D (Krawitt et al., 1977). Hepatic 25-hydroxylation of vitamin D may also be impaired in PBC (Wagonfeld et al., 1976), although most patients achieve normal plasma 25-OHD concentrations if given sufficient precursor vitamin D (Skinner et al., 1977).

Osteomalacia develops in some patients with PBC (Compston and Thompson, 1977; Long et al., 1978a) and large parenteral doses of vitamin D are usually given for its prophylaxis or treatment. However, very few studies of bone histology have been carried out after treatment, and the therapeutic dose range has not been strictly defined, nor is it certain that parenteral administration of the vitamin is necessary. In this study we report the biochemical and histological response in four patients with symptomatic PBC and osteomalacia to treatment with either parenteral vitamin D$_2$ or oral 25-OHD$_3$.

Methods

Patients

The four patients, three female and one male, aged 50–66 years (mean 60 years) with symptomatic PBC and histological osteomalacia have been described elsewhere (Compston and Thompson, 1977). Patients 1 and 2 were treated with oral 25-OHD$_3$, 50 µg/day, and patients 3 and 4 with intramuscular vitamin D$_2$, 150 000 units once weekly (Table 1). All except patient 3 were taking cholestyramine (4 g twice or thrice daily).

Control values for bone histology were obtained from post-mortem biopsies of seven females and
Plasma 25-OHD levels were measured by a competitive protein-binding assay (Edelstein et al., 1974) using normal human serum as binding protein. Serum calcium, phosphate, alkaline phosphatase, bilirubin, and albumin were measured on an SMA 12/60 autoanalyzer, and serum calcium concentrations were corrected for the serum albumin level (Berry et al., 1973). Urinary calcium was measured on an Eppendorf flame photometer.

Results

Bone histology (Table 1)
All patients responded well to vitamin D therapy, with a reduction in osteoid amount and increase in calcification fronts to normal values.

Plasma 25-OHD (Table 2)
Plasma 25-OHD concentrations were below normal in all patients before vitamin D therapy. After five to 12 months' treatment all had levels within the normal range.

Serum calcium and phosphate (Table 2)
Serum calcium and phosphate levels were higher after vitamin D therapy in patients 1, 3, and 4, but, in patient 2, levels of calcium and phosphate fell during therapy, the phosphate to less than normal. After correction for her much reduced serum albumin, serum calcium after treatment was mildly raised in patient 1.

Twenty-four hour urinary calcium (Table 2)
Twenty-four hour urinary calcium excretion was below normal in all patients before treatment. The values after treatment, measured in patients 2, 3, and 4, were normal in patients 2 and 4, but remained low in patient 3.

Serum alkaline phosphatase and bilirubin (Table 3)
After treatment with vitamin D, serum alkaline phosphatase had fallen in all patients, the greatest change occurring in patients 1 and 3. Serum bilirubin increased in all patients during the treatment period.

Discussion

This study has demonstrated that osteomalacia associated with symptomatic PBC may respond to oral or parenteral vitamin D, and that, even in the presence of severe liver disease, 25-hydroxylation of the parent vitamin, or intestinal absorption of orally administered 25-OHD₃, can be adequate to produce normal plasma 25-OHD levels and to promote bone healing, despite the continued cholestyramine therapy in three patients.

The value of histological examination of bone in the diagnosis of osteomalacia has recently been emphasised (Compston et al, 1978), and our findings show that it is also useful in assessing the response to treatment, as biochemical measurements may be misleading. Thus, in patient 2, serum calcium and phosphate levels were lower after vitamin D therapy than before, and urinary calcium excretion remained low in patient 3 after bone healing had occurred.

Table 1  Details of vitamin D therapy and quantitative bone histology before and after treatment

<table>
<thead>
<tr>
<th>Patient</th>
<th>Treatment</th>
<th>Duration of treatment (months)</th>
<th>Osteoid volume % total cancellous volume</th>
<th>Osteoid surface % total cancellous surface</th>
<th>Calcification fronts % osteoid surface</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Before</td>
<td>After</td>
<td>Before</td>
</tr>
<tr>
<td>1</td>
<td>25-OHD₃*</td>
<td>5</td>
<td>21.3</td>
<td>2.7</td>
<td>64.3</td>
</tr>
<tr>
<td>2</td>
<td>25-OHD₃*</td>
<td>12</td>
<td>14.3</td>
<td>0</td>
<td>78.1</td>
</tr>
<tr>
<td>3</td>
<td>Vitamin D₃†</td>
<td>6</td>
<td>12.3</td>
<td>0.5</td>
<td>24.8</td>
</tr>
<tr>
<td>4</td>
<td>Vitamin D₄†</td>
<td>13</td>
<td>8.9</td>
<td>2.0</td>
<td>18.1</td>
</tr>
<tr>
<td>Normal mean ± SD</td>
<td></td>
<td>2.9 ± 1.7</td>
<td>17.9 ± 10.9</td>
<td>63.3 ± 4.5</td>
<td></td>
</tr>
</tbody>
</table>

* By mouth.
† Intramuscularly.
‡ Bordier et al. (1975).
Treatment of osteomalacia associated with primary biliary cirrhosis

Table 2  Serum calcium and phosphate, plasma 25-OHD, and 24-hour urinary calcium before and after vitamin D therapy

<table>
<thead>
<tr>
<th>Patient</th>
<th>Corrected serum calcium (mmol/l)</th>
<th>Serum phosphate (mmol/l)</th>
<th>Plasma 25-OHD (nmol/l)</th>
<th>Urinary calcium (mmol/24 hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Before</td>
<td>After</td>
<td>Before</td>
<td>After</td>
</tr>
<tr>
<td>1</td>
<td>2.25</td>
<td>2.73</td>
<td>0.8</td>
<td>1.3</td>
</tr>
<tr>
<td>2</td>
<td>2.45</td>
<td>2.30</td>
<td>1.0</td>
<td>1.2</td>
</tr>
<tr>
<td>3</td>
<td>2.32</td>
<td>2.48</td>
<td>1.0</td>
<td>1.5</td>
</tr>
<tr>
<td>4</td>
<td>2.32</td>
<td>2.54</td>
<td>1.0</td>
<td>1.5</td>
</tr>
<tr>
<td>Normal</td>
<td>2.25-2.55</td>
<td></td>
<td>0.8-1.4</td>
<td></td>
</tr>
</tbody>
</table>

Table 3  Serum bilirubin, albumin, and alkaline phosphatase before and after vitamin D therapy

<table>
<thead>
<tr>
<th>Patient</th>
<th>Serum bilirubin (μmol/l)</th>
<th>Serum albumin (g/l)</th>
<th>Serum alkaline phosphatase (KAl/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Before</td>
<td>After</td>
<td>Before</td>
</tr>
<tr>
<td>1</td>
<td>28</td>
<td>460</td>
<td>36</td>
</tr>
<tr>
<td>2</td>
<td>24</td>
<td>30</td>
<td>40</td>
</tr>
<tr>
<td>3</td>
<td>216</td>
<td>400</td>
<td>41</td>
</tr>
<tr>
<td>4</td>
<td>58</td>
<td>164</td>
<td>36</td>
</tr>
<tr>
<td>Normal</td>
<td>5-20</td>
<td>38-56</td>
<td>4-13</td>
</tr>
</tbody>
</table>

However, the serum alkaline phosphatase concentration fell in all patients during treatment, especially in patients 1 and 3, despite deterioration of liver function, and although isoenzymes were not measured, this decrease in total alkaline phosphatase activity was probably due to a fall in the bone isoenzyme concentration. The normal plasma 25-OHD levels achieved by all patients during vitamin D therapy suggested that the dose administered was in the therapeutic range, although others have found that patients with normal or high-normal plasma 25-OHD concentrations may still have osteomalacia after many months’ treatment (Long et al., 1978a).

The prognosis of symptomatic PBC is poor (Sherlock, 1959), and in patients with advanced disease the diagnosis of osteomalacia may not be clinically important. None of our patients had severe bone pain or fractures, and two (1 and 4) have already died from their liver disease. However, three years after diagnosis patient 2 still has relatively mild, anicteric PBC, and treatment of her osteomalacia may well have prevented the onset of symptoms.

Recently there has been much interest in the therapeutic applications of vitamin D metabolites and their analogues. Parenteral 1,25-(OH)₂D₃ was claimed to heal osteomalacia in four patients with PBC (Long et al., 1978b), although calcification fronts were not examined, and these patients had been previously treated with parenteral vitamin D. Five patients with PBC and osteomalacia were treated with oral 25-OHD₃ (Reed et al., 1977) and all showed a reduction in osteoid surface, although calcification fronts were again not reported. Wagonfeld et al. (1976) found that large oral doses (100-200 μg/day) of 25-OHDS improved or stabilised bone mineral content in six of seven patients with PBC, whereas oral or subcutaneous vitamin D₂ was ineffective, but bone biopsy specimens were not examined, and the apparent lack of effect of vitamin D may have been due in some patients to the small doses given, and in others to the subcutaneous route of administration.

We conclude that both parenteral vitamin D₂ and oral 25-OHDS in the doses used in this study, may be effective in healing osteomalacia associated with primary biliary cirrhosis.

We thank J. Sainsbury Ltd and the Special Trustees, St Thomas’ Hospital, for generous financial support, Mr Adrian Webb for technical assistance, and Dr Norman Eve, Roussel Laboratories, for supplies of 25-OHDS.

References


Bordier, P. J., Marie, P., Arnaud, C. D., Gueris, J., Ferriere, C., and Norman, A. W. (1975). Early effects of vitamin D₃ and some analogues—25-OHDS, 1,25(OH)₂D₃, 1αOHDS₃—upon resorption, formation and mineralisation of bone in nutritional or malabsorption osteomalacia—with reference to serum 1.PTH, calcium, phosphate and alkaline phos-


Treatment of osteomalacia associated with primary biliary cirrhosis with parenteral vitamin D2 or oral 25-hydroxyvitamin D3.

J E Compston, L W Horton and R P Thompson

*Gut* 1979 20: 133-136
doi: 10.1136/gut.20.2.133

Updated information and services can be found at:
[http://gut.bmj.com/content/20/2/133](http://gut.bmj.com/content/20/2/133)

**Email alerting service**

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Topic Collections**

Articles on similar topics can be found in the following collections:

Pancreas and biliary tract (1949)

**Notes**

To request permissions go to:
[http://group.bmj.com/group/rights-licensing/permissions](http://group.bmj.com/group/rights-licensing/permissions)

To order reprints go to:
[http://journals.bmj.com/cgi/reprintform](http://journals.bmj.com/cgi/reprintform)

To subscribe to BMJ go to:
[http://group.bmj.com/subscribe/](http://group.bmj.com/subscribe/)