Vitamin D in Aging and Chronic Illness

Veronica A. Ralls¹, Alaina P. Boyer² and Consuelo H. Wilkins¹,²*¹
¹Meharry Medical College, Vanderbilt University Medical Center, Nashville, TN, USA
²Meharry-Vanderbilt Alliance, Vanderbilt University Medical Center, Nashville, TN, USA

Abstract

The role of vitamin D in calcium homeostasis and its impact on bone is well characterized and validated; however, the role vitamin D plays in non-bone physiology is less concrete. The Vitamin D Receptor (VDR) is expressed throughout the body and regulates many cellular processes. This discovery has led to vigorous research of vitamin D and the role it plays in many illnesses, especially those of high prevalence in the United States. Vitamin D deficiency has been implicated in numerous health conditions and thus is an intriguing target for therapeutic intervention. Studies examining the therapeutic effects of vitamin D in chronic disease and disease prevention have yielded conflicting results. Additionally, many publications on vitamin D result from studies in which vitamin D was not the primary focus. Given the increasing interest in the non-bone effects of vitamin D, we will review and summarize the recent literature related to older adults, a group with significantly increased risk of vitamin D deficiency. Older adults have substantial morbidity and mortality due to dementia, cancer and heart disease, all of which have been linked to vitamin D deficiency. We will explore current evidence of the expression of VDR and the effects of exposure to vitamin D that might impact these illnesses among older adults. We will review the most recent research on cognitive function and depression as a result of vitamin D deficiency. Through this work we aim to summarize the current data that sheds light on the possibility of clinical application of vitamin D therapy.

Keywords: Vitamin D; Heart disease; Cancer

Introduction

The purpose of this review it to summarize the evidence supporting the role of vitamin D in cognitive and physical health. We will summarize hormonal characteristics and describe the biological and physiological actions of vitamin D, the effects of vitamin D deficiency on cognitive and physical functions as well as depressive symptomatology. Finally, we will discuss the current state of evidence of the role of vitamin D deficiency in cardiovascular disease, cancer, and diabetes. Table 1 summarizes important information from each of the clinical studies reviewed, including a list of primary outcomes of interest.

Vitamin D

Hormone

Vitamin D is an essential sterol hormone which undergoes bioactivation in a tightly regulated process. In response to sunlight exposure, 7-dehydrocholesterol is converted to previtamin D3 in the skin by photolytic conversion which is then followed by thermal isomerization to form vitamin D3 [1]. Vitamin D3 is released into the blood where it is hydroxylated at carbon 25 primarily in the liver, although skin, intestine and kidney have been reported to catalyze this hydroxylation [2]. Of note, 25-hydroxyvitamin D (calcidiol) levels increase in proportion to oral vitamin D intake so plasma 25-hydroxyvitamin D levels are commonly used as an indicator of vitamin D status [1]. A second hydroxylation occurs by 1α-hydroxylase, primarily in the kidney, to form 1,25-dihydroxyvitamin D (calcitriol), the biologically active and hormone form of vitamin D (Figure 1). The presence of 1α-hydroxylase in the human brain has been confirmed suggesting a role of vitamin D in the brain [3]. Both Parathyroid Hormone (PTH) and serum calcium regulate renal synthesis of calcitriol in a feed-back loop to control bone mineral homeostasis.

Because dietary sources of vitamin D are limited, well-nourished people are still at risk for vitamin D deficiency. Foods with vitamin D include fatty fish such as sardines and salmon, liver, and egg yolks. Some foods, primarily milk and dry cereal, are fortified with vitamin D. Interestingly, the amount of vitamin D in fortified milk was determined based on the needs of preschool-aged children, not adults, and the amount is targeted to prevent rickets. Foods fortified with vitamin D often contain between 100 and 200 IU per serving while vitamin D that is generated from sunlight is estimated to be 3,000 to 5,000 units/day [4]. Vitamin D3 (cholecalciferol), from an animal source and vitamin D2 (ergocalciferol) from yeast and plants, can be administered orally and are subsequently converted to active vitamin D metabolites. Older adults, individuals with darker skin and those with extreme obesity (BMI<40 kg/m²) are at increased risk for vitamin D deficiency.

Biological and physiological action

The biologic actions of vitamin D may be considered as the classical and non-classical vitamin D responsive tissues. The classical tissues include the kidney, bone, parathyroid gland and intestine, which interact to maintain extracellular calcium homeostasis [2]. The non-classical tissues include immune, adipose, reproductive, endocrine, brain, liver, muscle, and skin [5-7]. The importance of vitamin D in bone mineralization and formation is well known and vitamin D deficiency alone can result in rickets, osteomalacia, and osteoporosis. In the small bowel, vitamin D improves absorption of calcium and phosphorus. Initially, calcium uptake is highly dependent on vitamin D [4]. Additionally, vitamin D is a potent modulator of parathyroid function. Vitamin D deficiency results in hyperparathyroidism and conversely, vitamin D administration inhibits PTH synthesis [8]. The expression of 1α-hydroxylase in numerous cell types unrelated to calcium homeostasis including cells of the central nervous system [9], suggests an autocrine or paracrine function of vitamin D in these tissues [3]. The non-classical vitamin D responsive tissues are unrelated to its effect on calcium homeostasis. These tissues include neurons, muscle, activated T cells, islet cells and aortic endothelial cells (Figure 1).

*Corresponding author: Consuelo H. Wilkins, Vanderbilt University School of Medicine, Nashville, TN, USA, Tel: 615-963-2820; Fax: 615-320-9457; E-mail: consuelo.h.wilkins@meharry-vanderbilt.org

Received July 02, 2013; Accepted September 09, 2014; Published September 16, 2014

Citation: Ralls VA, Boyer AP, Wilkins CH (2014) Vitamin D in Aging and Chronic Illness. Vitam Miner 3: 125.

Copyright: © 2014 Ralls VA, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.
<table>
<thead>
<tr>
<th>Reference</th>
<th>Sample</th>
<th>N</th>
<th>Measurements</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cognitive Function</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kipen [25]</td>
<td>Community-dwelling older women with mild dementia and NC</td>
<td>20</td>
<td>Bone density, PTH, Vitamin D₃</td>
<td>Bone density: NSD between groups PTH: levels were higher in dementia group (P&lt;0.01) Vitamin D₃ levels were lower in the dementia group (P=0.01)</td>
</tr>
<tr>
<td>Ogihara [26]</td>
<td>3 groups of hospitalized women; NC, DAT, VaD</td>
<td>60</td>
<td>Ca levels, PTH, Vitamin D₃</td>
<td>DAT group showed a significant decrease in Ca (P&lt;0.01) and increase in PTH (P&lt;0.05) compared to NC</td>
</tr>
<tr>
<td>Sato [27]</td>
<td>Nursing home dwelling women with DAT and NC controls from the community</td>
<td>46</td>
<td>BMD, Vitamin D₃, PTH</td>
<td>BMD, vitamin D₃ levels were lower in DAT (P&lt;0.0001); PTH was lower in DAT (P=0.001)</td>
</tr>
<tr>
<td>Jorde [28]</td>
<td>Adults, with secondary hyperparathyroidism (SHPT) and healthy controls</td>
<td>84</td>
<td>Cognitive test of working memory, information processing, memory, language, executive function</td>
<td>SHPT group had poorer performance on Digit Span Forward (P&lt;0.01); Stroop Test (P&lt;0.05), Word Association Test (P&lt;0.01) relative to controls</td>
</tr>
<tr>
<td>Wilkins [15]</td>
<td>African American and European American older adults with NC or mild cognitive impairment</td>
<td>60</td>
<td>Vitamin D₃, General cognitive function</td>
<td>Lower vitamin D₃ levels were associated with poorer cognitive performance (P&lt;0.01); African Americans had lower vitamin D₃ levels (P&lt;0.0001)</td>
</tr>
<tr>
<td><strong>Depression</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Przybelski [31]</td>
<td>Chart review of clinic records of older adult patients referred for memory loss</td>
<td>80</td>
<td>Vitamin D₃, General cognitive function</td>
<td>Vitamin D₃ levels were positively correlated with MMSE performance (P=0.006)</td>
</tr>
<tr>
<td>Wilkins [16]</td>
<td>Participants with NC or mild AD</td>
<td>80</td>
<td>Vitamin D₃, General cognitive function</td>
<td>Vitamin D₃ deficiency was not predictive of AD (P=0.33); was associated with poorer SBT performance (P&lt;0.04); higher CDR Sum of Boxes (P=0.02)</td>
</tr>
<tr>
<td>Oren [34]</td>
<td>Adults with SAD and age-and-sex matched controls</td>
<td>30</td>
<td>Vitamin D₃ levels with and without light therapy</td>
<td>NSD in vitamin D₃ between groups or between light conditions (P=0.05)</td>
</tr>
<tr>
<td>Lapid [36]</td>
<td>Primary care patients, older adults</td>
<td>1,618</td>
<td>Vitamin D₃, depression diagnosis</td>
<td>Lower vitamin D₃ levels were associated with increased risk for depression (P=0.01)</td>
</tr>
<tr>
<td>Schneider [37]</td>
<td>Adult schizophrenia patients, alcohol addiction patients, patients with major depression, and NC</td>
<td>34</td>
<td>Vitamin D₃, PTH, Calcium, Phosphate</td>
<td>Vitamin D₃ levels were lower in psychiatric patients than NC (P&lt;0.01)</td>
</tr>
<tr>
<td>Hoogendijk [39]</td>
<td>Community-dwelling older adults</td>
<td>1,282</td>
<td>Vitamin D₃, PTH, CES-D</td>
<td>Lower vitamin D₃ levels were associated with depression severity (P&lt;0.001), PTH levels were positively associated with depression (P=0.003)</td>
</tr>
<tr>
<td>Gloth [42]</td>
<td>Adults with SAD who received Vitamin D₃ (1000 IU) supplementation or phototherapy</td>
<td>15</td>
<td>Depression, Vitamin D₃ levels</td>
<td>Vitamin D₃ group improved on all depression outcomes (P&lt;0.05), vitamin D₃ levels improved in both groups 74% vitamin D₃ group, p&lt;0.005 and 36% phototherapy group, P&lt;0.01</td>
</tr>
<tr>
<td>Lansdowne [43]</td>
<td>Healthy adults received Vitamin D3 supplementation (400 IU/d, 800 IU/d, placebo)</td>
<td>44</td>
<td>Vitamin D₃ levels, Positive and Negative Affect</td>
<td>Vitamin D₃ groups expressed enhanced positive affect relative to controls (P&lt;0.001)</td>
</tr>
<tr>
<td><strong>Physical Function</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peterson [44]</td>
<td>Community-dwelling older adults</td>
<td>159</td>
<td>Falls, Motor function</td>
<td>Fallsers had lower vitamin D₃ levels (P&lt;0.01), NSD in motor function (P=0.05)</td>
</tr>
<tr>
<td>Sato [48]</td>
<td>Parkinson's disease (PD) patients and patients with vascular Parkinsonism (VP) who received ergocalciferol (1200 IU)</td>
<td>178</td>
<td>Fall, muscle strength, and hip fracture incidence</td>
<td>The number of falls decreased in the VP patients (P&lt;0.0001), hip fractures caused by falls occurred in seven cases in the PD group, and in one case in the VP group, increase in muscle strength occurred in both groups (P=0.001)</td>
</tr>
<tr>
<td>Dukas [47]</td>
<td>Older adults at increased risk for falls who received aflacidol (1µg)</td>
<td>237</td>
<td>Timed up and Go Test (TUG), Tandem Stand Test,(TST) Chair Rising Test (CRT), Falls</td>
<td>Increased muscle power (TUG, CRT)and balance (TST), and a decrease in falls (all P-values&lt;0.0001) was observed after supplementation</td>
</tr>
<tr>
<td>Schacht [48]</td>
<td>Older adults with reduced bone mass who received alfacalidol (1µg)</td>
<td>2,097</td>
<td>TUG, TST, CRT, fear of falls</td>
<td>Improvement in all muscle tests was observed (P&lt;0.0001) after supplementation</td>
</tr>
<tr>
<td>Sanders [49]</td>
<td>Community-dwelling older women who received cholecalciferol (500,000 IU/ yearly) or placebo</td>
<td>2,256</td>
<td>Falls and fractures (radiologically confirmed)</td>
<td>Treatment group experienced 15% more falls (P=0.03) and 26% more fractures (P=0.04) than the placebo group</td>
</tr>
<tr>
<td>Peterson [51]</td>
<td>Parkinson's disease patients</td>
<td>40</td>
<td>Balance function, Unified Parkinson's Disease Rating Scale (UPDRS)</td>
<td>Vitamin D₃ levels were inversely correlated with PD severity (P&lt;0.04), positively correlated with automatic posture responses to backwards translation (P&lt;0.05), specifically with response strength and stance weight (P&lt;0.05)</td>
</tr>
<tr>
<td>Stein [52]</td>
<td>Community-dwelling older adults and participants with mild-to-moderate cognitive impairment (MMSE score 12-24), all received vitamin D₃ supplementation (1000 IU/8 wk), then randomized to 6000 IU/8 wk (high dose group) or placebo</td>
<td>95</td>
<td>Disability assessment in dementia (DAD) questionnaire</td>
<td>NSD in disability after high dose (6000 IU) (P&gt;0.05)</td>
</tr>
</tbody>
</table>
Cardiovascular Disease

- Melamed [56]
  Adults ≥ 20 years old
  13,331 Vitamin D, incidence of cancer, cardiovascular disease, all-cause mortality
  Lowest quartile of vitamin D was associated with a 26% increased rate of all-cause mortality (1.26, 95% CI 1.08-1.46).

- Forman [57]
  African American adults aged 30-80 years who were randomized in to a 4-arm trial (placebo, 1000 IU, 2000 IU, 4000 IU/d) of cholecalciferol
  283 Blood pressure
  1-ng/mL increase in plasma vitamin D levels were associated 0.2 mmHg reduction in systolic blood pressure (P=0.02).

- Larsen [58]
  Adults with hypertension who were randomized to receive cholecalciferol (3000 IU/placebo)
  112 Blood pressure (24-hour, central), heart rate, pulse wave velocity, central augmentation,
  Cholecalciferol was associated with a decrease in central systolic blood pressure (P=.007) relative to controls.

- Judd [59]
  African Americans ≥ 30 years old with 25 ≤ vitamin D levels ≤ 75 were randomized to receive (placebo, calcitriol, or 200,000 IU cholecalciferol)
  9 Blood pressure
  Calcitriol group experienced a 9% decrease in systolic blood pressure compared to placebo (p<0.001).

- Snijder [60]
  Community-dwelling older adults
  1,205 Blood pressure, vitamin D, PTH
  Vitamin D levels were not associated with blood pressure (P>.05), PTH was positively associated with blood pressure (P<0.05).

- Jorde [61]
  Adults aged 25-69 years old
  27,159 Calcium intake, blood pressure
  A negative association between calcium intake and diastolic blood pressure (P<0.05).

- Jorde [62]
  Adults aged ≥ 25 years old
  4,125 Blood pressure, vitamin D levels
  Vitamin D levels were negatively associated with systolic blood pressure (P=.001), but did not predict future hypertension (P=.05).

- Jorde [63]
  Overweight and obese adults aged 21-70 years who were randomized for vitamin D supplementation (40000 IU/wk, 20000IU/wk, placebo)
  438 Cardiovascular risk factors (lipids, blood pressure, glucose metabolism)
  After 1 year, NSD were found between the 3 groups for measures of glucose metabolism or lipids (P>.05). In the 20000 IU group there was a slight increase in systolic blood pressure compared to placebo (P<.05).

- Wood [64]
  Postmenopausal women aged 60-70 years who were randomized for vitamin D supplementation (400 IU/d, 1000 IU/d, placebo)
  305 Cardiovascular disease risk (lipids, insulin resistance, markers of inflammation, blood pressure)
  After 1 year, daily supplementation with vitamin D had no effect on cardiovascular disease risk (P>.05).

- Correia [65]
  Patients with unstable angina in coronary care units
  206 Cardiovascular death during hospitalization
  Severe vitamin D deficiency was an independent predictor of in-hospital cardiovascular mortality (OR: 14, 95% CI: 1.2 to 158, P<.03).

Cancer Therapeutic

- Wagner [70]
  Patients with prostate cancer (Gleason score of 6 or 7) were randomized for vitamin D (400 IU/d, 10,000 IU, 40000 IU/d)
  66 Serum vitamin D, and prostate tissue levels of vitamin D metabolites, proliferation marker Ki67 (MB-1)
  Dose dependent increase in prostate tissue and serum levels of vitamin D metabolites (P<.03), NSD in Ki67 levels between groups (P>.05).

Diabetes Control

- Nikooyeh [73]
  Participants with T2D aged 30-60 years were randomized to consume plain yogurt, vitamin D-fortified yogurt (500 IU, 150 mg CA), or vitamin D (500 IU) + calcium-fortified yogurt (250 mg)/ 2xd/12 wks
  90 Glycemic status
  Daily intake of vitamin D-fortified yogurt with or without added calcium, improved glycemic status in T2D participants (P<.0001).

- von Hurst [75]
  South Asian women aged 23-68 years, with insulin resistance and vitamin D deficiency were randomized to receive 4000 IU or placebo for 6 months
  81 Insulin resistance (IR)
  Improvements were seen in insulin sensitivity (P=.003) and IR (P=.02) and fasting insulin decreased (P=.02) with supplementation compared with placebo.

- Aljabri [76]
  Participants ≥ 12 years old with T1D and vitamin D deficiency were randomized to receive 4000 IU/d or placebo for 12 weeks
  80 Glycosylated hemoglobin
  Vitamin D supplementation was associated with improved glycemic control (P=.04).

- Deleskog [77]
  Adults aged 33-56 years without T2D followed for 8-10years
  2,378 Incidence of prediabetes, T2D
  High vitamin D, levels predicted a reduced risk of T2D in individuals with prediabetes (OR 0.38, CI0.21, 0.71).

- Jorde [80]
  Adults aged 21-75 years, with T2D treated with insulin and metformin were randomized to receive supplementation (40000 IU/wk or placebo) for 6 months
  36 Glycemic control
  NSD from baseline values, NSD in change in glycemic control (P>.05).

Table 1: Study Design: A=randomized controlled trial (RCT); B=cross-sectional; C=longitudinal; D=Randomized trial; DAT=Dementia Alzheimer’s type; VaD=vascular dementia; NSD=no significant difference; NC=normal cognition; Ca=calcium; CA=calcium; CED=carotid intima-media thickness; CAC=crown arteries calciﬁcation; CH=coronary heart disease; PTH=parathyroid hormone; MMSE=Mini-mental State Examination; SBT=Short Blessed Test; CDR=Clinical Dementia Rating; SAD=Seasonal Affective Disorder; CES-D=Center for Epidemiologic Studies-Depression Scale; IU=International Units; T2D=type 2 diabetes; T1D= type 1 diabetes.

New mechanisms of vitamin D action have emerged and suggest effects on cell cycle and apoptotic processes which are critical events in the life cycle of a cell, but when deregulated can contribute to the pathology of disease development. Through a series of studies, VDR was found to suppress the oncogene c-myc while simultaneously inducing c-fos, two genes intricately involved in carcinogenesis [10]. Further validation is required to determine the exact mechanism of action in other model systems. Altered RNA expression of Bcl-2 genes has also been attributed to VDR activation by calcitriol in leukemia cells [11]. Vitamin D supplementation has been used to manage chronic...
conditions and diseases in older adults. Although dosing remains controversial, there are benefits to using vitamin D to manage diseases and conditions common in older adults. Vitamin D has been shown to regulate calcium-phosphate homeostasis and stimulate bone growth through binding to VDR, which is expressed on osteoblasts and osteoclasts. This regulation makes the hormone a valuable supplement in older adults, a group more at risk for falls and fractures. Vitamin D deficiency has also been implicated in chronic inflammation known to co-exist with the aging process. Vitamin D has demonstrated anti-inflammatory effects through inhibition of the activation of pro-inflammatory transcription factor NFκβ and interleukin-6 (IL-6) [12].

Effects of Vitamin D Deficiency

Cognitive function

Vitamin D plays an important role in brain function in vitro [13] and in vivo models [14]; recent evidence extends these findings to cognitive function in humans [15-17]. Vitamin D nuclear binding has been shown in the hippocampus [18], amygdala, and basal forebrain [19,20] which are important areas in memory processing. Additionally, vitamin D dependent proteins are reduced in the hippocampal cells of humans with Alzheimer’s disease (AD) [21] and transgenic mice with reduced vitamin D dependent proteins have impaired spatial learning [22]. Treatment with vitamin D stimulates choline acetyl-transferase in vitamin D-deficient rats [23] suggesting that vitamin D is important in the cholinergic systems associated with memory [24].

Observational studies report that levels of 25-hydroxyvitamin D are lower in participants with AD than non-demented controls [25-27]. The first paper to report an association between vitamin D deficiency and secondary hyperparathyroidism and impairment in neuropsychological function [28] in non-demented participants was published in 2006. This study found that participants with vitamin D deficiency and secondary hyperparathyroidism (without chronic renal disease) have significantly impaired performance in 3 of 14 cognitive tests including the Digit Span Forward, Stroop Test Part 1 and 2 and Word Association Test as well as significantly higher depression scores on the Beck Depression Inventory (BDI).

It is well established that persons with darker pigment, women and older adults are at greater risk of developing vitamin D deficiency. Further, African American women with vitamin D deficiency have worse outcomes than European women when performing cognitive assessments [15]. Secondary analysis of prospective research studies further confirms that poorer cognition is more strongly associated with vitamin D deficiency [29,30]. While these studies clearly support a relationship between cognition and vitamin D deficiency, the mechanism has not been fully elucidated.

Subsequently, other investigators have demonstrated that vitamin

Figure 1: Vitamin D Metabolism and Actions. Vitamin D is metabolized in multiple steps and is expressed in many tissues. The classical actions of vitamin D, bone health and thyroid regulation most often occur in specific tissues. Other non-classical functions of vitamin D include cell proliferation, effects on inflammatory proteins and effects on cognitive function.
D levels are associated with cognitive function in non-demented [16,31,32] and mildly demented [16] older adults. A recent finding in a cohort of community-dwelling older women further substantiates this relationship of vitamin D deficiency and cognition [33]. These findings suggest that a strong association exists between cognitive impairment and vitamin D status; therefore, supporting the hypothesis that vitamin D therapy may prevent or delay Age-Related Cognitive Decline (ARCD).

Depression

For years the association between vitamin D and mood disorders has been debated. Early studies found no association with depression and 1,25-dihydroxyvitamin D [34] but with the discovery of vitamin D receptors in the brain, researchers began to reconsider its role in mood disorders [35]. There have been several studies that support the notion that decreased levels of 25-hydroxyvitamin D levels are correlated with depression published in recent years. A recent study that enrolled a range of low to severely vitamin D deficient participants found the more severe the deficiency the greater the likelihood of depression (OR=2.45, 95% CI=1.24-4.64, p=0.006) [36]. One specific study found a correlation between 25-hydroxyvitamin D and Seasonal Affective Disorder (SAD) and depression [37,38]. Patients with a serum 25-hydroxyvitamin D concentration below 16 ng/ml at a reference point also had increased Beck Depression Inventory (BDI) scores compared to patients with higher serum levels [35]. Numerous populations have been assessed for their depressive symptoms associated with decreased 25-hydroxyvitamin D. Older adults with reasonable health showed an association between decreased 25-hydroxyvitamin D levels and depression according to the BDI [39]. Cardiovascular patients with no history of depression developed symptoms of depression in the presence of low 25-hydroxyvitamin D concentrations ranging from 15-30 ng/mL [35]. Depression status and severity was also associated with decreased vitamin D and increased PTH in a large population based study [40].

Since SAD has been associated with winter months and sunlight deprivation [41], vitamin D deficiency has been considered a possible contributor to SAD. Additionally, vitamin D was found to be superior to phototherapy in SAD [42] and in a placebo-controlled trial vitamin D enhanced positive affect [43].

Physical function

The risk of falling and injury in the aging population is a fear expressed in many older adults. Research focused on vitamin D deficiency and falls in the elderly have demonstrated significant (p<0.01) associations [44]. An improvement in fall risk was demonstrated in institutionalized women following a stroke with vitamin D supplementation who had low serum concentrations at baseline [45]. Another study showed significant reduction in fall risk in patients with vascular Parkinsonism [46]. It has further been demonstrated that treatment with alfalcacidol can increase muscle balance and power resulting in decreased risk of falls [47]. An additional study supported this claim [48]. On the contrary there have been studies that have shown no benefit in vitamin D supplementation and reduced fall risk [49].

Movement, stability, and falls go hand in hand. Osteoporotic fractures have a significant effect on movement, especially in the elderly population. Treatment of these fractures lies largely in prevention including screening and appropriate intake of vitamin D and calcium [50]. The use of vitamin D in movement disorders has also been implicated. Automatic postural control was shown to be correlated with vitamin D concentrations [51]. The association of Parkinson’s disease severity with vitamin D concentrations was demonstrated in the same study. Evidence is lacking to support the use of vitamin D in treatment of Alzheimer’s disease [52,53], but one recent study showed vitamin D intake to be associated with decreased risk of developing the disease in older women [54].

The overwhelming majority of recent data supports the use of vitamin D as treatment for certain movement disorders and to reduce fall risk. Vitamin D is already used in prevention of osteoporotic fractures and at least one phase III trial tested the use of vitamin D in prevention of falls [55]. The clinical application of vitamin D in conditions affecting movement is rapidly increasing and has the potential to have some major effects on the quality of life of patients living with these conditions.

Vitamin D and Disease

Cardiovascular illness

The role of vitamin D in Cardiovascular Disease (CVD) is debated among researchers. Vitamin D deficiency is associated with increased risk of mortality according to the National Health and Nutrition Examination Survey III (NHANES III) [35]. Vitamin D deficiency has also been linked to hypertension, diabetes, and insulin resistance, which are known risk factors for CVD. Groups at greater risk of becoming vitamin D deficient include older adults, women, African Americans, diabetics, current smokers, and high Body Mass Indices (BMI) [56]. One in three deaths were attributed to cardiovascular disease in 2009; however, the role that vitamin D deficiency plays in these deaths cannot be inferred. The association of cardiovascular mortality with 25-hydroxyvitamin D concentration may not be a causal relationship, but simply a reflection of the underlying health of the individuals of this group [35].

There is still more to understand about the relationship between 25-hydroxyvitamin D serum concentrations and blood pressure; although, it is known that vitamin D plays a role in the control of the Renin-Angiotensin-Aldosterone System (RAAS) and indirectly regulates intracellular calcium levels, which mutually work on smooth muscle vascular tone [35]. Two recent randomized, double-blind, placebo-controlled trials demonstrated that cholecalciferol administered during the winter months was able to significantly reduce systolic blood pressure [57,58]. No significant change was noted in diastolic blood pressure. Another recent trial demonstrated that vitamin D produced a short-term decrease in blood pressure [59]. Comparatively, a cross-sectional study and a longitudinal, population-based multipurpose study showed no correlation between vitamin D levels and blood pressure [60-62]. Another cross-sectional study did not support a correlation with vitamin D levels and blood pressure [63].

Although vitamin D therapy may improve muscle function and reduce cardiac remodeling [35], there is still much to learn regarding its ability to improve health outcomes for patients suffering from heart failure. One randomized control trial demonstrated improved B-type naturetic peptide after a 10 week treatment regimen of 10,000 IU ergocalciferol compared to placebo [35]. An observational study demonstrated that vitamin D3 supplementation over one year did not significantly change markers of cardiovascular disease, including: total HDL and LDL cholesterol, triglycerides, APO A-1, insulin resistance or inflammatory markers [64]. Another trial supported the correlation between severe vitamin D deficiency and cardiovascular mortality in hospitalized patients [65].

The most recent data seems to support the use of vitamin D treatment in hypertensive patients. While different analogs of vitamin D
have been clinically tested, the efficacies have not been fully determined. The future for use of vitamin D as treatment for cardiovascular illness is promising, but more trials are necessary to solidify its clinical purpose. More confirmatory data is necessary in order to discern the role that vitamin D can play in patients with heart disease.

**Cancer therapeutic**

The hormonally active form of vitamin D, calcitriol, demonstrates anti-proliferative effects *in vitro* by promoting differentiation in a variety of cancer cell types including prostate, breast, colon, skin, and hematological cells [66-68]; however, there remains much debate on the efficacy of vitamin D on reducing cancer risk and use as a cancer treatment [69]. In addition to the tissues previously mentioned, VDR is expressed in pancreatic islet cells, skin keratinocytes, mammary glands and other reproductive tissues.

Ubiquitous expression of VDR in multiple human tissues implicated in cancer development has made vitamin D a therapeutic target of interest. Clinical trials assessing the effectiveness of vitamin D have been conflicting. It has been suggested, one reason for the conflict may relate to the stage of disease when vitamin D treatment is initiated [69]. Vitamin D clinical trials involving patients with advanced stage disease were largely unsuccessful. Conversely, in a recent randomized clinical trial, vitamin D3 was shown to reduce prostate-specific antigen (PSA) in two dosing groups (10,000 IU/d, p=0.04 and 40,000 IU/d, p=0.19) and serum PTH in the highest dosing group (40,000 IU/d, p<0.0001) in men with localized prostate cancer [70]. One remaining concern is that the use of high dose vitamin D may result in hypercalcemia; however, controlled regulation of the hydroxylases involved in the metabolism of vitamin D may reduce this risk.

**Diabetes control**

The effect of vitamin D on glycemic control is still being discovered. There has been some debate about the role of vitamin D in the onset and treatment of diabetes [35,71,72], but recent studies are promising for its use in glycemic control. There have been numerous clinical trials that have successfully demonstrated its role. One study concluded that if patients consumed vitamin D-fortified yogurt twice a day for 12 weeks a significant improvement in glucose control was noted [73]. A meta-analysis found that there was a 55% reduction in risk of diabetes in patients with high serum 25-dihydroxyvitamin D concentration [74]. Another study showed a reduction of insulin resistance in south Asian women upon supplementation with vitamin D [75]. A recent prospective study also showed improvements in glycemic control with supplementation of vitamin D in Type I diabetic patients [76].

The pathogenesis of diabetes driven by insulin resistance or altered β-cell function has been linked to vitamin D deficiency in two recent studies [77,78]. In addition, 1,25-dihydroxyvitamin D has been shown to play a significant role in improving insulin sensitivity in critical tissues including liver, skeletal muscle, and adipose tissue, along with enhancing β-cell function [71]. Animal and human models have also demonstrated a correlation between vitamin D deficiency and Type 1 Diabetes [71,72]. Evidence continues to associate polymorphisms in the genes of vitamin D binding protein, VDR, and vitamin D 1α-hydroxylase with pathogenesis of insulin resistance [71,72].

Despite the overwhelming data to support vitamin D in diabetes, there have still been studies that show no benefit of vitamin D supplementation in diabetic patients [79,80]. The benefit of vitamin D for the treatment of diabetes is becoming clearer, although more studies are required to determine standard dosing. Additional clinical trials will reveal a more in depth role for vitamin D in the treatment of diabetes.

**Conclusion**

The clinical role of vitamin D for treatment of non-bone diseases is still being evaluated. The top two causes of death, heart disease and cancer, are among the conditions for which vitamin D treatment is being considered. Vitamin D appears to play a significant role in many conditions associated with aging including cognition, depression, cancer, diabetes and cardiovascular disease. While the vitamin D data is more established concerning its association with depression, cognition and diabetes, conflict still remains in the role vitamin D plays in cancer and cardiovascular disease. Additional trials are needed to establish whether vitamin D therapy can treat or prevent these conditions in older adults. Because the amount and type of vitamin D therapy has varied substantially among prior clinical trials, future studies should focus vitamin D therapy on correcting vitamin D deficiency or maintaining serum vitamin D levels at a specific goal. This will allow more individualized therapy, which is necessary given the variability in individual vitamin D levels. If successful, vitamin D treatment could be an inexpensive strategy to reduce morbidity and mortality and improve quality of life for older adults.

**References**

Citation: Ralls VA, Boyer AP, Wilkins CH (2014) Vitamin D in Aging and Chronic Illness. Vitam Miner 3: 125.


43. Sanders KM, Stuart AL, Williamson EJ, Simpson JA, Kotowicz MA, et al. (2010) Annual high-dose oral vitamin D and falls and fractures in older women: a randomized controlled trial. JAMA 303: 1815-1822.


