Predicted 25-hydroxyvitamin D score and incident type 2 diabetes in the Framingham Offspring Study

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ABSTRACT

Background: Accumulating evidence suggests that vitamin D is involved in the development of type 2 diabetes (T2D).

Objective: Our objective was to examine the relation between vitamin D status and incidence of T2D.

Design: We used a subsample of 1972 Framingham Offspring Study participants to develop a regression model to predict plasma 25-hydroxyvitamin D [25(OH)D] concentrations from age, sex, body mass index, month of blood sampling, total vitamin D intake, smoking status, and total energy intake. Using this model, we calculated the predicted 25(OH)D score for each nondiabetic participant at the cohort’s fifth examination to assess the association between the predicted 25(OH)D score and incidence of T2D by using Cox proportional hazards models.

Results: A total of 133 T2D cases were identified over a 7-y average follow-up. In comparison with individuals in the lowest tertile of the predicted 25(OH)D score at baseline, those in the highest tertile had a 40% lower incidence of T2D after adjustment for age, sex, waist circumference, parental history of T2D, hypertension, low HDL cholesterol, elevated triglycerides, impaired fasting glucose, and Dietary Guidelines for Americans Adherence Index score (hazard ratio: 0.60; 95% CI: 0.37, 0.97; P for trend = 0.03).

Conclusions: Our findings suggest that higher vitamin D status is associated with decreased risk of T2D. Maintaining optimal 25(OH)D status may be a strategy to prevent the development of T2D. Am J Clin Nutr 2010;91:1627–33.

INTRODUCTION

Vitamin D, a fat-soluble vitamin, plays a critical role in regulating plasma calcium concentration through effects on intestinal absorption and bone metabolism (1). Vitamin D is formed in the skin from 7-dehydrocholesterol during exposure to solar ultraviolet B (UVB) radiation (2). Although vitamin D can be derived from the diet, only a few foods naturally contain vitamin D, such as oily fish. However, in the United States and Canada, certain foods are fortified with vitamin D, such as milk, some cereals, bread, and orange juice (3). The circulating concentration of 25-hydroxyvitamin D [25(OH)D] is the common biomarker used to assess vitamin D status. Studies have shown that, in addition to vitamin D intake, 25(OH)D concentration is associated with age, sex, adiposity, latitude of residence, skin pigmentation, and season of blood sampling (4–7).

The relation between vitamin D status and bone health is well established, but there is accumulating evidence that vitamin D might have other functions, including involvement in the development of type 2 diabetes (T2D) (8). There are few prospective studies on the association between vitamin D status and risk of T2D, and evidence from these studies is limited by the use of vitamin D intake or a single measurement of 25(OH)D concentration as a surrogate for usual vitamin D status (9–11). Because vitamin D can be synthesized in human skin under sunlight exposure and thereby has seasonal variation, the use of vitamin D intake cannot capture an individual’s overall vitamin D status, whereas a single measurement of 25(OH)D does not take into account the within-person variation across different seasons.

In this study we sought to test the hypothesis that vitamin D status is inversely associated with subsequent risk of T2D using a predicted 25(OH)D score, which was derived from known potential determinants of plasma 25(OH)D concentration and aimed to remove the seasonal effect on vitamin D status by holding season constant. This method of developing a predicted 25(OH)D score has been previously used to relate vitamin D status to cancer risk (12).

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2 None of the sponsors had any role in the design of the study; the design was exclusively the work of the authors.
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SUBJECTS AND METHODS

Study population

The Framingham Study was initiated in 1948 as a longitudinal, population-based study of cardiovascular disease. In 1971, 5135 offspring of original participants of the study and their spouses were recruited to participate in the Framingham Offspring Study. Members of the Framingham Offspring Study have returned, on average, every 4 y for a physical examination, questionnaires, laboratory tests, and assessment of cardiovascular and other risk factors (13). A total of 3799 cohort members participated in the fifth examination cycle (1991–1995). Valid food-frequency questionnaires (FFQs) were available for 3418 participants.

Participants were excluded from these analysis on the basis of previous diagnosis of T2D (n = 208), fasting glucose concentration ≥7.0 mmol/L or 2-h postchallenge glucose concentration ≥11.1 mmol/L (n = 134), or current use of insulin or oral hypoglycemic medication (n = 4). We also excluded those individuals without body mass index (BMI) data (n = 6). After exclusions, 3066 participants (1402 men and 1664 women) remained eligible for the present investigation. The participants were essentially all white. Participants were followed across the sixth (1996–1998) and seventh (1998–2001) offspring cohort examinations for a mean of 7 y for the development of incident T2D (Figure 1). The Institutional Review Boards for Human Research at Boston University and Tufts Medical Center approved the study protocols.

Dietary assessment

Usual dietary intakes for the previous year were assessed at each examination by using the 126-item semiquantitative Harvard FFQ (version 88GP) (14). The questionnaires were mailed to participants before the examination, and the participants were asked to bring the completed questionnaire with them to their scheduled appointment. The FFQ consists of a list of foods with a standardized serving size and a selection of 9 frequency categories ranging from never or <1 serving/mo to >6 servings/d. Nutrient intakes were calculated at the Harvard Channing Laboratory by multiplying the frequency of consumption of each unit of food from the FFQ by the nutrient content of the specified portion. Separate questions about use of vitamin and mineral supplements and type of breakfast cereal most commonly consumed were also included in the FFQ. In addition to assessing specific vitamin D supplement use, we used information on vitamin D content from multivitamins. FFQs with reported energy intakes <2.51 MJ/d (600 kcal/d) for men and women, >16.74 MJ/d (4000 kcal/d) for women, or >17.57 MJ/d (4200 kcal/d) for men or with >12 food items left blank were considered invalid. The FFQ has been shown to be valid for both nutrients and foods (14, 15). The FFQ has also been validated specifically for vitamin D intake in relation to plasma 25(OH)D (16), and the key dietary sources of vitamin D (including milk and dark fish) correlated well between the FFQ and diet records (17).

Lifestyle variables

Height, weight, and waist circumference were measured while the subjects were standing. BMI was calculated as weight in kilograms divided by the square of height in meters (kg/m²). A physical activity score was calculated from the number of self-reported hours spent doing specific activities that were categorized and weighted according to oxygen consumption required to perform them (MET-h/d) (18). Additional measurements

FIGURE 1. A: Timeline for the Framingham Offspring cohort examinations. B: Participants included in the development and validation of predicted 25-hydroxyvitamin D [25(OH)D] score. C: Participants in the diabetic follow-up study. n a = number alive at exam, n p= number of participants at exam; FFQ, food-frequency questionnaire.
included age, sex, current smoking (none, <20 cigarettes/d, ≥20 cigarettes/d), and alcohol consumption.

**Laboratory measurements at the fifth examination cycle**

Blood samples were drawn after the participants had fasted for ≥8 h for measurement of fasting glucose and lipid profile at the fifth examination cycle. A 75-g oral-glucose-tolerance test was administered according to World Health Organization standards to measure 2-h postchallenge glucose (19). Plasma glucose concentrations were measured in fresh specimens with a hexokinase reagent kit (A-gent glucose test; Abbott Laboratories Inc, South Pasadena, CA); the intra-assay CV was <3%. Lipid measures included enzymatic measurement of triglyceride (20) and the measurement of the HDL-cholesterol fraction after precipitation of LDL and VLDL cholesterol with dextran sulfan (21). Blood pressure was measured twice after the precipitation of LDL and VLDL cholesterol with dextran sulfan and the measurement of the HDL-cholesterol fraction. Plasma glucose administered according to World Health Organization standards for the fifth examination cycle. A 75-g oral-glucose-tolerance test was administered to participants at fasting glucose.

**Ascertainment of incident T2D**

Incident T2D was defined as a fasting plasma glucose concentration ≥7.0 mmol/L or use of insulin or oral hypoglycemic drug therapy at the sixth or seventh study examinations.

**Statistical analysis**

**Development and validation of predicted 25(OH)D score**

Plasma 25(OH)D concentration was measured by radioimmunoassay (Diasorin, Stillwater, MN) in a subsample of 1728 participants in the Framingham Offspring Study during the period of July 1997 to May 1999, which included the latter half of the sixth and the beginning of the seventh examination cycles. Using data from the subsample with plasma 25(OH)D measurements, we developed a multiple linear regression model to predict plasma 25(OH)D concentration [predicted 25(OH)D score] from potential determinants of vitamin D status. Because we planned to apply this predicted score to a nondiabetic sample, we excluded subjects with T2D (n = 234) from the 1972 participants for the development of the multiple linear regression model. We also excluded 10 subjects with influential outlying values (plasma 25(OH)D concentrations >112.5 nmol/L). After these exclusions, a total of 1728 subjects were available for the development of the predicted 25(OH)D score. We randomly selected ~50% of the 1728 subjects (n = 883) for the development of the predicted 25(OH)D score. We used the remaining subjects (n = 845) in the subsample with 25(OH)D measurements to validate the predictive multiple linear model (Figure 1). Using multiple linear regression modeling, we related the measured 25(OH)D concentration to the following potential determinants of vitamin D status: age, sex, BMI, month of blood sampling, total vitamin D intake, physical activity score, smoking status, total energy intake, and alcohol consumption. To meet the assumption of a linear relation between our independent and dependent variables in our regression model, we transformed BMI using the reciprocal of BMI and total vitamin D intake using the square root of total vitamin D intake. We also examined all possible 2-way interactions between sex, BMI (reciprocal), and other potential determinants of vitamin D status. A potential determinant of vitamin D status was excluded from our final model if it was not a significant predictor of the plasma 25(OH)D concentrations in this subsample, and little additional variance in plasma 25(OH)D concentrations could be explained by adding it back into the model.

**Application of predicted 25(OH)D score**

We applied the multiple linear regression model to the entire cohort at the fifth examination cycle to calculate a predicted 25(OH)D score for each participant without T2D (n = 3066). Although 25(OH)D concentration is a good measure of contemporaneous vitamin D status, it may not be a good reflection of a person’s typical status throughout the year because of the strong seasonal effect of sunlight exposure on 25(OH)D concentration. Therefore, to remove the seasonal effect on the predicted 25(OH)D score, we treated month of the blood sample as a confounder and calculated the score assuming that all blood samples were collected in the month of May. We chose the month of May because the 25(OH)D concentrations from samples drawn in May for the subsample of individuals used in the development and validation of the score were similar to the average 25(OH)D concentration observed for these individuals (Table 1).

Subjects were followed from the fifth to the sixth and seventh examinations. The date of diagnosis was the examination visit date at which a new case of T2D was identified. We used hazard ratios (HRs) from Cox proportional hazards models to estimate relative risks and 95% CIs for incident T2D across the tertile category of the predicted 25(OH)D score. P values for trend were calculated by treating the median of the predicted 25(OH)D score in each tertile category as a continuous independent variable in the hazards models. The following risk factors, which have been previously shown to predict incidence of T2D in the Framingham Offspring Cohort (23), were included as covariates in the regression models: age, sex, parental history of T2D, hypertension, low concentration of HDL cholesterol, elevated concentration of triglycerides, and impaired fasting glucose. Given that BMI was included in the prediction model, we included waist circumference as a covariate in place of BMI. We also included the Dietary Guideline for Americans Adherence Index (DGAI) as a covariate to determine whether any association with vitamin D was a consequence of overall diet quality (24). Because of missing information for some covariates, our sample size for these analyses was 2956.

Proportional hazards assumptions were tested by modeling interaction terms between each of the variables and person-years of follow-up. None of the interaction terms was statistically significant (ie, P > 0.05), indicating that the HR for the variables was reasonably constant over time. On the basis of observations from an earlier study (8), we examined whether calcium intake from foods and supplements was an effect modifier of the association between the predicted 25(OH)D score and incident T2D. The interaction term was not statistically significant. Statistical analysis was performed using Statistical Analysis

RESULTS

Participant characteristics

Characteristics of the study participants used in the development, validation, and application of the predicted 25-hydroxyvitamin D [25(OH)D] score are shown in Table 1. The development and validation samples were comparable on all characteristics considered; none of the characteristics considered were significantly different between these 2 groups. Although no formal statistical comparison was made because of lack of independence between subsamples, participants in the application sample were younger than participants in the subsamples used to develop and validate the score, as would be expected based on the difference between the dates of the study examinations, and they also appeared to have lower vitamin D intakes and were more likely to be heavy smokers.

Development and validation of predicted 25(OH)D score

Our final predictive model included age, sex, reciprocal of the BMI, total energy intake, square root of total vitamin D intake, smoking status, and month of blood sampling as independent predictors of the 25(OH)D score. Physical activity and alcohol consumption were not significant independent predictors of 25(OH)D concentrations ($P > 0.05$), and together they explained only 0.03% of the additional variance in plasma 25(OH) D concentration. Therefore, we excluded them from the final model. No interaction was found between sex, BMI, and other variables in the model. After excluding those with missing values for the predictor variables, 805 subjects were left in the final model, and the model $r^2$ was 25.75% (Table 2). We then calculated the predicted 25(OH)D score for 769 of the 845 remaining participants with actual plasma 25(OH)D measurements and complete data on the predictor variables. In this validation subset, the correlation coefficient between the predicted 25(OH)D score (mean: 49.1 nmol/L) and the actual plasma 25(OH)D measurement (mean: 49.3 nmol/L) was 0.51 ($P < 0.001$). The slope relating actual to predicted 25(OH)D in the validation sample was 0.99 (SE = 0.06, $P < 0.001$). The regression constant was 0.77 (SE = 2.79, $P = 0.80$) (Figure 2).

Application of predicted 25(OH)D score

Using the multiple linear regression model, we calculated the predicted 25(OH)D score for 3066 participants without T2D at the fifth examination cycle. The mean of the predicted score was 47.8 and ranged from 18.8 to 68.8 nmol/L. Among the 2956 participants without T2D, we observed a significant correlation between the predicted and actual plasma 25(OH)D score ($r = 0.51$, $P < 0.001$). Our final model included age, sex, reciprocal of the BMI, total energy intake, square root of total vitamin D intake, smoking status, and month of blood sampling as independent predictors of the 25(OH)D score. Physical activity and alcohol consumption were not significant independent predictors of 25(OH)D concentrations ($P > 0.05$), and together they explained only 0.03% of the additional variance in plasma 25(OH) D concentration. Therefore, we excluded them from the final model. No interaction was found between sex, BMI, and other variables in the model. After excluding those with missing values for the predictor variables, 805 subjects were left in the final model, and the model $r^2$ was 25.75% (Table 2). We then calculated the predicted 25(OH)D score for 769 of the 845 remaining participants with actual plasma 25(OH)D measurements and complete data on the predictor variables. In this validation subset, the correlation coefficient between the predicted 25(OH)D score (mean: 49.1 nmol/L) and the actual plasma 25(OH)D measurement (mean: 49.3 nmol/L) was 0.51 ($P < 0.001$). The slope relating actual to predicted 25(OH)D in the validation sample was 0.99 (SE = 0.06, $P < 0.001$). The regression constant was 0.77 (SE = 2.79, $P = 0.80$) (Figure 2).
participants with all covariate information, we identified 133 incident T2D cases from the fifth to seventh examination cycle. The association between the tertile categories of the predicted 25(OH)D score and the risk of T2D is shown in Table 3. After adjustment for risk factors for T2D, including waist circumference, diet quality (as assessed by the DGAI), parental history of T2D, hypertension, low concentrations of HDL cholesterol, elevated concentrations of triglyceride, and impaired fasting glucose, individuals in the middle and highest tertile had a 30% and a 40% reduction in risk of development of T2D over the 7-y follow-up compared with individuals in the lowest tertile of the predicted 25(OH)D score, respectively (P for trend = 0.03). When we further adjusted for other potential confounders that were also used to calculate the 25(OH)D predicted score, including age and sex, the association persisted.

**DISCUSSION**

We used a subsample of participants with plasma 25(OH)D measurements to develop a model using known and suspected determinants of vitamin D status to predict 25(OH)D status in the larger cohort. This approach has been successfully used in other cohorts to examine the relation between predicted 25(OH)D concentrations and cancer risk (12). Consistent with the literature, we found that plasma 25(OH)D concentration was positively associated with total vitamin D intake and inversely associated with BMI (4, 25, 26). Men had higher plasma 25(OH)D than women, and smokers had lower 25(OH)D than nonsmokers (27, 28). Unlike previous studies (12, 26), physical activity, presumably a proxy of outdoor exposure to sunlight, was not a significant predictor for plasma 25(OH)D concentrations. Physical activity may not relate to UVB exposure in our sample because of the latitude of residence and the fact that only 11% of our 25(OH)D measurements were taken during the summer months (June, July, or August) when sunlight exposure is most effective for 25(OH)D production in the northeastern United States (29); we also did not measure factors such as clothing, use of sunscreen, and duration spent outdoors that affect the actual sunlight exposure during outdoor physical activity. In addition, we could not obtain information on winter residence for our study participants at the time of their study examinations.

By applying this model to the larger cohort, we found that the predicted 25(OH)D score was inversely associated the incidence of T2D after adjustment for established risk factors for T2D and insulin resistance, including waist circumference, parental history of T2D, hypertension, low HDL-cholesterol concentrations, elevated triglyceride concentrations, impaired fasting glucose, diet quality, age, and sex. The appropriateness of adjustments for age and sex, both components of the predicted score, is a potential limitation of the current study. Failure to adjust for these variables might result in residual confounding of the relation between the 25(OH)D score and T2D risk because age and sex are also established risk factors for T2D. However, such adjustment may also result in overadjustment, because these 2 variables are constituents of the predicted 25(OH)D score. Consequently, we presented the association without and with this adjustment. We found that the inverse association between predicted 25(OH)D and risk of T2D was not appreciably attenuated after further adjustment for age and sex. Therefore, the observed association between predicted 25(OH)D and T2D risk does not appear to be a consequence of confounding by age and sex.

The absolute values of the predicted 25(OH)D score should also be used cautiously because previous research has shown large differences in 25(OH)D concentrations using different assays or even using the same assay in different laboratories (30). However, we have only used our predicted score to rank participants and not to classify participants’ vitamin D adequacy.

The fact that we were only able to predict 26% of the variance in 25(OH)D concentrations using our score in the validation
TABLE 3
Median 25-hydroxyvitamin D [25(OH)D] score, number of cases and person-years, and hazard ratios (95% CIs) for type 2 diabetes across tertile (T) categories of predicted 25(OH)D score

<table>
<thead>
<tr>
<th>Predicted 25(OH)D score tertile categories</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>P for trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>Predicted 25(OH)D score (nmol/L)</td>
<td>42.3 (18.8–45.6)</td>
<td>47.9 (45.7–50.4)</td>
<td>54.3 (50.5–68.8)</td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>987</td>
<td>983</td>
<td>986</td>
<td></td>
</tr>
<tr>
<td>No. of cases</td>
<td>70</td>
<td>36</td>
<td>27</td>
<td></td>
</tr>
<tr>
<td>Person-years</td>
<td>6359</td>
<td>6441</td>
<td>6430</td>
<td></td>
</tr>
<tr>
<td>Model 1</td>
<td>1.00</td>
<td>0.72 (0.48, 1.10)</td>
<td>0.61 (0.38, 0.97)</td>
<td>0.03</td>
</tr>
<tr>
<td>Model 2</td>
<td>1.00</td>
<td>0.71 (0.46, 1.11)</td>
<td>0.60 (0.37, 0.97)</td>
<td>0.03</td>
</tr>
</tbody>
</table>

1 Values are medians; ranges in parentheses.
2 Values are hazard ratios; 95% CIs in parentheses.
3 Adjusted for waist circumference, parental history of diabetes, hypertension (≥130/85 mm Hg), low concentrations of HDL cholesterol (<1.04 mmol/L in men or <1.29 mmol/L in women), elevated concentrations of triglycerides (≥1.70 mmol/L), impaired fasting glucose (≥5.6 mmol/L), and Dietary Guidelines for Americans Adherence Index (n = 2956).
4 Adjusted as in model 1 and further adjusted for age and sex (n = 2956).

Our findings advance the hypothesis that vitamin D status is inversely associated with T2D risk. To further our understanding of this relation, future studies should examine the relation of long-term estimates of vitamin D status and T2D risk, identify the optimal vitamin D intake or 25(OH)D concentration associated with low risk of T2D, and elucidate the underlying mechanisms of the relation between vitamin D and T2D.
VITAMIN D STATUS AND TYPE 2 DIABETES

We are grateful to the Framingham Study participants and staff for data collection. We thank Gail Rogers for data management. The authors’ responsibilities were as follows—EL: study design, data preparation, statistical analyses, interpretation of results, and drafting of the manuscript; PJF, JBM, AGP, CDE, and NMM: study design, interpretation of the results, and editing of the manuscript; and SLB: provision of plasma 25(OH)D data and editing of the manuscript. All authors approved the manuscript. The authors stated that they had no conflicts of interest.

REFERENCES