Advances in the knowledge of the metabolism of vitamin D

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The secosteroid, vitamin D, functions in calcium homeostasis as a hydroxylated derivative: 1,25-dihydroxy-vitamin D (1,25-DOHD). This molecule becomes localized in the nuclei of the mucosal cells of the small intestine (Lawson, Wilson & Kodicek, 1969) where it appears to induce messenger RNA (Emtage, Lawson & Kodicek, 1973) for proteins which will subsequently increase the transport of Ca by these cells. Thus vitamin D seems to act in a manner similar to the classical steroid hormones of the adrenal cortex and the gonads.

It is notable that the steroid hormones all have oxygen functions at the extremities of the long axis of the molecule. For example, corticosterone, cortisol and aldosterone have 3-keto- and 21-hydroxy-groups while progesterone has keto-groups on both C-3 and C-20. This polar oxygenation enables these steroids to interact, with high affinity, with specific receptor proteins in target cells. The specificity of the interaction is enhanced by the presence of functional groups deployed elsewhere on the steroid structure. In the instance of 1,25-DOHD the polar oxygens are the hydroxy-groups on C-3 and -25 and the group giving the functional specificity is the hydroxyl on C-1.

The hydroxylation of vitamin D

Conversion of vitamin D to its functional state takes place in two stages: 25-hydroxylation in the liver (Ponchon, Kennan & DeLuca, 1969) followed by 1-hydroxylation in the kidney (Fraser & Kodicek, 1970). Hence vitamin D derived from the environment, either as the preformed vitamin in the diet or as a product of the ultraviolet irradiation of 7-dehydrocholesterol in skin tissue, is transported in the blood to the liver, where some of it is metabolized to 25-hydroxy-vitamin D (25-OHD). This reaction occurs in the endoplasmic reticulum of liver cells (Bhattacharyya & DeLuca, 1974) and the modified molecule is secreted back into the circulation, where in most mammals it is carried on a specific α-globulin protein (Edelstein, 1974). The hydroxylation of 25-OHD at C-1 takes place in the mitochondria of kidney cells (Fraser & Kodicek, 1970) and the dihydroxylated derivative passes back into the blood, from which it is removed by cells of the target tissues: the small intestine, bone and the kidney itself.

Since the discovery of this metabolic sequence, the hydroxylation of vitamin D has come to be regarded as an activating process to change a 'biologically inert'
molecule into a completed and now functional form. It is as if vitamin D were a metabolizable nutritional substrate requiring modification in the manner, for example, of the vitamin pyridoxine, which is converted to pyridoxal phosphate, the active coenzyme form. There is, however, a difference in these types of intermediary reactions. Because the function of vitamin D is to regulate Ca metabolism, the formation of the active molecule, 1,25-DOHD, must itself be regulated according to the requirements for maintaining Ca homoeostasis.

*Regulation of the hydroxylation of vitamin D*

After the administration of a single dose of vitamin D to man, the rat or the chicken, it has been demonstrated in many laboratories that, with time, the plasma concentration of the vitamin falls rapidly in comparison with that of its product 25-OHD. Thus with the usual low intake of vitamin D the major vitamin D substance in plasma is 25-OHD. The plasma can therefore be considered to be the body store of substrate for the formation of the active metabolite, 1,25-DOHD. The concentration of 25-OHD in plasma is related to the input of vitamin D and its production does not seem to be regulated according to need. Hence the higher the intake of vitamin D, the greater will be the concentration of 25-OHD in plasma. This can be shown experimentally in vitamin D-deficient chickens given varying doses of cholecalciferol (Table 1): 36 h after a dose of 5 mg cholecalciferol
the plasma 25-hydroxycholecalciferol (25-HCC) level has risen to over 100 μg/l and the cholecalciferol level itself is nearly five times greater again.

Table 1. Concentrations of cholecalciferol and 25-hydroxycholecalciferol (25-HCC) in plasma of 3-week-old, vitamin D-deficient chickens 36 h after a subcutaneous injection of [1,2-3H]cholecalciferol* (Values are for one chicken/dose level)

<table>
<thead>
<tr>
<th>Cholecalciferol dose (μg)</th>
<th>Plasma concentration (μg/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cholecalciferol</td>
</tr>
<tr>
<td>0.5</td>
<td>0.3</td>
</tr>
<tr>
<td>5</td>
<td>0.5</td>
</tr>
<tr>
<td>50</td>
<td>61.5</td>
</tr>
<tr>
<td>500</td>
<td>327.3</td>
</tr>
<tr>
<td>5000</td>
<td>505.5</td>
</tr>
</tbody>
</table>

*As a solution in propylene glycol.

On the other hand, the formation of 1,25-DOHD is very limited. In vitamin D-deficient chickens the concentration of 1,25-DOHD in the intestinal mucosa following a single injection of 2.5 μg cholecalciferol rises to no more than 3 ng/g (Lawson, 1971). However, the concentration in plasma of such birds is less than 300 ng/l. When the activity of 25-HCC-1-hydroxylase is measured in the chicken kidney, it is found to increase during the onset of vitamin D deficiency so that after 3-4 weeks the capacity to produce 1,25-DOHD may be 20 times greater than that of normal chickens (Fraser & Kodicek, 1973). Thus the ability to 1-hydroxylate 25-OHD increases according to the severity of the vitamin D deficiency, but the production of 1,25-DOHD cannot be raised by giving more vitamin D.

The 25-HCC-1-hydroxylase activity in kidney is correlated with a tendency for the extracellular Ca concentration to fall. Since 1,25-DOHD causes increased intestinal absorption of Ca (Kodicek, Lawson & Wilson, 1970; Myrtle & Norman, 1971; Omdahl, Holick, Suda, Tanaka & DeLuca, 1971), mobilization of Ca from bone (Reynolds, 1974) and probably increased tubular reabsorption of Ca in the kidney (Puschett, Fernandez, Boyle, Gray, Omdahl & DeLuca, 1972), then these combined actions of 1,25-DOHD, which raise the extracellular Ca concentration, will inhibit its own production (Fig. 2). Regulation of 25-HCC-1-hydroxylase is mediated primarily by a trophic action of parathyroid hormone (Garabedian, Holick, DeLuca & Boyle, 1972; Rasmussen, Wong, Bikle & Goodman, 1972; Fraser & Kodicek, 1973), but it is very likely that other factors, such as the intracellular concentrations of Ca (Larkins, MacAuley, Rapoport, Martin, Tulloch, Byfield, Matthews & MacIntyre, 1974) and phosphorus (Tanaka & DeLuca, 1973), also influence the enzyme activity.

The two stages in the functional metabolism of vitamin D are therefore fundamentally dissimilar. The 25-hydroxylation reaction proceeds according to the
intake of vitamin D, while the \( \text{r-hydroxylase} \) is firmly regulated in an endocrinological system as an integral part of the Ca homoeostatic mechanism.

**Vitamin D metabolism in normal animals**

Most vitamin D research has been done with vitamin D-deficient animals and has been concerned with the significance of \( 1,25\text{-DOHD} \) in Ca homoeostasis. However, vitamin D metabolites may have a role apart from that of maintaining a constant extracellular Ca concentration. If chickens are given sufficient radioactively labelled cholecalciferol to enable normal growth, then the pattern of vitamin D metabolites can be observed in a situation where there is no stress on the Ca homoeostatic mechanism. In such chickens the \( 25\text{-OHD} \) level in plasma is about \( 10 \) µg/l. This is 5–10 times higher than its concentration in any other tissue. The amount of \( 1,25\text{-DOHD} \) in plasma is about 500 times less than that of \( 25\text{-OHD} \), being about 20 ng/l. Yet this concentration maintains the \( 1,25\text{-DOHD} \) in intestinal mucosa at about 0.4 ng/g, which presumably is sufficient to allow adequate Ca absorption.

Other metabolites of vitamin D are also found in the plasma of these normal chickens, the most notable being \( 24,25\text{-dihydroxy-vitamin D} \) (about 2 µg/l) and \( 25,26\text{-dihydroxy-vitamin D} \) (about 0.7 µg/l). Like \( 25\text{-OHD} \), these dihydroxylated metabolites are more concentrated in plasma than elsewhere. It is possible that they are also concerned with Ca metabolism, but functions have not yet been defined for them.
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