Rat adipose tissue rapidly accumulates and slowly releases an orally-administered high vitamin D dose

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We investigated the effect of oral high-dose cholecalciferol on plasma and adipose tissue cholecalciferol and its subsequent release, and on plasma 25-hydroxyvitamin D (25(OH)D). Female Wistar rats (n 126) received 37.5 μg cholecalciferol/d for 14 d and were subsequently studied for a further 88 d. Two subgroups of eighteen rats each were fasted for 3 d immediately after treatment (days 14–17) and at the end of the study (days 98–101). During treatment, plasma cholecalciferol increased rapidly to reach a steady-state. Plasma 25(OH)D and adipose tissue cholecalciferol increased linearly for 1–2 d after treatment. Serum Ca and inorganic phosphate also increased. Subsequently half-lives of plasma cholecalciferol and 25(OH)D, and perirenal and subcutaneous adipose tissue were: 1.4, 22.5, 97.5 and 80.9 d respectively. Fasting, as compared with ad libitum feeding, caused increased plasma free fatty acids, weight loss up to 14 % and increased adipose tissue cholecalciferol (nmol/g wet weight). It did not affect plasma cholecalciferol immediately after cholecalciferol treatment, but raised plasma 25(OH)D. Fasting at the end of the study decreased plasma cholecalciferol and increased plasma 25(OH)D. We conclude that orally-administered cholecalciferol rapidly accumulates in adipose tissue and that it is very slowly released while there is energy balance. Fasting causes preferential loss of triacylglycerols from adipose tissue, as opposed to cholecalciferol, but nevertheless augments plasma 25(OH)D. Adipose tissue may act as a 'buffer to functional vitamin D status' by preventing, to a certain extent, unregulated production of 25(OH)D from dietary vitamin D, and by slowly releasing vitamin D under fasting conditions.

Cholecalciferol: Plasma 25-hydroxyvitamin D: Calcium: Blood

Adequate vitamin D status is essential for the prevention of rickets in children, and osteomalacia (Fraser, 1980) and probably osteoporosis (Compston, 1995) in adults. Based on, for example, the wide organ distribution of the vitamin D receptor, it is becoming increasingly clear that vitamin D metabolites are not only important in (bone) Ca homeostasis, but also for adequate functioning of many other systems (Reichel et al. 1989).

Present vitamin D supplementation recommendations concern post-menopausal women (Dawson-Hughes et al. 1991; Compston, 1995) and the elderly over 75 years (The Committee on Nutrition of the Elderly, Food and Nutrition Council, 1995) who live at high latitudes. These groups are recommended to take vitamin D supplements during winter. The daily dosage is, however, subject to discussion, because of remaining concern about (cumulative) toxicity (Schwartzman & Franck, 1987).

Plasma 25-hydroxyvitamin D (25(OH)D) level is widely regarded as the most reliable index of vitamin D status. However, oral supplementation of adults with either 10 or 20 μg cholecalciferol/d does not cause much difference in plasma 25(OH)D concentrations (Lips et al. 1988; van der Klis et al. 1996). On the basis of these results, it was suggested that adipose tissue functions as a vitamin D-buffering system that to a certain extent prevents uncontrolled synthesis of 25(OH)D in the liver (van der Klis et al. 1996). Mawer et al. (1972) had suggested previously that storage in tissue lipid may limit potential vitamin D toxicity, while Rosenstreich et al. (1971) suggested that slow release of vitamin D from adipose tissue may be an important factor in long-term vitamin D status. We investigated the effect of daily oral administration of high-dose (37.5 μg) cholecalciferol to rats on cholecalcif-
erol accumulation in adipose tissue and its subsequent release into plasma under *ad libitum* feeding conditions and during short-term fasting. Cholecalciferol was administered for 14 d and the levels of cholecalciferol in plasma and adipose tissue and of 25(OH)D in plasma were monitored from day 0 up to day 101. To promote adipose tissue mobilization, two subgroups of rats were fasted for 3 d, immediately after cessation of cholecalciferol treatment and 84 d after discontinuation of treatment. Serum Ca and phosphorus were monitored to detect biochemical signs of vitamin D toxicity.

**Materials and methods**

*Animals, vitamin D and chemicals*

Female Wistar rats, weighing 150 - 180 g, were obtained from Harlan (Zeist, The Netherlands). They were housed under standard laboratory conditions (12 h dark - 12 h light, 25°) with free access to acidified tap water (pH 2.5 - 3) and standard rodent chow, unless otherwise stated. The standard chow contained 37.5 μg cholecalciferol/kg, corresponding to a daily background intake of about 0.75 μg cholecalciferol. Cholecalciferol (cholecalciferol densitosem olesum, 50 mg/g) for administration was obtained from OPG (Utrecht, The Netherlands). For administration orally, it was diluted in arachid oil to 37.5 μg/ml. Chemicals for the analysis of cholecalciferol by HPLC were obtained from Merck (Darmstadt, Germany) and Rathburn (Brunswig Chemie, Amsterdam, The Netherlands). Ergocalciferol and cholecalciferol standards for quantification were obtained from Sigma (St Louis, MO, USA).

**Experimental**

A group of 126 rats received orally 37.5 μg cholecalciferol (dissolved in 1 ml arachid oil) daily for 14 d (days 0 - 13) at 22.00 hours. The last dose was administered 22.00 hours on day 13. Groups of six rats (a total of ninety rats) were killed at 10.00 hours under halothane anaesthesia on day 0 (i.e. before treatment), days 3, 6, 9 and 12 (i.e. during treatment) and days 14, 15, 16, 17, 40, 76, 98, 99, 100 and 101 (i.e. after treatment). The rats were weighed and samples of blood were drawn from the aorta both in the presence (to permit coagulation) and absence of EDTA. Serum and blood were drawn from the aorta both in the presence of EDTA (dissolved in 1 ml arachid oil) daily for 14 d (days 3, 6, 9 and 12). Isotonic elution of vitamin D was performed using a Waters M6000A pump (Waters, Milford, MA, USA) at a flow-rate of 1.5 ml/min, using a Spheri-5 RP18 column (Brownlee) and a Waters 486 tunable absorbance U.V. detector at 260nm. Plasma (nmol/l) and serum (nmol/l) 25-hydroxyergocalciferol and 25-hydroxycholecalciferol were measured as 25(OH)D by using a competitive radio-binding assay with 3H-labelled 25-hydroxycholecalciferol and vitamin D-binding protein from human serum after solid-phase extraction. Between-series CV at 16, 62 and 101 nmol/l were 13, 12 and 13 % respectively.

Plasma and adipose tissue cholecalciferol levels were determined using reversed-phase HPLC with u.v. detection, following saponification, organic solvent extraction and further prepurification with straight-phase HPLC, essentially according to Thompson *et al.* (1982). Ergocalciferol served as an internal standard. In short, the sample (1 ml plasma, 1 g adipose tissue) was saponified overnight in ethanolic KOH at 70° and extracted with hexane. For adipose tissue samples, the hexane layer was washed consecutively with KOH (50 g/l) ethanol (300 ml/l) in saline (9 g NaCl/l) and saline. After evaporation to dryness at room temperature under a stream of N2, the residue was dissolved in 1.5 ml hexane, followed by cholesterol precipitation with methanolic digitonin. After centrifugation, the hexane layer was collected, evaporated to dryness (at room temperature under N2), and redissolved in the straight-phase HPLC eluent (hexane-isopropanol; 99.5:0.5, v/v). Isotonic elution of vitamin D was performed using a Waters M6000A pump (Waters, Milford, MA, USA) at a flow-rate of 1.5 ml/min, using a Spheri-5 silica column (220 x 4.6 mm; Brownlee Applied Biosystems, Foster City, CA, USA) and a Waters 486 tunable absorbance u.v. detector at 260 nm. This system does not separate ergocalciferol and cholecalciferol. A cholecalciferol standard was used for the establishment of their retention time and their subsequent collection from sample extracts. The collected vitamin D fraction was evaporated to dryness (at room temperature under N2). The purified vitamin D extract was redissolved in the reversed-phase HPLC eluent (acetonitrile-tetrahydrofuran-water; 94:5:0.5, v/v). Separation and detection of ergocalciferol and cholecalciferol was performed using a Waters M6000A pump at a flow-rate of 1.0 ml/min, using a Spheri-5 RP18 column (Brownlee) and a Waters 486 tunable absorbance u.v. detector at 260 nm. Plasma (nmol/l) and adipose tissue (nmol/g wet weight) cholecalciferol levels were calculated by comparing the peak area ratios of ergocalciferol and cholecalciferol with those of equimolar...
Rat adipose tissue rapidly accumulates vitamin D standards. The within-series precision for cholecalciferol in plasma was calculated by measurement of six samples of a human plasma pool on six different occasions. The overall mean plasma cholecalciferol concentration was 134 nmol/l and the mean within-series CV amounted to 3.4 (range 1.7 - 7.7) %. The between-series precision and recovery were established by analysing a rat plasma pool with and without enrichment with 130 nmol cholecalciferol/l on four occasions. The mean endogenous cholecalciferol concentration amounted to 364 nmol/l (CV 2.7 %) and the mean recovery was 93.6 % (CV 2.7 %). The within-series precision of cholecalciferol in adipose tissue was measured by the 6-fold analysis of hog lard. The CV was 28.1 % at a mean cholecalciferol content of 32 pmol/g. The within-series recoveries were measured by enrichment of lard with 78, 156, 234 and 312 pmol cholecalciferol/g. The recoveries (%; n 6) amounted to: 73.6 (CV 5.2 % at 78 pmol/g), 92.5 (CV 3.9 % at 156 pmol/g), 94.6 (CV 3.0 % at 234 pmol/g) and 91.2 (CV 3.5 % at 312 pmol/g).

Data analyses

Body weight was calculated relative to the body weight on day 14 and day 98 (start of fast) for fasting and ad libitum-fed rats that were studied on days 15 – 17 and days 99 – 101 respectively. Serum and adipose tissue half-lives were estimated using standard pharmacokinetic calculations, using a program developed by Dr H. Proost, Department of Pharmacy, University of Groningen. The calculations assumed 1st-order kinetics without lag-time, one compartment and a multiple dosage regimen. Differences between groups were analysed using Student’s t test (Stevens, 1980). ANOVA (Stevens, 1980) was used to determine steady-state conditions. P < 0.05 was considered significant.

Results

Fig. 1 shows the mean plasma cholecalciferol and 25(OH)D concentrations, together with the subcutaneous and perirenal cholecalciferol levels before, during and after oral administration of cholecalciferol to rats. Cholecalciferol was given at 22.00 hours from day 0 to day 13 and samples were taken at 10.00 hours. Plasma cholecalciferol increased rapidly, to reach steady-state levels at day 3. After treatment, plasma cholecalciferol declined rapidly, with a calculated half-life of 1.4d. Levels at the end of the study (day 101, 31.3 (SD 1.5) nmol/l) were comparable with those on day 0 (31.8 (SD 2.3) nmol/l). Levels of 25(OH)D in

![Fig. 1. Time-courses of plasma cholecalciferol (a) and 25-hydroxyvitamin D (25(OH)D; (b)) and subcutaneous (c) and perirenal (d) adipose tissue cholecalciferol contents in rats, before, during and after supplementation with 37.5 pg cholecalciferol/d. Cholecalciferol was administered orally from day 0 to day 13 at 22.00 hours and samples were taken at 10.00 hours. ( ), Supplementation period. Data points represent means with their standard errors, represented by vertical bars, for groups of six rats. For details of procedures, see p. 528.](https://www.cambridge.org/core/terms). https://doi.org/10.1079/BJN19980091
plasma, and cholecalciferol in subcutaneous and perirenal adipose tissue increased linearly during treatment. They subsequently declined, with half-lives (d) of 22.5 (plasma 25(OH)D), 80.9 (subcutaneous adipose tissue), and 97.4 (perirenal adipose tissue). Plasma levels of 25(OH)D on day 101 (132 (SD 11)nmol/l) were below (<0.0001) those on day 0 (225 (SD 8)nmol/l). Subcutaneous and perirenal adipose tissue cholecalciferol contents at the end of the study (0.77 (SD 0.03) and 0.65 (SD 0.02)nmol/g respectively) were above (<0.0001) those on day 0 (0.30 (SD 0-04) and 0.24 (SD 0.03)nmol/g respectively).

Serum Ca and phosphate concentrations increased from baseline (day 0) to the end of the treatment (day 14); levels were: Ca 2.51 (SD 0.15) v. 2.96 (SD 0.10)nmol/l (P = 0.0001); phosphate 2.10 (SD 0.32) v. 2.69 (SD 0.32)nmol/l (P = 0.001). Serum phosphate reached baseline levels 3 days after treatment (2.14 (SD 0.23)nmol/l, day 17). Serum Ca was still elevated 3 days after treatment (2.68 (SD 0.07) mmol/l, P = 0.05, day 17) and reached baseline levels by 26 days after treatment (2.57 (SD 0.04)nmol/l, day 40).

Fig. 2 shows plasma free fatty acid concentrations and the relative body weights of fasting and ad libitum-fed rats. Rats were fasted immediately after vitamin D treatment (days 14–17) and at about 3 months after its discontinuation (days 98–101). Fasting rats had higher free fatty acid concentrations than their ad libitum-fed counterparts. Compared with the controls, fasted rats had lower relative body weights from 1 day after initiation of fasting. After a 3-day fast, they have lost 15.6% (days 14–17) and 13.0% (days 98–101) of their body weights; ad libitum-fed rats did not lose weight.

Fig. 3 shows the changes in plasma cholecalciferol and 25(OH)D concentrations, and in subcutaneous and perirenal adipose tissue cholecalciferol contents in fasting and ad libitum-fed rats. Plasma cholecalciferol and 25(OH)D concentrations of each of the groups decreased (ANOVA, P < 0.05) from day 14 to day 17 and from day 98 to day 101. When compared with ad libitum-fed rats, fasting rats showed no difference in plasma cholecalciferol concentrations from day 14 to day 17, but they had lower cholecalciferol concentrations from day 99 to day 101. Plasma 25(OH)D levels of fasting rats were higher on days 16 and 17, and on day 101.

In fasted rats cholecalciferol content (nmol/g wet tissue) of subcutaneous and perirenal adipose tissues increased between day 14 and day 17, but not from day 98 to day 101. Compared with ad libitum-fed rats, fasting rats had higher subcutaneous adipose tissue contents on days 16 and 17, and on days 100 and 101. Fasted rats had higher perirenal adipose tissue contents on days 17 and 101.

Neither fasting period caused differences in serum Ca and phosphate concentrations between fasting and ad libitum-fed rats.

Discussion

We monitored the levels of cholecalciferol in plasma and adipose tissue of rats that received 37.5μg cholecalciferol/d for 14 d. This dose amounted to about fifty times their usual daily intake via the diet. The cumulative dose equalled their usual intake in 700 d. After administration we monitored plasma and adipose tissue cholecalciferol under ad libitum feeding conditions for another 88 d, and studied the effect of a 3 d fast in the period immediately after treatment and at the end of the study. We also measured plasma 25(OH)D, since vitamin D is rapidly converted into 25(OH)D when it reaches the liver. Furthermore, 25(OH)D is generally considered to be a reliable index of functional vitamin D status because 25(OH)D has a long half-life and is the immediate precursor of 1,25-dihydroxyvitamin D.

During treatment, plasma cholecalciferol showed a very rapid increase and a subsequent steady-state, whereas adipose tissue cholecalciferol and plasma 25(OH)D showed linear increases up to 1–2 d after discontinuation of cholecalciferol treatment (Fig. 1). The steadily increasing plasma 25(OH)D, in the absence of any change of plasma cholecalciferol, seems of particular importance, since it is generally believed that vitamin D toxicity is caused by the interaction of high concentrations 25(OH)D with the vitamin...
D receptor. In agreement with this suggestion, we observed increases in serum Ca and inorganic phosphate during cholecalciferol treatment, while Ca levels at the end of the treatment (day 14: 2.96 mmol/l) were close to those (3.3 mmol/l) reported for rats that received toxic vitamin D doses (three oral doses of 2.5 mg; Świerczyński et al. 1987). The steady increase in 25(OH)D at 37.5 μg cholecalciferol/d seems to contrast with the observations of Lips et al. (1988) and van der Klis et al. (1996) in human subjects. From their studies, it can be concluded that healthy adults who receive 10 or 20 μg cholecalciferol/d show little or no difference in plasma 25(OH)D during 1 year of treatment, and that steady-state levels are reached in 1 week or less. The short-term conversion efficiency of orally-administered vitamin D to 25(OH)D may be dependent, therefore, on the vitamin D dose, and it is possible that above certain daily doses accumulation of 25(OH)D in plasma cannot be avoided. It is conceivable that the underlying mechanism of plasma 25(OH)D accumulation is increasing vitamin D uptake by the liver, as opposed to storage in adipose tissue, causing increasing exposure of vitamin D to the poorly regulated (Holick, 1994) hepatic 25-hydroxylase. A lower (physiological) vitamin D dose is likely, therefore, to exhibit different kinetics, and it also remains to be established whether present findings relating to a high dose can be extrapolated to human subjects. Ca and phosphate levels decreased rapidly after discontinuation of cholecalciferol administration (data not shown) concomitant with the decreases in plasma cholecalciferol and 25(OH)D (Fig. 1). Calculated half-lives of plasma cholecalciferol and 25(OH)D were in agreement with those reported for human subjects by Mawer et al. (1969, 1971; cholecalciferol 1 d, 25(OH)D 15–30 d). In contrast to plasma cholecalciferol and 25(OH)D, adipose tissue cholecalciferol contents were still well above baseline levels at the end of the study. A similar pattern showing a gradual decrease after cessation of cholecalciferol administration was also reported by Rosenstreich (1971) and our calculated half-life was in agreement with their observation (81 d). It raised the question of whether adipose tissue vitamin D should be regarded as a slow-release pool that supports long-term functional vitamin D status (i.e. plasma 25(OH)D), and whether massive vitamin D release from adipose tissue, e.g. during fasting, may cause an increase in plasma 25(OH)D and toxicity, as suggested by Connors et al. (1976).

Data in Fig. 1 show that the fifty times higher intake of cholecalciferol for 14 d was unable to increase plasma

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**Fig. 3.** Time-courses of plasma cholecalciferol (a) and 25-hydroxyvitamin D (25(OH)D; (b)) concentrations and subcutaneous (c) and perirenal (d) adipose tissue cholecalciferol contents in fasting (○—○) and ad libitum-fed (□—□) rats. All rats had previously received 37.5 μg cholecalciferol/d from day 0 to day 13. Rats were weighed and samples were taken at 10.00 hours. Data points represent means with their standard errors represented by vertical bars, for a group of six rats. 1 The last cholecalciferol administration (day 13, 22.00 hours) 2 the start of fasting (10.00 hours at days 14 and 98). Mean values for fasting rats were significantly different from those for ad libitum-fed rats: *P < 0.05. For details of procedures, see p. 528.
25(OH)D levels for longer than about 27 d after treatment, and that in this period only small changes occurred in adipose tissue cholecalciferol. Circulating 25(OH)D levels, therefore, seem particularly dependent on the continuous influx of vitamin D from the gastrointestinal tract, at least under the present condition of energy balance and level of vitamin D intake from the diet.

A different situation occurs, however, during negative energy balance and zero dietary vitamin D intake, as indicated by the study of fasting and ad libitum-fed rats. Fasting, in contrast to ad libitum feeding, was associated with high free fatty acid concentrations in plasma and continuing weight loss (Fig. 2). As a result, adipose tissue cholecalciferol content (nmol/g wet weight) increased, apparently because of the preferential loss of triacylglycerols from adipose tissue, as opposed to the loss of cholecalciferol (Fig. 3). Fasting and ad libitum-fed rats showed no differences in plasma cholecalciferol concentrations in the period immediately after cholecalciferol treatment, but 25(OH)D levels of fasting rats were clearly higher. This shows that, despite their zero intake of vitamin D, fasting rats were able to increase their circulating 25(OH)D levels, most probably by vitamin D mobilization from adipose tissue. The zero dietary vitamin D intake by fasting rats was reflected in their lower plasma vitamin D levels during the fasting period at the end of the study. Also these rats developed higher plasma 25(OH)D concentrations, although the difference was obviously less pronounced.

We conclude that orally-administered cholecalciferol rapidly accumulates in adipose tissue and that it is only slowly released in periods of energy balance. Under these conditions circulating 25(OH)D levels may be dependent on dietary vitamin D intake and synthesis in skin. Fasting causes preferential loss of triacylglycerols (and possibly water) from adipose tissue, as opposed to vitamin D, but nevertheless augments plasma 25(OH)D levels under these conditions of zero dietary vitamin D intake. Toxicity because of massive vitamin D release and subsequent 25(OH)D production does not take place. Adipose tissue is likely to act as a ‘buffer to functional vitamin D status’ by preventing unregulated production of 25(OH)D from dietary vitamin D up to a certain extent, and by slowly releasing vitamin D under ad libitum-fed and fasting conditions.

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