Vitamin D deficiency in postmenopausal, healthy women predicts increased cardiovascular events: a 16-year follow-up study

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Abstract

Objective: To investigate the relationship between vitamin D status in healthy women and cardiovascular outcome.

Design and methods: Between 1990 and 1993, 2016 healthy, recently postmenopausal women were enrolled in the Danish Osteoporosis Prevention Study. Serum levels of 25-hydroxyvitamin D (25(OH)D, nmol/l) were measured at baseline. Participants were followed for 16 years. The primary end point was a combination of death, heart failure, myocardial infarction (MI) and stroke. Vitamin D deficiency was defined as serum 25(OH)D < 50 nmol/l. The primary end point was adjusted for other risk factors of adverse cardiovascular events (age, smoking, blood pressure, hip–waist ratio, education and family history of MI).

Results: At baseline, mean age was 50 years and BMI 25. Women with vitamin D deficiency (n = 788) had more cardiovascular risk factors than vitamin D-replete women (n = 1225). Compared with vitamin D-replete women, women with low 25(OH)D levels had significantly higher BMI and triglycerides, lower HDL and hip–waist ratio and less education. More were smokers among the vitamin D deficient (47 vs 38%). A primary end point was experienced by 118 (15%) with vitamin D deficiency and by 125 (10%) of the vitamin D-replete. Hazard ratio (HR) was 1.49 (95% confidence interval: 1.16–1.92; P = 0.002) in the vitamin D deficient. Adjusted HR was 1.32 (1.02–1.71; P = 0.03). In total, 135 women died; of these, 65 (8%) were of the vitamin D deficient and 70 (6%) in the vitamin D-replete group; unadjusted HR was 1.44 (1.02–2.01; P = 0.04) for vitamin D deficiency.

Conclusion: Healthy women with vitamin D deficiency have increased risk of adverse cardiovascular outcome.

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Introduction

Recently, a large observational study found vitamin D deficiency to be associated with an increased risk of coronary artery disease, diabetes, cardiomyopathy, hypertension and all-cause death (1). The theory of a relationship between vitamin D and cardiovascular disease (CVD) and mortality is not new (2, 3, 4, 5, 6, 7, 8). The main dispute remains whether vitamin D is just a marker for a worse prognosis (9, 10) potentially explained by other factors, as vitamin D deficiency has also been associated with obesity (1, 2, 11), smoking (2), ageing (1, 12, 13) and less physical activity (2, 11). Though the major source of vitamin D is dermal production triggered by the sun (3, 14), vitamin D is also ingested and could thus be a marker of general diet composition. In this study, we have had the opportunity to adjust for various lifestyle factors including degree of education, leisure time physical activity and importantly diet. CVD may have a different pattern in men and women, and studying men and women separately eliminates some of the variation.

The aim was to investigate whether baseline vitamin D status in healthy postmenopausal women predicted cardiovascular outcome or mortality, when adjusting for important confounders that are common risk factors for vitamin D deficiency and CVD.

Materials and methods

Study subjects

The Danish Osteoporosis Prevention Study (DOPS) is a prospective multicentre trial evaluating the effect of postmenopausal hormone therapy (HT) as primary
prevention of osteoporotic fractures. Women included in the study fulfilled the following criteria: healthy, recent postmenopausal Caucasian women aged 45–58 years, with last menstrual bleeding 3–24 months before study entry or perimenopausal symptoms (including irregular menstruations) in combination with postmenopausal serum FSH (FSH more than 2 S.D. above the premenopausal mean). Hysterectomised women were included if aged 45–52 years, provided they had elevated postmenopausal serum FSH. Exclusion criteria were history of bone disease (including non-traumatic vertebral fractures on X-ray), uncontrolled chronic disease, previous or current cancer or thromboembolic disease, current or past treatment with glucocorticoids > 6 months, current or previous HT use within the past 3 months or alcohol or drug addiction. Between 1990 and 1993, 2016 women were enrolled in a prospectively followed cohort, and of these, 1006 were randomly allocated to receive either HT or no treatment (502 on HT); the remaining 1010 women had personal choice (221 opted for HT). Two thousand and thirteen had baseline measurements of 25-hydroxyvitamin D (25(OH)D). Recruitment has previously been described in detail (14). Subjects gave informed consent before the study, which was conducted according to Helsinki II. The study was approved by the Central and Regional Ethics Committee (# 1990/1821) and monitored by the Good Clinical Practice (GCP) unit at University of Aarhus, Denmark (ClinicalTrials.gov NCT00252408).

**Biochemistry**

Blood samples were collected at baseline after overnight fasting and serum was stored at −80 °C for later analysis. Serum levels of 25(OH)D (nmol/l) were measured by RIA, preceded by specific extraction procedures with ether, preliminary chromatography and a competitive binding assay, using rat kidney-derived receptor binding protein (15). This method includes both vitamin D2 and D3 metabolites. The limit of detection was 12.5 nmol/l; intra- and interassay coefficients of variation were 8.3 and 10.2% respectively.

Serum levels of intact parathyroid hormone (PTH) were measured at inclusion in 1095 subjects by DPC Immulite (chemiluminescense) with an interassay coefficient of variation of 11% and an intra-assay coefficient of variation of 6%. Due to the smaller fraction of women with baseline measurements of PTH, it is not included in the final model. Baseline level of triglycerides was measured in 1437 women (71%) and thus is also not included in the final model.

**Diet records**

The participants recorded daily intakes of food and beverages after being given oral and written instructions at baseline using 4- or 7-day diet records. Intakes did not differ between 4- and 7-day food records (16). A trained dietician used food models and photographs during a 15-min individual validation interview to evaluate serving sizes and cooking habits. The diet content of calcium and vitamin D was analysed using a computerised database (‘Dankost’, software version 1.3b), containing information on diet composition based on official Danish food tables. The variation coefficient for intake of calcium from one measurement to the next was 15%, and for vitamin D intake, it was 28%.

**Clinical measurements**

At baseline, BMI (kg/m²) was calculated from height and weight. Subjects were interviewed regarding lifestyle habits, including physical activity and gynaecological history. Family history of myocardial infarction (MI) was noted.

**Assessment of mortality and hospitalisation due to CVD**

The primary end point for this study was a combination of death, hospitalisation due to MI, heart failure (HF) or stroke. Secondary end points were the individual components of the primary end point.

On June 16, 2008, we ended our follow-up period by retrieving data on all studied subjects from the Danish Civil Registration System and the National Hospital Discharge Register. In the Danish Civil Registration System, we identified all women who had died or emigrated during follow-up, as this register has electronic records on all changes in vital status, including change of address and date of death or emigration for the entire Danish population since 1968.

We identified women who had been hospitalised due to a CVD using the Danish National Hospital Discharge Register, which covers all contacts to the hospitals. The register was founded in 1977 and includes information on discharge diagnoses and date of discharge assigned exclusively by the physician at discharge according to the Danish version of ICD-8 until the end of 1993 and to the Danish version of the ICD-10 from 1994. The register has nationwide coverage of hospitals with an almost 100% completeness of recordings and a high precision of diagnoses (17). Using this register, we identified all study participants who had been assigned a diagnosis of CVD – classified as MI (ICD-10 code I21), HF (ICD-10 code I42-I50) and stroke (ICD-10 code I60-I69). For the combined end point, the date of first incident was used.

**Statistical analysis**

Unless otherwise stated, all data are expressed as mean and S.D. Vitamin D deficiency is defined as 25(OH)D
Results

Baseline characteristics of vitamin D-deficient and vitamin D-replete women are given in Table 1. Mean age was 50 years, and 788 (39%) women had vitamin D deficiency.

Compared with vitamin D-replete women, women with vitamin D deficiency had lower hip–waist ratio and HDL and higher BMI, triglycerides, PTH and a trend towards higher fasting blood glucose. The vitamin D-deficient women were less physically active and less educated, more had a family history of MI and more were smokers. In addition, total daily energy intake as well as dietary intake of protein, calcium and vitamin D differed between groups with higher intakes in the vitamin D-replete compared with the vitamin D-deficient women (Table 1). Only 11 women took calcium supplements, while 55% took over-the-counter vitamin tablets – mainly multivitamins – containing 5 μg vitamin D. Dietary intake of vitamin D was primarily from meat, fish and eggs. Age was not correlated with 25(OH)D in this population. Levels of 25(OH)D was associated with intake of several nutrients, but none of the nutrients had any influence on the primary outcome. PTH correlated with 25-OHD and the Pearson correlation coefficient was −0.164, P < 0.001.

Clinical outcome

One hundred and twenty-five (10%) vitamin D-replete women and 118 (15%) vitamin D-deficient women experienced a primary end point. Overall CVD rates were low: 135 women died and few were hospitalised due to a MI (n = 35). HF (n = 18) or stroke (n = 89) during the 16 years of follow-up.

Primary end point

The composite end point (mortality, MI, HF or stroke) was experienced by 243 women: 118 (15.0%) with vitamin D deficiency and 125 (10.2%) who were vitamin D replete. Accordingly, as shown in Table 2 and Fig. 1, vitamin D deficiency was associated with an increased risk of the primary end point (HR: 1.32; CI: 1.02–1.71, P = 0.034; Table 2). Adding PTH to the final model did not change the risk estimate of vitamin D deficiency (HR: 1.30, CI: 0.92–1.85).

Levels of triglycerides at baseline were also associated with an increased risk of the primary end point; however, if triglycerides were added to the model, it did not change the risk estimate for vitamin D deficiency. Leisure time physical activity was significantly associated with 25(OH)D (P < 0.0001) but did not correlate with any of the outcome measures and hence was not included in the final model.
Smoking was in univariate analysis associated with a HR of 2.26 (CI: 1.75–2.93, P < 0.0001) and each cigarette or cheroot/day was associated with a HR of 1.043 (CI: 1.031–1.056, P < 0.0001) for the primary end point. Smoking (as a dichotome variable) as well as number of cigarettes/cheroots per day correlated inversely to 25(OH)D (P < 0.001). Table 3 shows the HRs associated with vitamin D deficiency for the primary end point, stratified by smoking status. For other predictors of the primary end point included in the final model, please see Table 4.

Secondary end points In univariate analysis, vitamin D deficiency was associated with an increased risk of death; 70 (5.7%) vitamin D-replete women and 65 (8.3%) vitamin D-deficient women died (HR: 1.44, CI: 1.02–2.01, P = 0.04; Fig. 2), but after adjustment, risk was no longer significantly increased (HR: 1.18, CI: 0.84–1.67, P = 0.34). HF was diagnosed in 18 participants, ten (0.7%) vitamin D replete and eight (1.3%) vitamin D deficient (HR: 1.97, CI: 0.78–4.99, P = 0.15), MI in 35 participants (22 (1.8%) and 13 (1.7%) respectively), HR: 0.93 (CI: 0.47–1.84, P = 0.82) and stroke in 89 women (42 (3.4%) vitamin D replete and 47 (6.0%) with vitamin D deficiency), HR: 1.76 (CI: 1.16–2.67, P = 0.0075). There was no interaction between vitamin D deficiency and any of the other variables in the model.

Table 2 shows the number of events and the risk of the individual component of the primary end point. In crude as well as adjusted analysis, risk of stroke was significantly increased in the vitamin D-deficient women, whereas vitamin D deficiency was not associated with increased risk of hospitalisation due to HF or MI.

Discussion

In this population of 2013 healthy postmenopausal women, we found baseline 25(OH)D to be a predictor of later mortality or cardiovascular events. Even though sufficient 25(OH)D levels are associated with a healthier lifestyle, adjusting for important parameters such as smoking, blood pressure, age, family history of MI and hip–waist ratio did not obliterate the association. Importantly, we had information about physical activity and diet, but these were not – as often suggested – significant confounders in relation to 25(OH)D and outcome.

As 25(OH)D levels are inversely correlated with increasing tobacco use (2), it could be speculated that 25(OH)D levels in regard to cardiovascular outcome is a marker of smoking habits. However, by dividing the population into smokers and non-smokers, we found that vitamin D deficiency was associated with a higher HR (not statistically significant) for the primary end point in non-smokers than in smokers, and thus no indication of the suggested interpretation.

It is not surprising that vitamin D deficiency is associated with mortality and adverse cardiovascular events. Vitamin D has important physiological functions beyond bone and calcium metabolism; thus, vitamin D receptors are expressed in skeletal and in cardiac muscle, indicating a potential functional role (18). Vitamin D deficiency leads to reduced calcium absorption and impaired muscle function (19, 20). Furthermore, the activated form of vitamin D, 1,25(OH)2D, has been found to inhibit rennin–angiotensin activation (21, 22), reduce cardiac myocyte hypertrophy (23) and reduce inflammation (24, 25).
The mean intake of vitamin D in DOPS women was 3.06 μg/day, which is markedly lower than the recommendations by IOM of 600 IU (15 μg) per day (recommending up to 4000 IU/day as tolerable) (26). Even the vitamin D-replete women had a mean intake of vitamin D of < 5 μg/day. We did not see any effect of the intake of vitamin D on the primary outcome in our material – this is in line with previous findings (27). Furthermore, there was no effect of calcium intake on the primary end point nor on mortality or MI. These findings may contrast with earlier studies finding increased risk of MI (and possibly, though not significantly, mortality) with calcium supplements (28). However, in a recent study of 38 772 older women, a reduced risk of mortality was reported with the use of calcium supplements (29). Also, a recent meta-analysis of randomised controlled trials supports the positive effects of vitamin D along with calcium, whereas vitamin D supplements alone were not found to reduce mortality (30). In another recent review and meta-analysis, the authors did not find significant reductions in mortality nor in cardiovascular risk associated with vitamin D supplementation (31). In most of the trials, the included subjects were not vitamin D deficient and doses were small (≤ 800 IU/day). Thus, only in a limited selection did vitamin D-deficient subjects reach vitamin D repletion, and intervention was not long. Together, the evidence of effect of vitamin D on mortality or cardiovascular outcome still remains to be evaluated in large randomised controlled trials including vitamin D-deficient subjects and larger doses (≥ 50 μg/day) of D3 for a prolonged period. Other meta-analyses describe reduced mortality with vitamin D supplements in randomised trials (32, 33).

The increase in cardiovascular morbidity with vitamin D deficiency seen in DOPS as well as in other populations, where up to a double 5-year risk of coronary heart disease with severe vitamin D deficiency (34) has been reported, seem biologically plausible. Vitamin D (25-OHD) is related to vascular endothelial function in healthy older adults (24). In 554 healthy subjects, 25(OH)D correlated with vascular function in subjects without traditional risk factors as well as in subjects with a number of risk factors, after adjusting for several confounders. Moreover, in subjects, that corrected vitamin D insufficiency an improvement in vascular function was seen (35). In a randomised trial, vitamin D supplements reduced arterial stiffness (36). Also in patients with type 2 diabetes and vitamin D deficiency, vitamin D treatment improved endothelial function (37). This may, partly, account for the association between vitamin D deficiency and hypertension (1, 38, 39).

Vitamin D supplements to elderly women have shown to increase the oxidative muscle enzymes citrate synthase and 3-hydroxyacyl-CoA-dehydrogenase – this is a marker of improved muscle function (40). Also in children, the association between vitamin D and cardiac morbidity seems to be present – as seen in the NHANES study where vitamin D deficiency was associated with hypertension, low HDL-cholesterol levels and elevated PTH levels (39). Another possible cardio-protective effect of vitamin D may be via lipid regulation, as the association between vitamin D deficiency and a less favourable lipid profile has been reported earlier (1, 2, 11, 13, 41). The positive correlation between 25(OH)D and HDL in postmenopausal women also seen in DOPS has previously been investigated and the subclass of large HDL was accounting for the difference (42). The positive correlation between 25(OH)D and triglycerides are in line with the increased intima media thickness seen in subjects with carotid plaques from The Northern Manhattan Study (43). Finally, vitamin D may also act – indirectly – through PTH. Vitamin D deficiency leads to reduced calcium absorption with a tendency towards hypocalcaemia, which stimulates PTH secretion. Thus, decreasing vitamin D leads to increasing levels of PTH (44). Increased PTH for longer periods of time may indeed have harmful effects on the cardiovascular system (7, 45, 46, 47, 48, 49). The possible effects of vitamin D on the cardiovascular system have been discussed thoroughly in several overviews (50, 51, 52, 53).

Some studies have found the highest levels of 25-OHD to be associated with increasing mortality (26, 54, 55), but we did not find this association in our material (data not shown). This may be due to residual confounding or differences of the populations, as the DOPS women were healthy at baseline, whereas other studies have included patients referred to testing for different reasons (54) or were population based (55).

### Table 3
**Occurrence of composite end point divided according to smoking habits (number (%)) in vitamin D-sufficient and -deficient groups with hazard ratio (HR) of composite end point or mortality in vitamin D-deficient compared with vitamin D-replete participants.**

<table>
<thead>
<tr>
<th>Composite end point</th>
<th>n (%) of subgroup</th>
<th>Vitamin D deficient</th>
<th>Vitamin D replete</th>
<th>HR (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Smokers</td>
<td></td>
<td>74 (20.1)</td>
<td>72 (15.8)</td>
<td>1.32 (0.97–1.83)</td>
<td>0.091</td>
</tr>
<tr>
<td>Non-smokers</td>
<td></td>
<td>44 (10.5)</td>
<td>53 (7.0)</td>
<td>1.52 (1.02–2.26)</td>
<td>0.041</td>
</tr>
</tbody>
</table>

### Table 4
**Predictors of the primary end point – results from multivariable analysis.**

<table>
<thead>
<tr>
<th>Full model</th>
<th>Composite end point</th>
</tr>
</thead>
<tbody>
<tr>
<td>n/events</td>
<td>1972 (240)</td>
</tr>
<tr>
<td>Vitamin D deficiency</td>
<td>1.321 (1.022–1.707)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>1.084 (1.037–1.133)</td>
</tr>
<tr>
<td>Blood pressure (mmHg)</td>
<td>1.014 (1.008–1.020)</td>
</tr>
<tr>
<td>Hip–waist ratio</td>
<td>0.248 (0.091–0.672)</td>
</tr>
<tr>
<td>Family history of MI</td>
<td>1.664 (1.069–2.527)</td>
</tr>
<tr>
<td>Smoking (y/n)</td>
<td>2.321 (1.775–3.007)</td>
</tr>
<tr>
<td>Degree of education</td>
<td>0.819 (0.675–0.993)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>HR (CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.003</td>
<td></td>
</tr>
<tr>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td>0.017</td>
<td></td>
</tr>
<tr>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td>0.043</td>
<td></td>
</tr>
</tbody>
</table>
A limitation is the observational nature of this study, which makes the causality uncertain. The population is – on the other hand – very well described at baseline, with several risk factors believed to be shared between vitamin D deficiency and cardiovascular disease, including family history, hip–waist ratio, blood pressure, smoking status, educational level, sedentary lifestyle and nutrition. Additionally, changes in intake of calcium and vitamin D supplements may have changed over time as these are available over-the-counter. This could potentially explain the lack of association between calcium intake and prognosis. However, it cannot explain the association between baseline 25(OH)D levels and prognosis. If important changes in intake occurred, this would tend to reduce the association between baseline values and long-term prognosis. Also, baseline vitamin D was only measured once at inclusion. Thus, patients may have seasonal fluctuations that would tend to diminish the association between 25(OH)D and outcome.

In conclusion, vitamin D deficiency is associated with lifestyle factors that have also been linked to adverse cardiovascular outcome. Even after adjusting for the variables that were important for the primary outcome, vitamin D deficiency was, in this population of healthy postmenopausal women, associated with increased risk of mortality, MI, HF or stroke.

Declaration of interest

PEiken: Advisory board of Nycomed, Novartis; speaker’s bureau with Amgen, Novartis, Eli Lilly, and GSK. J-EB Jensen: Advisory board and speaker’s bureau of Nycomed, Eli Lilly, Novartis, MSD, Amgen.

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Author contribution statement

LMosekilde contributed to the original design. This study was planned by LEJensen, LLSchierbeck and LKober. LMosekilde, PEiken, LS tilgren, CLTofteng, LRejnmark and JEB Jensen implemented the study at different sites. LRejnmark retrieved data from national registers. LRejnmark, LKober, CLTofteng and LISchierbeck organised the database. LLSchierbeck and LKober had principal responsibility for the statistical analysis of the data. LLSchierbeck, LKober, LRejnmark and JEB Jensen were active in the interpretation of the results. The report was drafted by LLSchierbeck and reviewed by all authors. All authors have approved the final report.

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Vitamin D deficiency and cardiovascular outcome


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