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Vitamin D: beyond boneSylvia Christakos,¹ Martin Hewison,² David G. Gardner,³ Carol L. Wagner,⁴ Igor N. Sergeev,⁵ Erica Rutten,⁶ Anastassios G. Pittas,⁷ Ricardo Boland,⁸ Luigi Ferrucci,⁹ and Daniel D. Bikle¹⁰

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In recent years, vitamin D has been received increased attention due to the resurgence of vitamin D deficiency and rickets in developed countries and the identification of extraskeletal effects of vitamin D, suggesting unexpected benefits of vitamin D in health and disease, beyond bone health. The possibility of extraskeletal effects of vitamin D was first noted with the discovery of the vitamin D receptor (VDR) in tissues and cells that are not involved in maintaining mineral homeostasis and bone health, including skin, placenta, pancreas, breast, prostate and colon cancer cells, and activated T cells. However, the biological significance of the expression of the VDR in different tissues is not fully understood, and the role of vitamin D in extraskeletal health has been a matter of debate. This report summarizes recent research on the roles for vitamin D in cancer, immunity and autoimmune diseases, cardiovascular and respiratory health, pregnancy, obesity, erythropoiesis, diabetes, muscle function, and aging.

Keywords: vitamin D; cancer; immunity; pregnancy; obesity; diabetes; pulmonary disease; muscle; aging; cognitive function

Introduction

Traditionally, vitamin D has been considered almost exclusively in the context of its role in calcium homeostasis. Vitamin D can either be taken in the diet, largely through fortification of dairy products, or synthesized in the skin from 7-dehydrocholesterol upon exposure to UV irradiation. Whether ingested or synthesized, vitamin D is transported to the liver, where it is hydroxylated at position 25 to form 25-hydroxyvitamin D (25(OH)D₂ or 25(OH)D₃), the major circulating form of vitamin D. Next, 25(OH)D₃ is transported to the kidney, where it is hydroxylated at position 1 by the enzyme CYP27B1 to form 1,25-dihydroxyvitamin D (1,25(OH)₂D₃), the most active form of vitamin D, which is then transported to target tissues, where it functions like a steroid, binding to the vitamin D receptor (VDR).

The VDR heterodimerizes with the retinoid X receptor (RXR), and the VDR/RXR complex binds to VDR-responsive elements found in or around target genes and, in association with various co-activators, results in the transcription of target genes. When there is a need to increase blood calcium levels (e.g., during growth or pregnancy), 1,25(OH)₂D₃ acts in the intestine to increase calcium absorption. If this increased intestinal absorption is insufficient to restore normal calcium levels, 1,25(OH)₂D₃ works in concert with the parathyroid hormone (PTH) in the kidney to promote calcium reabsorption from the distal tube, and in the skeletal system to release calcium from bones.

While it was long held that vitamin D acted only at the intestine, kidney, and skeleton, and that its function was limited to calcium homeostasis, the possibility of extraskeletal effects has been

considered for decades as a result of the discovery of the VDR in tissues that have no involvement in calcium homeostasis (e.g., skin, placenta, pancreas, breast, prostate and colon cancer cells, and activated T cells). Discovery of the VDR and CYP27B1 in these tissues led to exploration of the roles and mechanisms of vitamin D function in each. On September 21, 2012, the Abbott Nutrition Health Institute and the Sackler Institute for Nutrition Science at the New York Academy of Sciences sponsored a conference, "Vitamin D: Beyond Bone," that gathered researchers investigating vitamin D in a wide range of tissues and diseases.

Nonclassical effects of vitamin D

Vitamin D: beyond bone

Sylvia Christakos (UMDNJ-New Jersey Medical School) began the meeting by summarizing the metabolism and activity of vitamin D and introducing the emerging awareness of roles for vitamin D beyond calcium homeostasis and bone metabolism. Evidence in the laboratory, including the use of animal models, indicates that $1,25(\text{OH})_2\text{D}_3$ generates a number of extraskelatal effects, including inhibition of cancer progression, effects on the cardiovascular system and skin, modulation of innate immunity with subsequent killing of bacteria, and inhibition of certain autoimmune diseases (reviewed in Ref. 1). For example, in rats treated with the chemical carcinogen *N*-methyl *N*-nitrosourea (NMU), $1,25(\text{OH})_2\text{D}_3$ inhibits the progression of breast tumors. In addition, when $1,25(\text{OH})_2\text{D}_3$ is given prior to NMU, tumor incidence is prevented or reduced.¹ $1,25(\text{OH})_2\text{D}_3$ also reduces the incidence and severity of prostate neoplasia in a mouse model (Ndx 3.1; *Pten* mutant mouse), and $1,25(\text{OH})_2\text{D}_3$ has tumor inhibitory activity in a mouse model of colorectal adenoma (*Apc*^{min}). In order to determine mechanisms involved in inhibition of breast tumor growth, Christakos' lab showed that C/EBP α , a transcription factor that has been shown to play a critical role in growth arrest of other cell types, is induced by $1,25(\text{OH})_2\text{D}_3$ in MCF-7 human breast cancer cells.² C/EBP α was found to induce transcription of the vitamin D receptor in MCF-7 cells.² Since the levels of the VDR correlate with the antiproliferative effects of $1,25(\text{OH})_2\text{D}_3$, and since it has been suggested that C/EBP α can be considered a potential tumor suppressor, these findings suggest mechanisms whereby $1,25(\text{OH})_2\text{D}_3$ may

act to inhibit growth of breast cancer cells. These findings also identify C/EBP α as a $1,25(\text{OH})_2\text{D}_3$ target in breast cancer cells and provide evidence for C/EBP α as a candidate for breast cancer treatment.²

With regard to autoimmune diseases, $1,25(\text{OH})_2\text{D}_3$ has been shown to suppress type 1 diabetes in the non-obese diabetic (NOD) mouse model, to suppress experimental autoimmune encephalomyelitis (EAE) (a mouse model of multiple sclerosis (MS)), and to suppress mouse models of inflammatory bowel disease and systemic lupus erythematosus.¹ Recent studies from Christakos' lab have shown that inhibition of EAE is associated with inhibition of interleukin (IL)-17, a cytokine that plays a critical role in numerous inflammatory conditions and autoimmune diseases including MS. The mechanism of $1,25(\text{OH})_2\text{D}_3$ suppression of IL-17 was found to be transcriptional and to involve blocking of nuclear factor for activated T cells (NFAT, which is important for T cell receptor-mediated transcriptional regulation of IL-17), recruitment of histone deacetylase to the IL-17 promoter, and sequestration of Runt-related transcription factor 1 (Runx1) by the VDR.³ $1,25(\text{OH})_2\text{D}_3$ was also found to have a direct effect on the induction of Foxp3, a transcription factor that associates with NFAT and Runx1 for transcriptional repression.³ These results describe novel mechanisms and new concepts with regard to vitamin D and the immune system and suggest therapeutic targets for the control of autoimmune diseases.

Unlike the association between vitamin D deficiency and rickets, causal links between vitamin D deficiency and specific extraskelatal diseases have yet to be identified. However, the evidence in the laboratory of beneficial effects of $1,25(\text{OH})_2\text{D}_3$ beyond bone is compelling (summarized in Fig. 1). Findings in animal models may suggest mechanisms involving similar pathways in humans that could lead to the identification of new therapies.

Vitamin D in immune function and disease prevention

Martin Hewison (the David Geffen School of Medicine, University of California) detailed one of the most prominent of the so-called nonclassical effects of vitamin D: its ability to act as a potent modulator of human immune responses. Evidence for this initially stemmed from two observations.

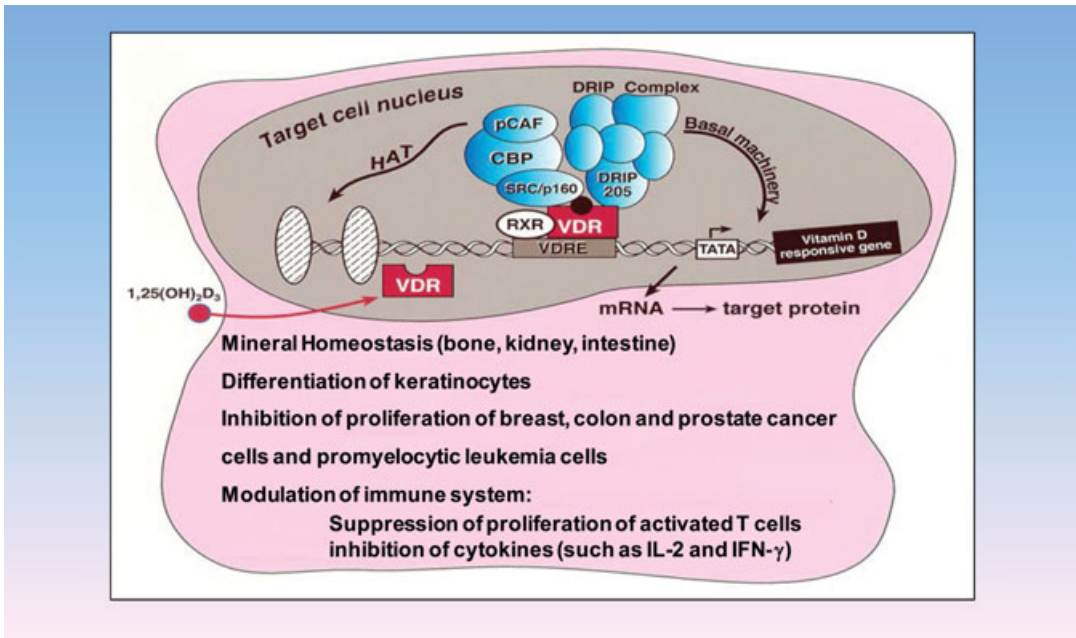


Figure 1. Genomic mechanism of vitamin D action. Mechanism of action of $1,25(\text{OH})_2\text{D}_3$ in target cells. The VDR heterodimerizes with the RXR. After interaction with the VDRE (vitamin D response element), transcription proceeds through the interaction of the VDR with coactivators and with the transcription machinery. The histone acetyltransferase (HAT) activity-containing complex (SRC/p160 and CBP), the DRIP complex, and additional coactivators not shown (including specific methyltransferases) are recruited by liganded VDR. $1,25(\text{OH})_2\text{D}_3$ is known to maintain calcium homeostasis and to affect numerous other cell types. Effects on other cell systems, including modulation of the immune system and inhibition of proliferation of cancer cells, are discussed. With permission from Christakos.⁶⁰

First, many cells from both the innate and adaptive immune systems express the VDR. Second, antigen cells from the innate immune system, such as macrophages or dendritic cells (DCs), also express the vitamin D activation enzyme 1α -hydroxylase, also known as CYP27B1. As such, these cells are able to convert precursor $25(\text{OH})\text{D}_3$, the major circulating form of vitamin D, to active $1,25(\text{OH})_2\text{D}_3$ that can then induce responses in the cells by binding to their VDRs and promoting transcriptional regulation.

This localized intracrine mechanism appears to be central to two key features of immune function: innate antibacterial activity and the presentation of antigen to cells from the adaptive immune system such as T lymphocytes (T cells). In macrophages and monocytes, cellular sensing of pathogens, such as *Mycobacterium tuberculosis*, by pattern recognition receptors, such as Toll-like receptors, enhances expression of CYP27B1 and the VDR, thereby increasing intracrine synthesis and activity of $1,25(\text{OH})_2\text{D}_3$.⁴ The resulting effects, including enhanced expression of the antibacterial

proteins cathelicidin and β -defensin 2, and enhanced formation of autophagosomes, facilitate improved bacterial killing in a vitamin D-dependent fashion. Vitamin D insufficiency or deficiency may therefore impair innate immune responses and predispose individuals to infection, and low serum concentrations of $25(\text{OH})\text{D}_3$ have been linked to infectious diseases such as tuberculosis. However, it is important to recognize that similar antibacterial responses to vitamin D have also been reported for other cell types within tissues, such as the skin, lungs, GI tract, and placenta, that may broadly be termed *barrier sites*. Thus, it is possible that vitamin D deficiency has a generalized detrimental impact on antibacterial responses and may therefore play a role in many types of infectious diseases.⁵ An important note by Hewison is that current data suggest that this particular immune response to vitamin D is restricted to primates, underlining a potential role for vitamin D in the evolution of primate immunity, but also limiting the scope of further *in vivo* studies to explore this activity.

The immunomodulatory effects of vitamin D also involve the adaptive immune system. Intracrine synthesis of $1,25(\text{OH})_2\text{D}_3$ by DCs decreases their maturation, thereby suppressing antigen presentation and decreasing T cell proliferation.⁶ Recent studies have shown that intracrine expression and activity of CYP27B1 by DCs is particularly effective in supporting the generation of tolerogenic regulatory T (T_{reg}) cells, while simultaneously suppressing inflammatory IL-17-expressing T cells (Th17 cells).⁷ In this way, localized metabolism of $25(\text{OH})\text{D}_3$ may play a pivotal role in prevention and/or treatment of autoimmune diseases, such as multiple sclerosis, rheumatoid arthritis, and inflammatory bowel disease. As with antibacterial responses to infection, it has been proposed that adaptive immune responses may be compromised in patients with low serum concentrations of $25(\text{OH})\text{D}_3$, and association studies have shown links between autoimmune disease and vitamin D deficiency. The extent to which vitamin D deficiency is a cause of immune disease has yet to be demonstrated and will require new prospective clinical trials of vitamin D supplementation. Hewison stated that it will be important to define whether enhanced vitamin D-mediated immunity prevents immune disease or whether it can be used as therapy for these diseases. Moreover, it is possible that some diseases, such as leprosy and HIV infection, may be associated with dysregulation of the core vitamin D intracrine immune mechanism, and may therefore require higher circulating levels of $25(\text{OH})\text{D}_3$ to reestablish normal immune responses. These and other facets of vitamin D and immune function will be important research challenges for the vitamin D community.

Vitamin D in the heart and vascular system

David G. Gardner (University of California, San Francisco) expanded the discussion of nonclassical effects of vitamin D to include evidence of a significant role for vitamin D in the cardiovascular system. While the heart and vasculature have only recently been identified as potential targets of vitamin D action, a growing body of evidence suggests that the activated VDR plays an important role in regulating cardiovascular function.⁸ First, the VDR and the ligand-generating 25 -hydroxyvitamin D_3 1 - α -hydroxylase have been shown to be present in cardiac myocytes, cardiac fibroblasts, vascular

smooth muscle cells, and vascular endothelial cells. Second, vitamin D deficiency in rodents has been shown to elicit increases in blood pressure and cardiac hypertrophy. Third, $1,25(\text{OH})_2\text{D}_3$ and its analogs have been shown to reverse agonist-induced myocyte hypertrophy *in vitro* and endogenous cardiac hypertrophy in the Dahl S rat, the spontaneous hypertensive rat (SHR), the spontaneous hypertensive heart failure-prone rat (SHR-HF), and the 5/6 nephrectomy model of chronic renal failure, *in vivo*. Fourth, a variety of cardiovascular disorders including congestive heart failure, coronary artery disease, and peripheral vascular disease have been linked to reduced circulating levels of $25(\text{OH})\text{D}_3$. Fifth, and perhaps most important, a mouse with a complete deletion of the VDR gene demonstrates both hyperreninemic hypertension and cardiac hypertrophy.⁹

Gardner and colleagues examined the ability of paricalcitol, a bioactive analog of $1,25(\text{OH})_2\text{D}_3$, to prevent cardiac hypertrophy in rats infused with moderate doses of angiotensin II (800 ng/kg/min) over a 14-day period. Infusion of angiotensin II led to increased blood pressure, increased myocyte hypertrophy, increased expression of the hypertrophy-sensitive fetal gene program (i.e., atrial natriuretic peptide, B-type natriuretic peptide, and alpha skeletal actin gene expression), and increased cardiac interstitial fibrosis with augmented procollagen 1 and 3 expression. In each case, coadministration of paricalcitol (intraperitoneal injection of 300 ng/kg every 48 h), resulted in partial reversal of the angiotensin II effects.¹⁰

In an effort to define the role of the VDR in the myocardium, Gardner and colleagues created a mouse with a selective deletion of the fourth exon of the murine *Vdr* gene in cardiac myocytes. This mouse displayed an increase in myocyte size, left ventricular weight/body weight, and hypertrophy-dependent fetal gene expression compared with control littermates. There was no demonstrable increase in interstitial fibrosis. The mouse also demonstrated a reduction in end-diastolic and end-systolic volume by echocardiography and increased expression of modulatory calcineurin inhibitory protein 1 (MCIP1), a direct downstream target of calcineurin and NFAT that has been linked to the development of cardiac hypertrophy. Gardner's group showed that isoproterenol treatment of neonatal rat cardiac myocytes *in vitro* resulted

in myocyte hypertrophy and increased MCIP1 expression. Coadministration of $1,25(\text{OH})_2\text{D}_3$ resulted in a dose-dependent reduction in MCIP1 expression.¹¹

Vitamin D and the VDR have also been implicated in the support of normal endothelial function. Endothelial dysfunction, regarded by many as a harbinger of cardiovascular disease, has been linked to a number of disorders that are also associated with low circulating levels of $25(\text{OH})\text{D}_3$. In some cases, treatment of these disorders with exogenous vitamin D or vitamin D metabolites has been shown to restore endothelial function. To establish a link between the VDR and endothelial function, Gardner and colleagues created a mouse with selective deletion of the *Vdr* gene in vascular endothelial cells. This mouse displays many of the phenotypic features normally associated with endothelial dysfunction.

Collectively, these data demonstrate that the vitamin D system—specifically, the ligand-bound VDR—plays an important role in the maintenance of cardiovascular health and suggest that vitamin D repletion may have important public health implications in controlling the prevalence of, and morbidity associated with, cardiovascular disease. Of equivalent importance, therapy with high potency vitamin D metabolites or their analogs may subservise an important role in the management of these diseases.

Vitamin D during pregnancy and lactation

Vitamin D as a prohormone has effects that extend beyond calcium metabolism and homeostasis throughout life but which are most vivid during pregnancy when vitamin D metabolism is disengaged from its usual constraints.¹² At no other time during the lifecycle is $25(\text{OH})\text{D}_3$ linked directly to $1,25(\text{OH})_2\text{D}_3$ production, the latter rising more than 2.5 times above nonpregnant levels.¹² This elevation of $1,25(\text{OH})_2\text{D}_3$ likely serves the purpose of immune regulation, though there have been only a few studies that have evaluated the effect of vitamin D status on immune function during pregnancy to support this hypothesis. In addition, few randomized controlled trials have been conducted in pregnant women to determine optimal vitamin D status.^{13–15}

Carol L. Wagner (Medical University of South Carolina) discussed two recent studies performed

by her group in collaboration with Bruce Hollis's group—the National Institute of Child Health and Human Development (NICHD) Vitamin D Pregnancy Trial¹² and the Thrasher Research Fund Vitamin D Community-Based Supplementation Trial¹⁶—that indicate that 400 IU vitamin D, the amount found in most prenatal vitamins, is inadequate and that 4000 IU/day is necessary to optimize $1,25(\text{OH})_2\text{D}_3$ production, achieved when total circulating $25(\text{OH})\text{D}_3$ is at least 40 ng/mL (Fig. 1).¹² When analyzed by maternal $25(\text{OH})\text{D}_3$ concentration at various time points—i.e., baseline, by trimester, throughout pregnancy as mean, median, and area under the curve, or at delivery—the group taking 4000 IU/day attained optimal $1,25(\text{OH})_2\text{D}_3$ conversion from $25(\text{OH})\text{D}_3$ throughout pregnancy at considerably higher rates than groups taking lower doses. Both studies provide evidence that 4000 IU/day is not only safe but is associated with fewer adverse events of pregnancy when taken together as comorbidities of pregnancy, compared with the lower dose groups.

Comorbidities of pregnancy appear to be directly linked to lower $25(\text{OH})\text{D}_3$, and, conversely, increasing $25(\text{OH})\text{D}_3$ appears to afford protection to the mother and developing fetus, with lower risk of preterm labor, preterm birth, or infection for each 10 ng/mL increase in total circulating $25(\text{OH})\text{D}_3$ concentration.¹⁶ Such findings suggest a simple approach to reduce morbidities during pregnancy. Yet, before such findings are fully embraced, the mechanism(s) of action of vitamin D and its protective effect during pregnancy must be better delineated. Additional research that addresses the impact of vitamin D on immune homeostasis during pregnancy and its effects during early infancy and later childhood becomes essential.

These findings extend to lactation and early infancy. It was thought for decades that human milk was minimally sufficient in vitamin D;¹⁷ yet, preliminary results of the recently completed NICHD vitamin D supplementation trial during lactation show that when a mother is vitamin D replete, with an intake of 6400 IU vitamin D₃/day, breast milk is replete and a breastfeeding infant has excellent vitamin D status without infant supplementation that is comparable to combined maternal and infant supplementation of 400 IU vitamin D₃/day.¹⁸ The implications of safely achieving vitamin D sufficiency for both mother and breastfeeding infant

solely through maternal supplementation is just beginning to be understood and will likely challenge researchers for decades to come.

Vitamin D and obesity

Igor N. Sergeev (South Dakota State University) discussed the possibility of integrating vitamin D supplementation with current strategies in the prevention and treatment of obesity. The induction of adipocyte death through apoptosis is emerging as a promising strategy for addressing obesity.^{19,20} Increased adipose tissue mass is the result of both hypertrophy (an increase in adipocyte size) and hyperplasia (an increase in adipocyte number), and once adipocytes achieve a maximum size, further increase in adipose tissue mass involves an increase in adipocyte number. Thus, weight loss can result from not only a decrease in adipocyte size but also adipocyte number. Even a small increase in the amount of adipocyte apoptosis will prevent excessive accumulation of adipose tissue and can result in a significant loss of adipose tissue mass over time. Therefore, removal of adipocytes through apoptosis reduces body fat and can help in long-lasting maintenance of weight loss.

The effects of the hormonal form of vitamin D, 1,25(OH)₂D₃, on apoptotic cell death are mediated via multiple signaling pathways that involve common regulators and effectors converging on cellular Ca²⁺.^{19,21} Sergeev and colleagues have shown that the critical characteristic of the apoptotic Ca²⁺ signal is a sustained increase in the concentration of intracellular Ca²⁺, reaching elevated but not cytotoxic levels.²² However, 1,25(OH)₂D₃-regulated, Ca²⁺-dependent apoptotic molecular targets have not been identified in adipose tissue. Thus, Sergeev and colleagues have been investigating the mechanism by which 1,25(OH)₂D₃ regulates apoptosis in adipocytes. Their results have demonstrated that 1,25(OH)₂D₃ induces, in a concentration- and time-dependent fashion, a sustained, prolonged increase in concentration of intracellular Ca²⁺ (the apoptotic Ca²⁺ signal) in mature mouse 3T3-L1 adipocytes.²³ The 1,25(OH)₂D₃-induced increase in cellular Ca²⁺ is associated with activation of Ca²⁺-dependent μ -calpain and Ca²⁺/calpain-dependent caspase-12.²³ Activation of these proteases is sufficient to effect morphological and biochemical changes consistent with apoptosis. The 1,25(OH)₂D₃-induced

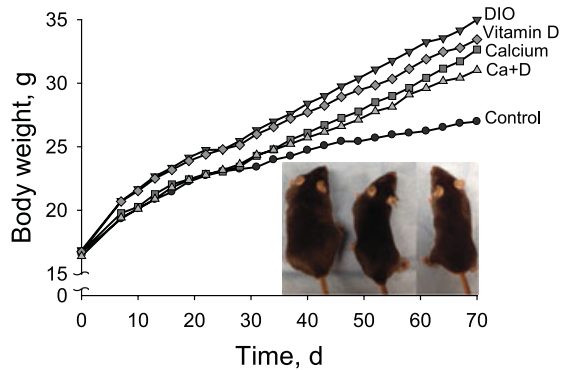


Figure 2. Body-weight gain in mice fed high-fat diets with increased levels of Ca and vitamin D. Weight matched mice were randomly assigned to the experimental high-fat (HF) diets containing 60% energy as fat (the normal control diet contained 10% energy as fat). The treatment groups on HF diet were high-calcium (Ca diet, 1.2% Ca), high-vitamin D₃ (D diet; vitamin D₃ intake 10 times higher than the recommended level of 1000 IU/kg) and high-Ca plus high-D₃ (Ca + D diet). Data are means for each group and time point; *n* = 7–8 per group from week 1 to week 10. The insert shows (from left to right) mice from the DIO, control, and Ca + D groups at week 10. Data are unpublished and from work in progress.

increase in cellular Ca²⁺ is also associated with reduced lipid accumulation in mature adipocytes.²³ In preliminary but ongoing studies, a murine diet-induced obesity (DIO) model is being used to evaluate the role of vitamin D in adiposity. DIO mice (C57BL/6J) fed a high-vitamin D₃ diet, in particular a high-vitamin D₃ plus high-calcium diet, show decreased body and fat weight gain (Fig. 2; unpublished data) and improved markers of adiposity and vitamin D status (plasma concentrations of glucose, insulin, adiponectin, 25(OH)D₃, 1,25(OH)₂D₃, and PTH), but an increased plasma Ca²⁺. High vitamin D₃ and calcium intake is associated with activation of the Ca²⁺-dependent apoptotic proteases calpain and caspase-12 in adipose tissue of DIO mice. These preliminary findings suggest that the 1,25(OH)₂D₃-induced cellular Ca²⁺ signal can act as an apoptotic initiator that directly recruits Ca²⁺-dependent apoptotic effectors capable of executing apoptosis in adipose tissue. The preliminary results also suggest that high vitamin D and calcium intake decreases body and fat weight gain in diet-induced obesity, and that the potential mechanism of these effects involves activation of a Ca²⁺-mediated apoptotic pathway in adipose tissue (specifically, normalization of the activities of apoptotic proteases

Ca^{2+} -dependent calpain and Ca^{2+} /calpain-dependent caspase-12 that are reduced in obesity). Targeting Ca^{2+} signaling and the vitamin D/ Ca^{2+} -dependent calpains and caspases in adipocytes with vitamin D supplementation may therefore be an effective and affordable approach for chemoprevention and treatment of obesity. Additional studies are warranted to evaluate this approach from a safety point of view and to identify the optimal levels of calcium and vitamin D intakes in obesity.

Vitamin D and lung function in patients with chronic obstructive pulmonary disease

Erica Rutten (Ciro+, Centre of Expertise for Chronic Organ Failure, Netherlands) discussed her work investigating the association between plasma vitamin D levels and lung function in chronic obstructive pulmonary disease (COPD) patients. COPD is characterized by a persistent and, usually, progressive airflow limitation and is associated with an enhanced local inflammatory response. By 2030, COPD is predicted to become the fourth leading cause of death worldwide, and therefore accounts for about 8% of total deaths. Today, in addition to lung function impairment, COPD is recognized as a systemic disease with multiple comorbidities, such as osteoporosis, cardiovascular disease, renal impairment, and psychological disorders. In relation to the systemic pathology of COPD the role of vitamin D has generated interest, as vitamin D deficiency is often present in patients with COPD,²⁴ dependent on the severity of COPD.²⁵ Indeed, the prevalence of vitamin D deficiency, defined as plasma 25(OH) D₃ concentrations below 20 ng/ml (50 nmol/L), was 58% in a cross-sectional study of 151 COPD patients entering pulmonary rehabilitation during summer.²⁶ In the Bergen COPD Cohort Study, the prevalence of vitamin D deficiency was indeed higher in the COPD patients than in the control subjects, independent of season, age, smoking, comorbidity, and body mass index.²⁷ Several potential explanations for the increased risk of vitamin D deficiency in patients with COPD include poor diet, less outdoor activity and thus less exposure to sunlight, accelerated skin aging due to smoking, renal dysfunction, and treatment with corticosteroids. Vitamin D is related to skeletal health in patients with COPD, and vitamin D deficiency is associated with low bone mineral density.^{26,28} Additionally, low vitamin D has been related to decreased exercise performance^{26,29} and

exacerbation rate³⁰ in COPD patients. And while vitamin D concentration has been related to lung function in the general population, the relation between lung function and vitamin D concentration has yet to be investigated in COPD patients only.

Rutten and colleagues performed a secondary analysis of the publication by Romme *et al.*²⁶ to investigate whether there is an independent association between lung function parameters and plasma vitamin D levels in a group of 151 COPD patients admitted for pulmonary rehabilitation at Ciro+. Plasma 25(OH)D₃ levels, body mass index (BMI; weight in kg/(height in m)²), and lung function parameters (forced expiratory volume in 1s (FEV1), forced vital capacity (FVC), and diffusing capacity of the lung (DLCO)) were measured. The study group comprised 57% males, with mean age 65 ± 9 years, mean FEV1 47.2 ± 17.9% predicted, mean FVC 94.6 ± 20.6% predicted, and mean DLCO 53.2 ± 19.7% predicted; 58% of the patients were vitamin D deficient (vitamin D concentration <20 ng/ml). Pearson correlation coefficient showed significant correlations between plasma vitamin D concentration and FEV1 ($r = 0.31$, $P < 0.01$), FVC ($r = 0.29$, $P < 0.01$) and DLCO ($r = 0.20$, $P = 0.01$). After stratification for vitamin D deficiency, the correlations persisted only in the patients with vitamin D deficiency: FEV1 ($r = 0.37$, $P < 0.01$; Fig. 3, data are from an unpublished study), FVC ($r = 0.25$, $P < 0.01$), and DLCO ($r = 0.32$,

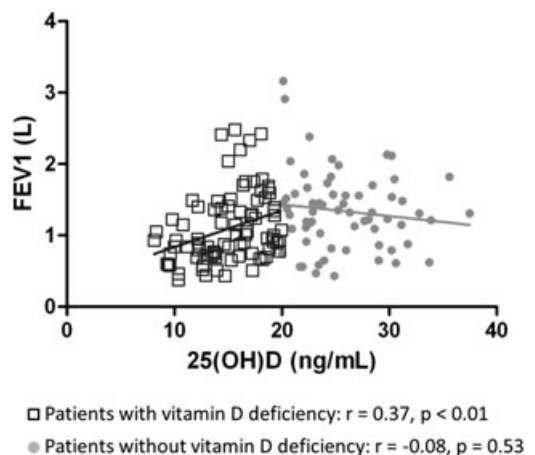


Figure 3. Lung function and plasma vitamin D levels in a group of 151 COPD patients. FEV1, forced expiratory volume in 1 s. These data are from an unpublished secondary analysis by Rutten and colleagues.

$P = 0.02$). Multivariate regression analyses revealed that, after correction for age, gender, and BMI, vitamin D concentration remained independently associated with FEV1 ($\beta = 0.26$, $P < 0.01$) and DLCO ($\beta = 0.25$, $P < 0.01$), and there was a trend for FVC ($\beta = 0.15$, $P < 0.08$) in the subjects with vitamin D deficiency. From their unpublished secondary analysis, Rutten and colleagues conclude that there is evidence suggesting that vitamin D plays a role in the lung pathology of patients with COPD, which requires further investigation.

Impact of cholecalciferol repletion on erythropoietin requirements in hemodialysis patients

Lily Li (Cleveland Clinic Lerner College of Medicine of Case Western Reserve University and Mount Sinai School of Medicine) presented pilot data from a randomized control trial investigating the impact of cholecalciferol repletion on erythropoietin requirements in vitamin D-deficient hemodialysis patients. Li mentioned that immune cells contain the machinery needed to convert $25(\text{OH})\text{D}_3$ to $1,25(\text{OH})_2\text{D}_3$, and that there is evidence that this local production of $1,25(\text{OH})_2\text{D}_3$ has immunomodulatory properties. In a setting of decreased $25(\text{OH})\text{D}_3$, the absence of $1,25(\text{OH})_2\text{D}_3$ production may enhance local inflammation.

Among patients with end-stage renal disease (ESRD) on hemodialysis, $25(\text{OH})\text{D}_3$ deficiency is common, reported at greater than 80%.³² ESRD patients are commonly treated with $1,25(\text{OH})_2\text{D}_3$ for prevention of secondary hyperparathyroidism and bone loss. However, $25(\text{OH})\text{D}_3$ is rarely repleted within this population, and the impact of $25(\text{OH})\text{D}_3$ deficiency remains obscure. Anemia is also associated with ESRD and is the result of reduced erythropoietin production due to kidney disease as well as iron deficiency and inflammation-mediated abnormalities in iron metabolism. In this population, anemia not only causes impaired quality of life but also left ventricular hypertrophy, myocardial infarction, and increased mortality. Patients with ESRD-associated anemia are commonly treated with erythropoiesis-stimulating agents (ESA), which are very effective in the majority of the population. However, a subset of these patients develop erythropoietin hyporesponsiveness, which is thought to be caused by inadequate iron mobilization in the setting of inflammation, medi-

ated in part by hepcidin, a small molecule produced by hepatocytes that is known to inhibit macrophage iron release and intestinal iron absorption. Since hepcidin production appears to be induced by IL-6, and $25(\text{OH})\text{D}_3$ has been shown to suppress IL-6 production in monocytes,³³ Li's group developed the hypothesis that in hemodialysis patients deficiency of $25(\text{OH})\text{D}_3$, which is required for local production of $1,25(\text{OH})_2\text{D}_3$ in immune cells, leads to dysregulation of innate immunity and inflammation (e.g., enhanced IL-6 production), which alters iron metabolism and contributes to erythropoietin resistance.

They have initiated a randomized controlled trial of $25(\text{OH})\text{D}_3$ repletion in vitamin D-deficient hemodialysis patients to evaluate efficacy and safety and the effects on inflammation and ESA requirements. Patients are given either no repletion (standard of care) or 50,000 IU/week of cholecalciferol for six weeks, or until $25(\text{OH})\text{D}_3$ levels exceeded 35 ng/mL, and 10,000 IU/week afterwards. Their preliminary results show that $25(\text{OH})\text{D}_3$ repletion was both safe and effective, with $25(\text{OH})\text{D}_3$ levels increasing significantly in the treatment arm and no patients experiencing hypercalcemia or other adverse effects. In addition, patients in the treatment arm exhibit a decrease in ESA requirement at six months. Li and colleagues have isolated monocytes from patients and examined expression of a number of genes related to vitamin D metabolism and immune responses. Their observations include a decrease in IL-6 expression following $25(\text{OH})\text{D}_3$ repletion at six months. These preliminary data indicate that $25(\text{OH})\text{D}_3$ repletion treatment is safe, effective, and may result in lower ESA requirements, a result with significant economic implications.

Translating epidemiological data into policy and clinical applications

Vitamin D for diabetes

Anastassios G. Pittas (Tufts Medical Center) highlighted the potentially important role of vitamin D in a variety of non-skeletal medical conditions, including type 2 diabetes. To systemically appraise the available evidence on the role of vitamin D in diabetes, Pittas reviewed the original Bradford Hill criteria, which have been subsequently modified to include three general categories (mechanistic studies, direct evidence, and parallel evidence),³⁴

and then applied these criteria to the available evidence.

From a biological perspective, the hypothesis that vitamin D may be a determinant of diabetes risk is plausible, as both impaired insulin secretion and action have been reported with vitamin D insufficiency. The effect of vitamin D may be direct, as supported by the expression of the VDR and the local production of $1,25(\text{OH})_2\text{D}_3$ in pancreatic β cells, or indirect via its role in regulating calcium homeostasis and calcium flux through cell membranes.

In humans, the role of vitamin D in type 2 diabetes is suggested by a large number of cross-sectional studies that have consistently found an inverse association between vitamin D status and prevalent hyperglycemia, including a reported seasonal variation in the control of glycemia in patients with diabetes, being worse in the winter when hypovitaminosis D is more prevalent.³⁵ In the majority of longitudinal observational studies, lower vitamin D status is also associated with increased risk of incident type 2 diabetes. In the most recent systematic review of observational studies, Song *et al.*³⁶ reported a 38% relative risk reduction in incident diabetes among individuals with the highest versus the lowest category of blood $25(\text{OH})\text{D}_3$ concentration. In dose-response analyses, the risk of type 2 diabetes was reduced by 4% for each 4 ng/mL increment in blood $25(\text{OH})\text{D}_3$ concentration.

However, Pittas cautioned that one should resist making inferences on the basis of observational studies because of the possibility of confounding issues, which may be especially true when assessing vitamin D status, because while $25(\text{OH})\text{D}_3$ concentration is an excellent marker of good health many of the variables that contribute to $25(\text{OH})\text{D}_3$ concentration (e.g., skin pigmentation, physical inactivity, aging, unhealthy dietary patterns) are also risk factors for the development of diabetes. Therefore, evidence from intervention studies—which is the most direct evidence—is critically important before one can conclude that vitamin D has a role in the prevention or treatment of diabetes. The results from small clinical trials and *post hoc* analyses of larger trials on the effect of vitamin D supplementation on diabetes-related parameters have been inconclusive, although vitamin D appears to have beneficial effects in persons at risk for diabetes.^{36–39} However, no firm conclusions can be drawn from the available intervention studies because of several limitations:

(1) nearly all studies were underpowered; (2) the minority of studies were designed specifically for glycemic outcomes; (3) most studies did not report use of diabetes medications at baseline or during the study; (4) some trials used large infrequent doses of vitamin D, which may be metabolized differently compared to daily doses and may provide either no benefit or result in an unfavorable benefit/risk ratio.

Pittas concluded that although the observation studies strongly suggest an important role for vitamin D in type 2 diabetes, and such a role is biologically plausible, there is lack of evidence from intervention studies to support the contention that type 2 diabetes can be improved or prevented by raising $25(\text{OH})\text{D}_3$ concentration. On numerous earlier occasions, encouraging findings from observational studies were not confirmed by well-designed clinical trials (e.g., hormone replacement therapy, β -carotene, vitamins C and E, and folic acid) and prevailing clinical practice was overturned. Therefore, confirmation of a potential beneficial effect of vitamin D in type 2 diabetes is needed in trials conducted in well-defined populations (e.g., pre-diabetes and early diabetes) specifically designed to test the compelling but yet unproven hypothesis that vitamin D status is a direct contributor to the pathogenesis of type 2 diabetes.

Molecular aspects of the role of vitamin D in muscle

Ricardo Boland (Universidad Nacional del Sur, Argentina) discussed molecular aspects of the role of vitamin D in skeletal muscle health and function. This is a topic of clinical significance because myopathy characterized by proximal muscle weakness and atrophy is a common symptom in vitamin D deficiency states.⁴⁰ The loss of muscular strength and a reduction in muscle mass increases the risk of falls and leads to fractures. It has been reported that administration of the hormonal vitamin D metabolite $1,25(\text{OH})_2\text{D}_3$ decreases the number of falls by improving muscle strength in elderly subjects.⁴¹ Consistent with this observation, there is evidence that $1,25(\text{OH})_2\text{D}_3$ regulates muscle growth and development and contractility.⁴⁰ Moreover, the presence of the VDR in avian, murine, and human skeletal muscle has been demonstrated.^{40,42}

As in other target cells, $1,25(\text{OH})_2\text{D}_3$ elicits long-term and short-term responses in skeletal muscle that involve genomic and non-genomic modes

of actions, respectively. In the first, more classical mechanism, the hormone stimulates muscle cell proliferation and differentiation through nuclear VDR-mediated gene transcription, expressed by increased myoblast DNA synthesis followed by the induction of muscle specific myosin- and calcium-binding proteins. The non-genomic effects of $1,25(\text{OH})_2\text{D}_3$ are involved in the fast regulation of the calcium messenger system and growth-related signal transduction pathways in skeletal muscle cells. The hormone interacts with a membrane receptor, which leads to stimulation of adenylyl cyclase and phospholipases C, D, and A2 and activation of MAPK cascades.⁴³ Boland pointed out that there is a wealth of evidence indicating that $1,25(\text{OH})_2\text{D}_3$ mainly acts in muscle through these membrane-initiated events and that the VDR plays a role in the activation of intracellular signaling pathways.⁴⁴

Boland then focused his presentation on the signal transduction mechanisms of $1,25(\text{OH})_2\text{D}_3$ in skeletal muscle. $1,25(\text{OH})_2\text{D}_3$ -induced transmembrane activation of adenylyl cyclase/cAMP/PKA and PLC/DAG plus IP_3 /PKC regulates Ca^{2+} influx mediated by voltage-dependent Ca^{2+} channels (VDDC).⁴³ Boland's group showed that, in muscle cells, the hormone also modulates Ca^{2+} influx through SOC channels that are activated by the PLC-mediated Ca^{2+} release from intracellular stores via IP_3 —a process known as capacitative calcium entry (CCE).⁴⁵ Boland discussed the finding that $1,25(\text{OH})_2\text{D}_3$ rapidly induces reverse translocation of the VDR from the nucleus to the plasma membranes.⁴⁶ In accordance with this observation, a complex is formed between the VDR and TRCP3, an integral protein of capacitative Ca^{2+} entry, suggesting an association between both proteins and a functional role of the VDR in $1,25(\text{OH})_2\text{D}_3$ activation of CCE. Further supporting this interpretation, transfection with a VDR antisense oligonucleotide inhibited Ca^{2+} influx through SOC channels. In some vertebrate systems, as in invertebrate photoreceptor cells, TRP proteins have been shown to interact with the PDZ domain-containing INAD proteins, which serve as acceptors for a molecule with a regulatory function, which Boland hypothesized could be the VDR. Of additional mechanistic importance, when muscle cells are treated with $1,25(\text{OH})_2\text{D}_3$ in the presence of a specific anti-INAD antibody or transfected with an anti-INAD

antisense oligonucleotide, the hormone-dependent CCE was almost totally suppressed.^{43,45}

These data enabled Boland and colleagues to propose a non-genomic mechanism by which $1,25(\text{OH})_2\text{D}_3$ regulates intracellular Ca^{2+} levels in skeletal muscle cells: by acting on voltage-dependent channels via protein kinase-mediated phosphorylation and on SOC/TRCP3 channels through VDR-joining membrane supramolecular complexes. Alterations in these mechanisms during vitamin D deficiency states may account for skeletal muscle weakness, in view of the key role that Ca^{2+} plays in the regulation of contractility.

In addition, $1,25(\text{OH})_2\text{D}_3$ modulates growth-related signal transduction pathways in skeletal muscle cells. It has been established that the hormone activates MAPKs (ERK1/2) and Akt. Of note, the formation of complexes between the VDR and Src is involved in the activation of these pathways by the hormone.^{44,45} Experimental data demonstrates that $1,25(\text{OH})_2\text{D}_3$ induces the translocation of activated ERK to the nucleus, where it phosphorylates the transcriptional factors Elk and CREB, which coregulate the expression of genes that mediate mitogenic effects of the hormone.⁴³ Akt is activated by $1,25(\text{OH})_2\text{D}_3$ through PI3K, as the inhibitors Ly294002 and wortmannin block the increase in its phosphorylation in response to the hormone. Moreover, Akt mediates $1,25(\text{OH})_2\text{D}_3$ -induced differentiation of skeletal muscle myoblasts to myotubes. With regard to the mechanism by which the hormone initiates its action at the muscle cell plasma membrane, preliminary work using M β CD and siRNA technology has shown that intact caveolae and caveolin-1 expression are involved in ERK1/2 and Akt activation by $1,25(\text{OH})_2\text{D}_3$ (C. Buitrago & R. Boland, submitted). Boland concluded that the information described contributes to the elucidation of the mechanisms involved in the regulation of contractility and myogenesis by $1,25(\text{OH})_2\text{D}_3$.

Vitamin D and physical and cognitive function in older persons

Luigi Ferrucci (National Institute on Aging) discussed recent evidence linking vitamin D deficiency to the pathogenesis and clinical progression of chronic diseases highly prevalent in older persons. In large population-based studies, low levels of vitamin D predict loss of mobility and disability in basic

activities of daily living, independent of age and other potential confounders.⁴⁷ Based on these findings, some gerontologists claim that vitamin D has antiaging effects. Indeed, beyond its roles in bone health and calcium homeostasis, vitamin D affects many other physiological domains that are critically modified by aging. Unfortunately, the question of whether vitamin D prevents or slows aging cannot be easily tested. In spite of the intuitive nature of aging as a phenomenon, scientists have been unable to agree on good measures of biological aging because of great interindividual variability. Specific phenotypic changes occur in almost all aging individuals: changes in body composition, imbalance between production and utilization of energy, loss of redundancy of the homeostatic network, and loss of neurons and neuronal plasticity. Shifting attention from aging to the aging phenotype, the question of whether vitamin D affects major aging phenotypes becomes more approachable.

Data from the literature can generate some answers. Major changes in body composition that occur with aging include a decline in lean body mass (mostly muscle) and increase in fat mass. There is strong evidence that obesity is associated with poor vitamin D status due to fat sequestration.⁴⁸ In fact, compared with the non-obese, higher doses of vitamin D are required in obese individuals. Also, regardless of dietary intake, weight loss results in increased serum 25(OH)D₃ levels in overweight or obese women. However, some recent data suggest that low vitamin D predicts an accelerated increase in fat mass and incident obesity.

The importance of vitamin D for muscle has been discussed at length. Interestingly, VDR expression in human muscle tissue decreases with aging, and vitamin D level is strongly correlated with muscle quality and lower muscle fat infiltration. Consistently, in the Longitudinal Aging Study Amsterdam, low vitamin D and high parathyroid hormone were strong, independent risk factors for loss of muscle strength and muscle mass (sarcopenia).⁴⁹ Low vitamin D levels also appear to be correlated with collagen deposition and are related to arterial stiffness.⁵⁰ The role of vitamin D in energetics has been less thoroughly investigated. A recent study demonstrated that vitamin D level correlates with maximal oxygen consumption, and that supplementation of vitamin D increases fitness in athletes. Interestingly, a recent study found that VDRs are highly con-

centrated in the cristae of the mitochondria inner membrane.⁵¹ The role of vitamin D in homeostatic signaling is complex and pervasive. Vitamin D affects several pathways related to inflammation and cell proliferation, including inhibition of aromatase (conversion from testosterone to estrogens), COX2, prostaglandin receptor, NF- κ B, HIF-1 and VEGF, and stimulation of IGFBP-3 and E-cadherin.⁵² In addition, poor vitamin D status is a risk factor for several autoimmune diseases, confirming the role of this vitamin in immune function.

The role of vitamin D in neurologic function and pathology has been studied from multiple perspectives. Vitamin D receptors are found in high concentration in various areas of the brain. Low vitamin D levels have been associated with accelerated cognitive decline in epidemiological studies, and low vitamin D dietary intake is associated with increased risk of Alzheimer's disease and depression.^{53,54} Mechanisms suggested for this association include β -amyloid formation/aggregation, dysregulation of GABAergic activity and NMDA receptor, and increased calcium entry into neurons. Overall, vitamin D appears to be implicated in many of the physiological and pathological changes that occur with aging. Whether vitamin D supplementation may positively impact the aging process remains unknown and will require further long-term intervention studies.

Vitamin D dietary reference intakes

Daniel Bikle (University of California and VA Medical Center, San Francisco) addressed the recent Institute of Medicine (IOM) recommendations for dietary intake of vitamin D and evaluated whether they were sufficient in the context of emerging data about the roles of vitamin D and the prevalence of deficiency. Recommendations for the appropriate amount of vitamin D to be taken for optimal health—be it for musculoskeletal strength or for the numerous nonskeletal presumptive benefits of vitamin D—remain contentious. There is general consensus that the earlier recommendation of 400 IU of vitamin D/day was insufficient. Recently an IOM expert panel established to formulate new guidelines recommended that 600 IU/day would suffice for the general public aged 1–70 years, with the dose increased to 800 IU/day for those older than 70 years. However, doses up to 4000 IU/day were considered safe.⁵⁵ The task force compiling the Endocrine

Society Clinical Practice Guidelines (ESCPG) reached similar conclusions.⁵⁶ Vitamin D status is best assessed by the circulating levels of 25(OH)D₃. There is considerable variation among individuals with regard to the serum level of 25(OH)D₃ achieved with a given dose of oral vitamin D, and it is the serum level that counts with respect to biologic effect. The IOM expert panel concluded that most Americans (97.5%) had adequate levels of vitamin D, based on their conclusion that a 25(OH)D₃ of 20 ng/mL or greater was sufficient, raising the question of whether supplementation was even needed.

This conclusion was based on data from the 2001–2004 NHANES survey⁵⁷ that primarily sampled Caucasians, with limited sampling in Northern latitudes during the winter, the time and region that people would most likely have lower vitamin D levels. In that survey the average 25(OH)D₃ level was over 20 ng/mL. People of color, especially African-Americans, have substantially lower levels of 25(OH)D₃. Moreover, a number of studies have shown that prevailing levels of 25(OH)D₃, especially in older subjects, the institutionalized, and in most other countries, are well below those documented in the NHANES survey.⁵⁸ As parts of the aging process, the skin becomes less able to make vitamin D for a given amount of UVB exposure, the kidneys become less able to produce the active metabolite 1,25(OH)₂D₃, and the intestine becomes less able to respond to the 1,25(OH)₂D₃ with respect to calcium absorption. Furthermore, the conclusion by the IOM expert panel, that 20 ng/mL is sufficient, differs from that reached by the task force compiling the ESCPG. This task force recommended a higher target level, namely 30 ng/mL (75 mM), and pointed out that many individuals with a variety of conditions, including obesity, dark skin, malabsorption, renal failure, lactose intolerance, ingestion of certain drugs, and the institutionalized, are likely not to achieve adequate 25(OH)D levels by ingesting only 600 IU/day. If 30 ng/mL is deemed the optimal level (and neither set of recommendations considers this level unsafe), then the percent of Americans that maintain the optimal level falls well below 50%.

A further complication in trying to achieve a given level of 25(OH)D₃ is that 25(OH)D₃ assays have not proven consistent, making decisions regarding supplementation all the more difficult. This situation is improving with the availability of standards from the National Institute of Standards and Technol-

ogy. However, those assays relying on binding proteins or antibodies tend to report lower 25(OH)D levels than assays employing chromatography/mass spectroscopy.⁵⁹ Moreover, the latter can separately determine 25(OH)D₂ and 25(OH)D₃ levels, as well as the level of 3-epimer of 25(OH)D₃, capabilities that are lacking in the binding protein/antibody assays. While the clinical significance of these differences in assays is not clear, they do complicate decision making when a fixed target level of 25(OH)D₃ is being sought. In practical terms, young, healthy, light-skinned individuals with good diets that include dairy products and who have adequate exposure to sunlight are not likely to be vitamin D deficient or need to be tested. Individuals of color, the aged, those with gastrointestinal disorders, the institutionalized, and those with poor nutrition should be tested and treated appropriately to achieve a 25(OH)D₃ level of at least 20 ng/mL, or perhaps closer to 30 ng/mL. Daily doses appear to be safer and more efficacious than large doses administered once or twice a year.

Conclusions

The conference “Vitamin D: Beyond Bone” explored a considerable range of functions for vitamin D and potential therapeutic implications outside of calcium homeostasis and bone health, highlighting the increased understanding of vitamin D, as well as how much is yet to be learned. The scope of vitamin D research and its therapeutic possibilities, as presented by researchers across many different disciplines, generated enthusiasm tempered with some skepticism. Continuing research is needed to better understand the relationships between vitamin D and extraskelatal health and to determine the optimal dose of vitamin D for individuals based on age, health conditions, and other factors.

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Conflicts of interest

The authors declare no conflicts of interest.

References

- Christakos, S. & H.F. DeLuca. 2011. Minireview: Vitamin D: is there a role in extraskelatal health? *Endocrinology* **152**: 2930–2936.
- Dhawan, P., R. Wieder & S. Christakos. 2009. CCATT enhancer binding protein alpha is a molecular target of 1,25-dihydroxyvitamin D3 in MCF-7 breast cancer cells. *J. Biol. Chem.* **284**: 3086–3095.
- Joshi, S., L.-C. Pantalena, X. K. Liu, *et al.* 2011. 1,25-Dihydroxyvitamin D3 ameliorates Th17 autoimmunity via transcriptional modulation of interleukin-17A. *Mol. Cell. Biol.* **31**: 3653–3669.
- Liu, P.T. *et al.* 2006. Toll-like receptor triggering of a vitamin D-mediated human antimicrobial response. *Science* **311**: 1770–1773.
- Hewison, M. 2011. Antibacterial effects of vitamin D. *Nat. Rev. Endocrinol.* **7**: 337–345.
- Hewison, M. *et al.* 2003. Differential regulation of vitamin D receptor and its ligand in human monocyte-derived dendritic cells. *J. Immunol.* **170**: 5382–5390.
- Jeffery, L.E. *et al.* 2012. Availability of 25-Hydroxyvitamin D3 to APCs Controls the Balance between Regulatory and Inflammatory T Cell Responses. *J. Immunol.* **189**: 5155–5164.
- Gardner, D.G., S. Chen, D.J. Glenn & W. Ni. 2011. Vitamin D and the Cardiovascular System. In: *Vitamin D*, 3rd Edition. D.J.W. Feldman, Pike and J.S. Adams, eds: 541–563. Elsevier: New York.
- Li, Y.C., J. Kong, M. Wei, *et al.* 2002. 1,25-dihydroxyvitamin D(3) is a Negative Endocrine Regulator of the Renin-Angiotensin System. *J. Clin. Invest.* **110**: 229–238.
- Chen, S. & D.G. Gardner. Liganded Vitamin D Receptor Displays Anti-Hypertrophic Activity in the Murine Heart. *J. Steroid. Biochem. Mol. Biol.*, in press.
- Chen, S., C.S. Law, C.L. Grigsby, *et al.* 2011. Cardiomyocyte-specific deletion of the vitamin D receptor gene results in cardiac hypertrophy. *Circulation* **124**: 1838–1847.
- Hollis, B.W., D. Johnson, T.C. Hulsey, *et al.* 2011. Vitamin D supplementation during pregnancy: double-blind, randomized clinical trial of safety and effectiveness. *J. Bone. Miner.* **26**(10): 2341–2357. PMID: 3183324.
- Mahomed, K. & A.M. Gulmezoglu. 1999. *Vitamin D supplementation in pregnancy (Cochrane Review)*. *Cochrane Database of Systematic Reviews*. John Wiley & Sons, Ltd. Chicester, UK.
- De-Regil, L.M., C. Palacios, A. Ansary, *et al.* 2012. Vitamin D supplementation for women during pregnancy. *Cochrane Database of Systematic Reviews* **2**: CD008873.
- Hollis, B.W. & C.L. Wagner. 2012. Vitamin D and pregnancy: skeletal effects, nonskeletal effects, and birth outcomes. *Calcif. Tissue. Int.*
- Wagner, C.L., R. McNeil, S.A. Hamilton, *et al.* 2012. A randomized trial of vitamin D supplementation in 2 community health center networks in South Carolina. *Am. J. Obstet. Gynecol.*
- Wagner, C.L., T.C. Hulsey, D. Fanning, *et al.* 2006. High-dose vitamin D3 supplementation in a cohort of breastfeeding mothers and their infants: a 6-month follow-up pilot study. *Breastfeed Med.* **1**: 59–70.
- Wagner, C.L., C. Howard, T.C. Hulsey, *et al.* 2012. Preliminary Results of a Randomized Controlled Trial of Maternal Supplementation with 6400 IU Vitamin D/day Compared with Maternal & Infant Supplementation of 400 IU/day in Achieving Sufficiency in the Breastfeeding Mother-Infant Dyad. *Intl. Society Res. Human Milk Lactation*. in press (abstract).
- Sergeev, I.N. 2009. Novel mediators of vitamin D signaling in cancer and obesity. *Immun. Endoc. Metab. Agents Med. Chem.* **9**: 153–158.
- Song, Q. & I.N. Sergeev. 2012. Calcium and vitamin D in obesity. *Nutr. Res. Rev.* **25**: 130–141.
- Sergeev, I.N. 2012. Vitamin D and cellular Ca²⁺ signaling in breast cancer. *Anticancer Res.* **32**: 299–302.
- Sergeev, I.N. 2005. Calcium signaling in cancer and vitamin D. *J. Steroid Biochem. Mol. Biol.* **97**: 145–151.
- Sergeev, I.N. 2009. 1,25-Dihydroxyvitamin D3 induces Ca²⁺-mediated apoptosis in adipocytes via activation of calpain and caspase-12. *Biochem. Biophys. Res. Commun.* **384**: 18–24.
- Janssens, W. *et al.* 2009. Vitamin D Beyond Bones in COPD: Time to Act. *Am. J. Respir. Crit. Care Med.*
- Janssens, W. *et al.* Vitamin D deficiency is highly prevalent in COPD and correlates with variants in the vitamin D-binding gene. *Thorax.* **65**(3): 215–220.
- Romme, E.A. *et al.* 2012. Vitamin D status is associated with bone mineral density and functional exercise capacity in patients with chronic obstructive pulmonary disease. *Ann. Med.*
- Persson, L.J. *et al.* 2012. Chronic obstructive pulmonary disease is associated with low levels of vitamin d. *PLoS One* **7**(6): e38934.
- Forli, L. *et al.* 2004. Vitamin D deficiency, bone mineral density and weight in patients with advanced pulmonary disease. *J. Intern. Med.* **256**(1): 56–62.
- Hopkinson, N.S. *et al.* 2008. Vitamin D receptor genotypes influence quadriceps strength in chronic obstructive pulmonary disease. *Am. J. Clin. Nutr.* **87**(2): 385–390.
- Lehouck, A. *et al.* 2012. High doses of vitamin D to reduce exacerbations in chronic obstructive pulmonary disease: a randomized trial. *Ann. Intern. Med.* **156**(2): 105–114.
- Black, P.N. & R. Scragg. 2005. Relationship between serum 25-hydroxyvitamin d and pulmonary function in the third national health and nutrition examination survey. *Chest* **128**(6): 3792–3798.
- Kumar, V.A., D.A. Kujubu, J.J. Sim, *et al.* 2011. Vitamin D supplementation and recombinant human erythropoietin utilization in vitamin D-deficient hemodialysis patients. *J. Nephrol.* **24**(1): 98–105.
- Andrews, N.C. 2004. Anemia of inflammation: the cytokine-hepcidin link. *J. Clin. Invest.* **113**(9): 1251–1253.
- Howick, J., P. Glasziou & J.K. Aronson. 2009. The evolution of evidence hierarchies: what can Bradford Hill's 'guidelines for causation' contribute? *J. Royal Soc. Med.* **102**: 186–194.
- Mitri, J., M.D. Muraru & A.G. Pittas. 2011. Vitamin D and type 2 diabetes: a systematic review. *Eur. J. Clin. Nutr.* **65**: 1005–1015.
- Song, Y., L. Wang, A.G. Pittas, *et al.* Blood 25-hydroxyvitamin D concentration and incident Type 2

- Diabetes: A meta-analysis of prospective studies diabetes care (in press).
37. Pittas, A.G. *et al.* 2007. The effects of calcium and vitamin D supplementation on blood glucose and markers of inflammation in nondiabetic adults. *Diabetes Care* **30**: 980–986.
 38. von Hurst, P.R. *et al.* 2008. Study protocol—metabolic syndrome, vitamin D and bone status in South Asian women living in Auckland, New Zealand: a randomised, placebo-controlled, double-blind vitamin D intervention. *BMC Public Health* **8**: 267.
 39. Mitri, J. *et al.* 2011. Effects of vitamin D and calcium supplementation on pancreatic beta cell function, insulin sensitivity, and glycemia in adults at high risk of diabetes: the calcium and vitamin D for diabetes mellitus (CaDDM) randomized controlled trial. *Am. J. Clin. Nutr.* **94**: 486–494.
 40. Boland, R. 1986. Role of vitamin D in skeletal muscle function. *Endocr. Rev.* **7**: 434–448.
 41. Gallagher, J.C. 2004. The effects of calcitriol on falls and fractures and physical performance tests. *J. Steroid Biochem. Mol. Biol.* **89–90**: 497–501.
 42. Bischoff, H.A., M. Borchers, F. Gudat, *et al.* 2001. In situ detection of 1,25-dihydroxyvitamin D₃ receptor in human skeletal muscle tissue. *Histochem. J.* **33**: 19–24.
 43. Boland, R. 2005. Vitamin D and Muscle. In: *Vitamin D*. 2nd Edition; D. Feldman, G. Glorieux & W. Pike, eds: 885–898. Academic Press. CA, USA. Chapter 55.
 44. Boland, R. 2011. VDR activation of intracellular signaling pathways in skeletal muscle. *Mol. Cell. Endocrinol.* **34**: 11–16.
 45. Boland, R., A.R. de Boland, C. Buitrago, *et al.* 2002. Non-genomic stimulation of tyrosine phosphorylation cascades by 1,25(OH)₂D₃ by VDR-dependent and -independent mechanisms in muscle cells. *Steroids* **67**: 477–482.
 46. Capiati, D., S. Benassati & R. Boland. 2002. 1,25(OH)₂-vitamin D₃ induces translocation of the vitamin D receptor (VDR) to the plasma membrane in skeletal muscle cells. *J. Cell. Biochem.* **86**: 128–135.
 47. Houston, D.K., R.H. Neiberg, J.A. Tooze, *et al.* 2013. Low 25-hydroxyvitamin D predicts the onset of mobility limitation and disability in community-dwelling older adults: The health ABC study. *J. Gerontol. A Biol. Sci. Med. Sci.* **68**(2): 181–187.
 48. Rock, C.L., J.A. Emond, S.W. Flatt, *et al.* 2012. Weight loss is associated with increased serum 25-hydroxyvitamin D in overweight or obese women. *Obesity*. **20**(11): 2296–2301.
 49. Visser, M., D.J. Deeg, P. Lips, Longitudinal Aging Study Amsterdam. 2003. Low vitamin D and high parathyroid hormone levels as determinants of loss of muscle strength and muscle mass (sarcopenia): the longitudinal aging study amsterdam. *J. Clin. Endocrinol. Metab.* **88**(12): 5766–5772.
 50. Giallauria, F., Y. Milaneschi, T. Tanaka, *et al.* 2012. Arterial stiffness and vitamin D levels: the Baltimore longitudinal study of aging. *J. Clin. Endocrinol. Metab.* **97**(10): 3717–3723.
 51. Silvagno, F., E. De Vivo, A. Attanasio, *et al.* 2010. Mitochondrial localization of vitamin D receptor in human platelets and differentiated megakaryocytes. *PLoS One.* **5**(1): e8670.
 52. Krishnan, A.V. & D. Feldman. 2011. Mechanisms of the anti-cancer and anti-inflammatory actions of vitamin D. *Annu. Rev. Pharmacol. Toxicol.* **51**: 311–336.
 53. Llewellyn, D.J., I.A. Lang, K.M. Langa, *et al.* 2010. Vitamin D and risk of cognitive decline in elderly persons. *Arch. Intern. Med.* **170**(13): 1135–1141.
 54. Milaneschi, Y., M. Shardell, A.M. Corsi, *et al.* 2010. Serum 25-hydroxyvitamin D and depressive symptoms in older women and men. *J. Clin. Endocrinol. Metab.* **95**(7): 3225–3233.
 55. Institute of Medicine. 2011. *Dietary reference intakes for calcium and vitamin D*. Washington DC: The National Academies Press.
 56. Holick, M.F., N.C. Binkley, H.A. Bischoff-Ferrari, *et al.* 2011. Evaluation, treatment and prevention of vitamin D deficiency: an endocrine society clinical practice guideline. *J. Clin. Endocrinol. Metab.* **96**: 1911–1930
 57. Looker, A.C., C.L. Johnson, D.A. Lacher, *et al.* 2011. Vitamin D status: United States, 2001–2006. National Center for Health Statistics Data Brief no 59.
 58. Kuchuk, N.O., N.M. van Schoor, S.M. Pluijm, *et al.* 2009. Vitamin D status, parathyroid function, bone turnover, and BMD in postmenopausal women with osteoporosis: global perspective. *J. Bone. Min. Res.* **24**: 693–701.
 59. Binkley, N., D.C. Krueger, S. Morgan & D. Wiebe. 2010. Current status of clinical 25-hydroxyvitamin D measurement: an assessment of between-laboratory agreement. *Clin. Chim. Acta.* **411**: 1976–1982
 60. Christakos, S., P. Dhawan, Y. Liu, *et al.* 2003. New Insights into the mechanisms of vitamin D action. *J. Cell. Biochem.* **88**: 695–705.