Plasma appearance and disappearance of an oral dose of 25-hydroxyvitamin D2 in healthy adults

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Abstract

25-Hydroxyvitamin D (25(OH)D) half-life is a potential biomarker for investigating vitamin D metabolism and requirements. We performed a pilot study to assess the approach and practical feasibility of measuring 25(OH)D half-life after an oral dose. A total of twelve healthy Gambian men aged 18–23 years were divided into two groups to investigate the rate and timing of (1) absorption and (2) plasma disappearance after an 80 nmol oral dose of 25(OH)D2. Fasting blood samples were collected at baseline and, in the first group, every 2 h post-dose for 12 h, at 24 h, 48 h and on day 15. In the second group, fasting blood samples were collected on days 3, 4, 5, 6, 9, 12, 15, 18 and 21. Urine was collected for 2 h after the first morning void at baseline and on day 15. 25(OH)D2 plasma concentration was measured by ultra-performance liquid chromatography-tandem MS/MS and corrected for baseline. Biomarkers of vitamin D, Ca and P metabolism were measured at baseline and on day 15. The peak plasma concentration of 25(OH)D2 was 9.6 (SD 0.9) nmol/l at 4.4 (SD 1.8) h. The terminal slope of 25(OH)D2 disappearance was identified to commence from day 6. The terminal half-life of plasma 25(OH)D2 was 13.4 (SD 2.7) d. There were no significant differences in plasma 25(OH)D2, total 1,25(OH)2D, parathyroid hormone, P, Ca and ionised Ca and urinary Ca and P between baseline and day 15 and between the two groups. The present study provides data on the plasma response to oral 25(OH)D2 that will underpin and contribute to the further development of studies to investigate 25(OH)D half-life.

Key words: 25-Hydroxyvitamin D: Absorption: Half-life: Gambia

Vitamin D is required for skeletal health and Ca homeostasis. Increasing evidence suggests a role for vitamin D in other aspects of health and disease prevention(1,2). Biomarkers link health or disease outcomes to nutrient intake or supply and provide a means to assess nutrient status at the individual or population level(1–3). The most frequently used biomarker and provide a means to assess nutrient status at the individual

**Abbreviations:** 25(OH)D, 25-hydroxyvitamin D; DBP, vitamin D-binding protein; iCa, ionised calcium; LC, liquid chromatography; PTH, parathyroid hormone; uCa, urinary calcium; uCr, urinary creatinine; uP, urinary phosphate.

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half-life for 25(OH)D₃ (10–40 d)⁷⁻¹² is in part determined by the molar excess of vitamin D-binding protein (DBP) and strong binding affinity to DBP that protects 25(OH)D from degradation and provides a circulating store of vitamin D. Vitamin D metabolites with lower binding affinities for DBP have shorter half-lives.¹³,¹⁴ A number of experimental factors may affect the usefulness of methods to determine half-life, including specificity of measurement, size of the tracer dose, rapid mixing with endogenous pool and correctly defined terminal slope. In some studies, the 25(OH)D₃ half-life may have been calculated on the basis of data that included time points that represented not only the terminal slope but also the distribution phase.¹²,¹³,¹⁴ A number of experimental factors may affect the usefulness of methods to determine half-life, including specificity of measurement, size of the tracer dose, rapid mixing with endogenous pool and correctly defined terminal slope. In some studies, the 25(OH)D₃ half-life may have been calculated on the basis of data that included time points that represented not only the terminal slope but also the distribution phase.¹²,¹³,¹⁴,¹⁶,¹⁷ In other studies, tracer molecules were used with labelled atoms in metabolically active positions that can affect 25(OH)D and/or 1,25(OH)₂D metabolism in vitro and in vivo.¹⁶,¹⁷

Alternatives to radio labelled tracers are to use either stable isotope or other suitable tracers with similar metabolism, together with modern mass spectrometric methods. Here, we report the use of 25(OH)D₂ as a tracer, together with a sensitive liquid chromatography (LC)-tandem MS method, in a pilot study to investigate vitamin D metabolism. We aimed to characterise the absorption, distribution and terminal slope of the plasma response to an oral dose of 25(OH)D₂, a readily available and cost-effective tracer. In addition, the effect of the dose on vitamin D, Ca and P metabolism was evaluated.

The study was performed at MRC Keneba, a field station in rural Gambia at latitude 13°N, where there is no or only limited seasonality in vitamin D status due to the year-round UVB exposure from the sun,¹⁹ and thus subjects are expected to be in a steady state for vitamin D status for the duration of the study. Furthermore, the dietary intake of vitamin D₂ was expected to be absent or very low in this population. We used 25(OH)D₂ rather than the parent compound, vitamin D₂, in order to minimise inter-individual differences in absorption, tissue distribution and hydroxylation.

Dose preparation

25(OH)D₂ (product number: 17 937; purity ≥ 98%; Sigma-Aldrich, Poole, UK) was weighed and dissolved in olive oil. The solution was protected from light and mixed on a rotary mixer at room temperature. Aliquots were frozen at −20°C and were used within 2 weeks of preparation. Each dose consisted of 80 nmol (33 µg) 25(OH)D₂ in 810 µl olive oil.

Study protocols

The volunteers were divided into two groups, the ‘absorption’ group (n 6) and the ‘half-life’ group (n 6), with different sample timings but otherwise identical protocols. The size of the dose and sampling protocols were chosen based on data from the few previous estimates of absorption, kinetics and turnover that used unlabelled and radiolabelled compounds in human subjects and animals, combined with compartmental modelling.²⁰⁻¹¹,²⁰⁻²³ Biochemical markers of vitamin D, Ca and P metabolism and liver and kidney function were measured in fasting blood and urine samples collected at baseline (day 0) and on day 15 after the dose. After voiding of the first morning urine, a 2 h fasting urine was collected from approximately 07:00 hours. The subject was asked to drink 500 ml of water at both 30 min and 1.5 h in the 2 h period (total water intake, 1000 ml). Blood was collected 1 h into the urine collection. On day 1 (within 5 d of the baseline sample), the 25(OH)D₂ dose was administered in the morning at approximately 07.30 hours after an overnight fast. The dose was removed from the freezer, protected from light and brought to room temperature on a rotary mixer. Each dose was pipetted onto a small piece of local white bread. The volunteer ate the bread under supervision followed by a standardised meal of bread with mayonnaise (Armanti brand; NGM International Ltd, Rotterdam, The Netherlands). Water was provided ad libitum. Subjects were supervised for 5 h following the dose and were allowed to sit or move about freely within the field station but were asked to refrain from lying down or exercise during this period to minimise the influence of gastric emptying. After 5 h, a meal was provided. Blood samples in the first group were collected from an in-dwelling cannula inserted into a forearm or by venepuncture directly before and every 2 h after administration of the dose for a total of 12 h. After the blood sample at 12 h, a meal was provided and the volunteer was free to leave. Further fasting blood samples were collected after 24 and 48 h and on day 15 following the dose. For individuals in the second group, fasting blood samples were collected by venepuncture at approximately 08.00 hours on day 3 (corresponding to the 48 h sample in the absorption group), 4, 5, 6, 9, 12, 18 and 21.

Sample collections and laboratory analyses

Blood samples were collected into tubes containing EDTA or lithium heparin (S-Monovettes; Sarstedt Limited, Leicester, UK) and immediately placed on ice. Lithium-heparin blood samples for the measurement of ionised Ca (iCa) were measured within two minutes of collection using an ABL77
blood gas analyser (Radiometer, Bronshoj, Denmark) and the remaining samples were placed on ice. Corrected iCa values at pH 7.4 were used. Plasma was separated within 30 min of collection in a refrigerated centrifuge at 1800g for 20 min, and stored at −70°C. Collected urine was mixed thoroughly, and non-acidified and acidified (HCl, 10 ml/l, laboratory reagent grade, SG 1.18, Fisher Scientific, Loughborough, UK) aliquots were taken and stored at −20°C. After completion of the study, samples were packed and shipped on dry ice to MRC Human Nutrition Research, Cambridge for analysis. All assays were performed in duplicate with the exception of parathyroid hormone (PTH). Total Ca, P, Cr, total alkaline phosphatase, albumin and cystatin C were measured in heparinised plasma using colorimetric methods on the Kone Lab 20i clinical chemistry analyser platform (Kone, Espoo, Finland). On the same platform, urinary Ca (uCa), phosphate (uP) and creatinine (uCr) were measured in acidified urine. Within- and between-assay CV for all Kone assays were <2 and <4%, respectively. Concentrations of uCa and uP are expressed as a ratio relative to uCr to adjust for urine volume. EDTA plasma was used for the analysis of PTH by immunoassay (Immulite; Siemens Healthcare Diagnostics Limited, Camberley, UK). Between-assay CV was 4-7%. Total plasma 1,25(OH)2D was measured with a RIA (IDS Limited, Tyne and Wear, UK). Within- and between-assay CV were 7.5 and 9.0%, respectively. Cross-reactivity of the assay was 100 and 91% for 1,25(OH)2D3 and 1,25(OH)2D2, respectively. Assay performance was monitored using kit and in-house controls and under strict standardisation according to ISO 9001:2000. Quality assurance of 25(OH)D, 1,25(OH)D and PTH assays was performed as part of the Vitamin D External Quality Assessment Scheme (www.deqas.org) and the National External Quality Assessment Scheme (www.ukneqas.org.uk), and was within the accepted limits.

**MS**

25(OH)D2 and 25(OH)D3 were measured using ultra-performance LC-tandem MS. In brief, plasma proteins were precipitated with acetonitrile, dried and then reconstituted in deionised water, subjected to solid-phase extraction and, again, dried. The samples were then derivatised with 50 μl 4-phenyl-1,2,4-triazoline-3,5-dione (0.5 mg/ml) solution in acetonitrile. Then, 10 μl were injected into the ultra-performance LC system (Waters, Elstree, Herts, UK) consisting of Acquity UPLC BEH C18 1.7 μm, 2.1 x 50 mm columns (Waters) and a mobile-phase gradient of 0.1% formic acid of Acquity UPLC BEH C18 1.7
m/z 625 and 5 mM-methylamine in water and 0.1 % formic acid (Waters) and a mobile-phase gradient of 0.1 % formic acid

m/z 50 and, again, dried. The samples were then derivatised with

m/z 25(OH)D3 and 613

were precipitated with acetonitrile, dried and then reconstituted to commence from day 6.

**Data handling**

Data modelling and statistics were performed in Microsoft Excel 2003 (Microsoft Corporation, Seattle, WA, USA) and Data Desk 6.0 (Data Description, Ithaca, NY, USA). Within- and between-group differences were analysed with paired and unpaired t tests, respectively.

In order to describe the absorption and initial distribution of 25(OH)D2, we used the following equation:

\[
\frac{dA}{dt} = \begin{cases} 
F - k_Aq_A & t \leq T \\
-k_A & t > T
\end{cases}
\]

where \(q_A\) is the amount of 25(OH)D2 in a single compartment, \(F\) is the input constant, \(k_A\) is the fractional rate constant of loss and \(T\) is the duration of the input. This model assumes a single compartment, a constant input of the 25(OH)D2 dose and first-order disappearance. Initial estimates of the model parameters were then subsequently adjusted with the Solver function in Excel to minimise the differences between the modelled (predicted) and observed data. The absorption parameter \(T\) is equivalent to \(T_{max}\) (the time taken to reach the maximum plasma concentration, \(C_{max}\)). \(C_{max}\) was calculated from the absorption parameters \(F\), \(T\) and \(k_A\) using the following equation:

\[
C_{max} = \frac{F}{k_A} \times 1 - e^{-k_T t}
\]

Plasma 25(OH)D2 half-life was calculated from the following equation:

\[
T_1/2 = \frac{\ln(2)}{k_B}
\]

From visual inspection of the logarithmically transformed curves, \(k_B\) (the slope of terminal disappearance) was identified to commence from day 6. \(k_B\) was then derived from data obtained between days 6 and 21 and was calculated from the slope of the line of best-fit of the natural logarithm of 25(OH)D2 concentration against time.

**Results**

**Biochemical markers of vitamin D, calcium and phosphate metabolism**

There were no significant differences in any of the subjects or biochemical parameters between the groups at day 0 or 15, thus the groups were combined for further analysis of biochemical parameters (Table 1). There were no significant changes in plasma iCa, total Ca, P, PTH and 1,25(OH)2D3, or uCa/uCr and uP/uCr between days 0 and 15 (Table 1). A significant decrease was observed in total alkaline phosphatase (\(P=0.012\)) and a significant increase in plasma creatinine (\(P=0.047\), but not in cystatin C. However, all values for kidney and liver function were within the normal clinical
25-Hydroxyvitamin D2 absorption and half-life

Table 1. Baseline and day 15 post-dose plasma and urinary concentrations of calcium (uCa) and phosphate (uP), and calcitropic hormones† (Mean values and standard deviations, n 12)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Baseline (day 0)‡</th>
<th>Post-dose (day 15)‡</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>sd</td>
</tr>
<tr>
<td>25(OH)D2 (nmol/l)</td>
<td>1·1</td>
<td>0·5</td>
</tr>
<tr>
<td>25(OH)D3 (nmol/l)</td>
<td>54·3</td>
<td>10·4</td>
</tr>
<tr>
<td>1,25(OH)D2 (pmol/l)</td>
<td>290·3</td>
<td>57·6</td>
</tr>
<tr>
<td>PTH (pg/ml)</td>
<td>Geometric mean</td>
<td>71·8</td>
</tr>
<tr>
<td></td>
<td>CI</td>
<td>48·0, 107·6</td>
</tr>
<tr>
<td>iCa§ (mmol/l)</td>
<td>1·06</td>
<td>0·05</td>
</tr>
<tr>
<td>Ca (mmol/l)</td>
<td>2·29</td>
<td>0·09</td>
</tr>
<tr>
<td>P (mmol/l)</td>
<td>1·34</td>
<td>0·25</td>
</tr>
<tr>
<td>Albumin (g/l)</td>
<td>38·80</td>
<td>2·45</td>
</tr>
<tr>
<td>Creatinine (µmol/l)</td>
<td>82·47</td>
<td>7·54</td>
</tr>
<tr>
<td>Cystatin C (mg/l)</td>
<td>0·92</td>
<td>0·14</td>
</tr>
<tr>
<td>TALP (U/l)</td>
<td>262·8</td>
<td>76·6</td>
</tr>
<tr>
<td>uCa/uCr (mmol/l)</td>
<td>0·08</td>
<td>0·08</td>
</tr>
<tr>
<td>uP/iCr (mmol/l)</td>
<td>1·11</td>
<td>0·40</td>
</tr>
</tbody>
</table>

25(OH)D, 25-hydroxyvitamin D; 1,25(OH)2D, 1,25-dihydroxyvitamin D; PTH, parathyroid hormone; iCa, ionised calcium; TALP, total alkaline phosphatase; uCr, urinary creatinine.
† There were no significant differences between the absorption group (n 6) and the half-life group (n 6) in any of the subjects or biochemical characteristics.
‡ Mean values were significantly different from baseline values (day 0); *P=0·0001; **P=0·047; ***P=0·05 (paired t test).
§ Corrected value at pH 7·4.

range and changes were small and clinically insignificant. No biochemical abnormalities were detected and values were comparable with other data obtained from this age group and population. Total 25(OH)D was calculated as the sum of 25(OH)D2 and 25(OH)D3 plasma concentrations. There was no significant difference in total 25(OH)D or 25(OH)D3 plasma concentrations. The plasma concentration of 25(OH)D3 increased significantly from 1·1 (sd 0·5) at day 0 to 3·1 (sd 0·2) nmol/l at day 15 (P>0·0001). During the study, there were no reports of side effects or sickness.

Absorption, distribution and terminal slope of 25-hydroxyvitamin D2

Low but detectable concentrations of 25(OH)D2 were observed in all subjects and time points; therefore, all subsequent concentrations used for data modelling and half-life calculations were corrected for baseline 25(OH)D2. Cmax was 9·6 (sd 0·9) nmol/l and was observed at Tmax, 4·4 (sd 1·8) h (Figs. 1 and 2). The input constant, F, was 2·6 nmol/h. The initial disappearance rate constant (k0) calculated from the peak of the absorption curve and using the remaining data points after Tmax until 48 h was 0·36 (sd 0·14). Individual plasma response curves are plotted in Fig. 2 and highlight the consistent pattern between individuals. For the second group sampled between days 3 and 21, values are plotted on a semi-logarithmic scale. From a visual inspection of Fig. 2, it is noted that the terminal phase commenced at day 6. The increase in the plasma concentration of 25(OH)D2 at the commencement of the terminal slope was 3·4 nmol/l, approximately 6% of the total plasma concentration of 25(OH)D2. This, together with the lack of evidence of significant changes in Ca, P or vitamin D metabolism, demonstrates that there was no perturbation of the steady state.

The absorption and half-life of 25(OH)D3 in healthy individuals have been measured in a number of studies (summarised in Tables 2 and 3). Absorption studies frequently used very

This terminal disappearance phase (k0) between days 6 and 21 had a slope of 0·05 (sd 0·01). The mean half-life as calculated from individual slopes was 13·4 (sd 2·7) d. Individual plasma curves (Fig. 4) highlight again the consistent pattern between individuals and clearly show that by day 6, 25(OH)D2 disappearance was in the terminal phase in all subjects.

We explored relationships between Cmax, Tmax and half-life and selected physiological and biochemical parameters. A significant positive correlation was found between Cmax and BMI (r 0·87; P=0·02) but not with height (r 0·03; P=0·96) and weight (r 0·73; P=0·10). There was no association between either Tmax or Cmax and baseline 25(OH)D3 plasma concentration. There were no significant relationships between half-life and measures of plasma iCa, total Ca, P, TALP, 1,25(OH)D or 25(OH)D3 at day 15.

Discussion

The half-life of 25(OH)D may provide a useful biomarker to investigate vitamin D requirements. In the present pilot study, we characterised the absorption, distribution and terminal disappearance of 25(OH)D2 as a tracer to investigate vitamin D metabolism and to obtain baseline data for future studies. The protocol was feasible under field conditions and was well accepted by the subjects. The absorption parameters obtained in the present study indicated that the dose was consistently well absorbed. The data showed that the terminal slope commenced at day 6. The increase in the plasma concentration of 25(OH)D2 at the commencement of the terminal slope was 3·4 nmol/l, approximately 6% of the total plasma concentration of 25(OH)D2. This, together with the lack of evidence of significant changes in Ca, P or vitamin D metabolism, demonstrates that there was no perturbation of the steady state.

The absorption and half-life of 25(OH)D2 in healthy individuals have been measured in a number of studies (summarised in Tables 2 and 3). Absorption studies frequently used very
large unlabelled doses that may have perturbed normal vitamin D metabolism (Table 2). Studies reporting 25(OH)D$_3$ half-lives using unlabelled, radiolabelled and stable isotope-labelled compounds after oral or intravenous administration have reported widely different values (Table 3). Nevertheless, within-study comparisons demonstrated differences in 25(OH)D$_3$ half-life between controls and anephric$^{(9)}$ and between individuals on either a high- or normal-fibre diet$^{(11)}$. The wide variation between methods may have been caused by methodological differences and limitations in the experimental design or quantification techniques, such as an improperly defined terminal slope or measurement of total radioactivity.

There are advantages to using oral 25(OH)D$_3$ rather than vitamin D$_3$ to measure 25(OH)D$_3$ half-life that are based on a number of factors which relate to differences in absorption, tissue distribution and hepatic metabolism$^{(21,26–29)}$. In view of the similarities in the absorption and distribution of vitamin D$_2$ and vitamin D$_3$$^{(30–32)}$, these advantages may also apply to 25(OH)D$_2$. Absorption of 25(OH)D$_3$ is greater, faster and more consistent than that of vitamin D$_3$$^{(33)}$. Net absorption of vitamin D$_3$ in healthy human subjects has been reported to be between 55 and 99% of the dose$^{(21,34)}$, whereas 25(OH)D$_3$ absorption ranged from 88 to 100% in healthy subjects. This may be related to the difference in absorption mechanism, i.e. for vitamin D$_3$, this is primarily into the enterocytes and the lymph system whereas the primary route for 25(OH)D$_3$ absorption is via the portal vein$^{(35,36)}$, although another study has suggested that the absorption mechanism is the same for vitamin D and 25(OH)D$^{(37)}$. Differences in the principal absorption mechanism are also supported by evidence from studies in patients with various gastrointestinal disorders affecting chylomicron formation, in which absorption of vitamin D is affected to a greater extent than 25(OH)D$^{(20,21,26)}$. Vitamin D metabolites in the circulation are predominantly bound to DBP; however, a proportion of an oral dose, particularly for the less-polar vitamin D, will initially be associated with chylomicrons that may further influence initial metabolism and sequestration into body tissues. Therefore, in terms of delivering the maximum and most consistent plasma response from a minimum tracer dose, 25(OH)D is more effective than vitamin D.

For the determination of 25(OH)D half-life, the use of a tracer or labelled 25(OH)D rather than tracer or labelled precursor (i.e. vitamin D) is also likely to reduce variability because factors are removed that will influence the rate of appearance of 25(OH)D in the circulation. The appearance of 25(OH)D after a bolus dose of vitamin D is determined by the rate and extent of hepatic hydroxylation and sequestration and release of vitamin D from body stores. Little is known

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Fig. 2. Plasma appearance and disappearance over 48 h after an oral dose of 25-hydroxyvitamin D$_2$ (25(OH)D$_2$) in six subjects (A–F) in the absorption group. ○, Observed plasma concentration of 25(OH)D$_2$; —, modelled plasma curve.

Fig. 3. Semi-logarithmic plot of plasma concentration between days 3 and 21 after an oral dose of 25-hydroxyvitamin D$_2$ (25(OH)D$_2$) in the half-life group (n 6). Values are means, with standard deviations represented by vertical bars, and the terminal slope (0·05 (SD 0·01)) calculated from day 6 onwards represented by a dashed line.
about the relationship between the serum concentration of vitamin D and its tissue distribution. Appreciable amounts of vitamin D may be taken up by tissues (principally adipose tissue) and this may be influenced by the plasma concentration of 25(OH)D and saturation of the 25-hydroxylase enzyme(38). The majority of 25(OH)D is found in plasma, with smaller amounts in fat and fat-free mass(29,38). The potential slow release of tracer vitamin D from body stores and the consequent further and additional hepatic production of labelled 25(OH)D into the circulation would cause fluctuations in labelled 25(OH)D concentration and may distort the 25(OH)D half-life. The use of tracer 25(OH)D prevents the continued production of labelled 25(OH)D from labelled vitamin D and allows the establishment of faster equilibrium between body pools. In some earlier studies, increased variability may have been caused by measurement of total radioactivity that included the vitamin D3 dose and all downstream metabolites (predominantly 25(OH)D 3, 1,25(OH) 2D3 and 24,25(OH) 2D3), and therefore affected the accuracy of the calculated half-life. However, it was reported that between 4 and 40 d after a dose of vitamin D3 or 25(OH)D3, more than 80% of radioactivity is associated with 25(OH)D3. Up to 5% remains associated with vitamin D3 and up to 10% with more polar metabolites(8,9). Modern LC-MS/MS, as used in the present study, enhances the measurement specificity.

In the present study, as expected, the oral dose increased 25(OH)D2 concentrations (Figs. 1 and 2). Cmax was 9·6 (SD 0·9) nmol/l and occurred at Tmax, 4·4 (SD 1·8) h. These data support the notion of rapid and efficient absorption in the upper gastrointestinal tract. The results of absorption parameters Cmax and Tmax from the present study are similar to those previously observed for 25(OH)D3 (Table 2). This observation is consistent with comparisons of vitamin D2 and vitamin D3 absorption where no differences have been observed in the initial plasma response after oral doses(30–32,39). The small standard deviations in the absorption parameters demonstrate

Table 2. Absorption of 25-hydroxyvitamin D (25(OH)D) measured with oral doses of unlabelled or radiolabelled 25(OH)D (Mean values and standard deviations or standard errors)

<table>
<thead>
<tr>
<th>Dose</th>
<th>n</th>
<th>Tmax (h)</th>
<th>Cmax (nmol/l)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>The present study</td>
<td>25(OH)D3 33 μg (0·5–0·7 μg/kg)</td>
<td>6</td>
<td>4·4†</td>
</tr>
<tr>
<td>Haddad &amp; Rojanasathit(10)</td>
<td>Unlabelled 25(OH)D3 1·5, 5 or 10 μg/kg</td>
<td>22</td>
<td>4–8</td>
</tr>
<tr>
<td>Compston &amp; Creamer(66)</td>
<td>[*H]25(OH)D3 (1·8 nmol) + 26 μg reference 25(OH)D3</td>
<td>3</td>
<td>6–10</td>
</tr>
<tr>
<td>Compston et al.(27)</td>
<td>Unlabelled 25(OH)D3 10 μg/kg body weight</td>
<td>12</td>
<td>4–8</td>
</tr>
<tr>
<td>Davies et al.(21)</td>
<td>[*H]25(OH)D3 26 μg</td>
<td>5</td>
<td>7·2</td>
</tr>
<tr>
<td>Leichtmann et al.(22)</td>
<td>[*H]25(OH)D3 with 100 μg unlabelled 25(OH)D3</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>Sitrin &amp; Bengoa(20)</td>
<td>[*H]25(OH)D3 with 100 μg unlabelled 25(OH)D3</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>The present study</td>
<td>[3H]25(OH)D3 with 100 μg unlabelled 25(OH)D3</td>
<td>8</td>
<td>NR</td>
</tr>
</tbody>
</table>

NR, not reported.
* To convert nmol/l to μg/l, multiply by 0·400 for 25(OH)D2 and 0·412 for 25(OH)D3.
† se 1·8 h.
‡ se 0·9 nmol/l.
§ se 33 nmol/l.
|| se 0·5 h.
† se 0·9 nmol/l.
Table 3. Terminal plasma half-life of 25-hydroxyvitamin D (25(OH)D) (Mean values and standard deviations or standard errors)

<table>
<thead>
<tr>
<th>Dose Route</th>
<th>n</th>
<th>Measured over</th>
<th>Half-life (d)</th>
<th>Quantification</th>
<th>RA of polar metabolite measured</th>
<th>Total RA</th>
<th>RA of polar metabolite measured</th>
<th>Total RA</th>
<th>LC-MS/MS</th>
<th>Competitive binding radioassay</th>
</tr>
</thead>
<tbody>
<tr>
<td>The present study</td>
<td>33</td>
<td>Oral</td>
<td>6–21 d</td>
<td>27</td>
<td>Mean 11·4</td>
<td>SD 2·7</td>
<td>LC-MS/MS</td>
<td>RA</td>
<td>6–21 d</td>
<td>13·4 2·7</td>
</tr>
<tr>
<td>Mawer et al.</td>
<td>7</td>
<td>Oral</td>
<td>4–10 d</td>
<td>400</td>
<td>Mean 20·7</td>
<td>SD 0·0</td>
<td>LC-MS/MS</td>
<td>RA</td>
<td>4–10 d</td>
<td>20·7 0·0</td>
</tr>
<tr>
<td>Smith &amp; Goodman</td>
<td>8</td>
<td>Oral</td>
<td>4–10 d</td>
<td>51</td>
<td>Mean 10·7</td>
<td>SD 0·7</td>
<td>LC-MS/MS</td>
<td>RA</td>
<td>4–10 d</td>
<td>11·3 0·7</td>
</tr>
<tr>
<td>Vicchio et al.</td>
<td>7</td>
<td>Oral</td>
<td>4–10 d</td>
<td>2·0</td>
<td>Mean 2·0</td>
<td>SD 0·0</td>
<td>LC-MS/MS</td>
<td>RA</td>
<td>4–10 d</td>
<td>2·0 0·0</td>
</tr>
</tbody>
</table>

* Values are expressed as standard errors.

We identified the terminal phase to commence from day 6 (Figs. 3 and 4) and the terminal plasma half-life of 25(OH)D2 was determined to be 13·4 (sd 2·7) d (range 9·6–17·3 d). This is somewhat less than half-lives reported earlier in healthy individuals after doses of radiolabelled vitamin D3 or 25(OH)D3 (Table 3), which ranged from 10 to 40 d. The only study that used a stable isotope-labelled compound has reported a value of 10·4 d(12). Half-life was not associated with the underlying plasma concentration of 25(OH)D3 and may not be surprising given the relatively narrow range of concentrations (range 38·4–69·0 nmol/l). This observation is consistent with some studies(41,42) but not with others(7,43), and is an area that requires further investigation. The short half-life in the present study may be due to differences in half-life between 25(OH)D2 and 25(OH)D3, the study population or the experimental approach between the present and other studies (Table 3). Evidence from studies using unlabelled compounds suggests that 25(OH)D2 is cleared more rapidly from plasma than 25(OH)D3(30,31,59), and this is thought to be due to their relative binding affinities for DBP(13,14,44). Alternatively, correction for 25(OH)D2 concentration at baseline and subsequent inconsistent intake during the study period may have introduced variability in plasma concentrations of 25(OH)D2. In the absorption phase, fluctuation in 25(OH)D2 plasma concentration is likely to have had only a very minor effect because of the higher concentration from the dose and shorter time period. However, fluctuations in vitamin D3 intake may have contributed more significantly to variability in 25(OH)D2 half-life because of the lower concentrations during the terminal slope. In addition, the unexpected baseline levels of 25(OH)D2 and subsequent subtraction may have introduced additional error in the determination of half-life. This population in rural Gambia is known to have a low dietary Ca intake (300–400 mg/d)(25,45), a high fibre intake and elevated 1,25(OH)2D and secondary hyperparathyroidism(25), and are consistent with the findings of the present study. All these factors are known to increase 25(OH)D turnover(11,46,47). Furthermore, longitudinal bone growth and bone mineral accrual still occurs in Gambian men of this age(48), thus their tissue requirement for Ca and vitamin D may be higher than an older age group(49–51). Total 25(OH)D was calculated as the sum of 25(OH)D2 and 25(OH)D3 concentrations as measured by LC-MS/MS. In the present study, the mean baseline total 25(OH)D plasma concentration was about
55 nmol/l. Notwithstanding known assay differences, this value is somewhat lower than previous measurements in this and other age groups in The Gambia and Guinea-Bissau, and may be surprising given the year-round ability for cutaneous vitamin D production and plentiful sunshine. The relatively low values could be attributable to a number of causes as discussed above (i.e. skeletal development and low Ca intake).

Plasma 25(OH)D3, total 1,25(OH)2D, PTH, Ca, iCa and P and uCa and uP did not change from baseline, indicating that the tracer did not perturb Ca, P and vitamin D metabolism. This contrasts with another study in which a much higher dose (1 mg) resulted in elevated uCa and hydroxyproline but without changes in plasma values of Ca or P. This indicates alterations of Ca and bone metabolism and the resultant half-life may therefore not be reflective of habitual vitamin D turnover.

25(OH)D2 can only be used as a tracer in a population with absent or very low 25(OH)D plasma concentration, or after a period of weeks or months of vitamin D2 depletion, and thus its use may not be suitable in all populations, although recent evidence suggests that circulating plasma levels in the UK are similar to those observed in the present study (H. Macdonald, personal communication). This study was performed in a rural Gambian village where vitamin D2 intake was not expected. This was supported by recent work from Guinea-Bissau in which 25(OH)D2 measured by LC/MS was detected in only seven out of 856 subjects. Surprisingly, however, low but detectable concentrations of 25(OH)D2 (ranging from 0.6 to 2.5 nmol/l) were found in all baseline samples from all subjects. An explanation for this observation may be the non-identity of vitamin D2 and vitamin D3 to DBP are not the same and may affect plasma half-life suitable for use in adults and children. In populations with higher Ca intake and therefore lower vitamin D usage, a longer half-life might be expected. Additional studies are required to optimise the duration of sample collection to estimate half-life in different populations. 25(OH)D half-life may be useful as a biomarker of vitamin D metabolism to investigate environmental or physiological factors that may affect vitamin D usage and to better assess vitamin D requirements.

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