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Vitamin D supplementation for prevention of cancer: The D2d Cancer Outcomes (D2dCA) study

Ranee Chatterjee^a, John K Erban^b, Paul Fuss^c, Rowena Dolor^d, Erin LeBlanc^e, Myrlene Staten^f, Patricia Sheehan^g, Anastassios Pittas^h, and D2d Research Group

^aDivision of General Internal Medicine, 200 Morris Street, 3rd Floor, Box 104427, Durham, NC 27701, USA

^bCancer Center, Tufts Medical Center, 800 Washington St, Box 245, Boston, MA 02111, USA

^cDivision of Endocrinology, Tufts Medical Center, 800 Washington Street, Box 268, Boston, MA 02111, USA

^dDivision of General Internal Medicine, 200 Morris Street, 3rd Floor, Box 104427, Durham, NC 27701, USA

^eKaiser Permanente Center for Health Research NW, 3800 N Interstate, Portland, OR 97229, USA

^fKelly Government Services for the National Institute of Diabetes and Digestive and Kidney Diseases, 6701 Democracy Boulevard, Room 6107, Bethesda, MD 20817, USA

^gSpaulding Rehabilitation Network, 300 1st Ave, Charlestown, MA 02129, USA

^hDivision of Endocrinology, Tufts Medical Center, 800 Washington Street, Box 268, Boston, MA 02111, USA

Abstract

Evidence on biological plausibility from mechanistic studies and data from observational studies suggest that vitamin D may be linked to risk of several types of cancer. However, evidence from clinical trials evaluating the effect of vitamin D supplementation on cancer risk is limited. The Vitamin D and Type 2 Diabetes (D2d) study is a multi-center, randomized, placebo-controlled clinical trial conducted to examine the causal relationship between oral vitamin D supplementation and development of diabetes among overweight adults with prediabetes. The D2d study provides a unique opportunity to assess the effect of vitamin D supplementation at a higher dose (4,000 IU/day) than has been used in other clinical trials with cancer outcomes, in a population at higher than

corresponding author ranee.chatterjee@duke.edu.

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average risk for cancer. This paper provides: (1) baseline characteristics of the D2d population included in the D2d cancer outcomes secondary study (D2dCA) and comparison to other large trials of vitamin D supplementation and cancer risk; (2) description of data that are being collected during the trial and the planned statistical analyses to test whether vitamin D supplementation at a dose of 4,000 IU/day has an effect on incident cancer overall, on incidence of certain types of cancer, and on incidence of precancerous lesions. Results of D2dCA will help guide future research and clinical recommendations related to vitamin D and cancer risk.

Keywords

vitamin D; cancer risk; D2d study; prediabetes

INTRODUCTION

Evidence on biological plausibility from mechanistic studies and data from observational studies suggest that vitamin D status may be linked to risk of several types of cancer.(1-19) However, whether vitamin D supplementation lowers risk of cancer is unknown. Vitamin D is an appealing approach for prevention of cancer because of its ease of administration and low cost. However, most studies reporting significant associations between vitamin D and cancer incidence are observational studies, which do not establish causality. Few large trials have been conducted to examine whether vitamin D supplementation influences risk of cancer, and many of these trials have significant limitations: relatively low dose of vitamin D; small sample size; short follow up time; limited target population (e.g., women only). (20-22) One trial in the U.S. tested the combined effects of 2,000 IU/day of vitamin D₃ and 1,500 mg/day of calcium compared to placebo on cancer risk in 2,303 post-menopausal women over 4 years of follow-up.(23) Women randomized to vitamin D₃ had a 30% lower risk of incident cancer, which was nearly statistically significant (hazard ratio 0.70; 95%CI 0.47, 1.02).(23) A post-hoc analysis of the Vitamin D Assessment (ViDA) study among 5,108 people from New Zealand found no difference in incidence of cancer with 100,000 IU/month of vitamin D₃ compared to placebo.(24) More recently, in a cohort of 25,874 older adults, the VITamin D and OmegA-3 Trial (VITAL) reported no effect of 2,000 IU/day of vitamin D₃ supplementation on incident cancer.(25) An ongoing large trial of over 21,000 Australian adults 65-84 years old will assess the health effects, including cancer, of 60,000 IU/month of vitamin D₃ vs placebo.(26)

While large trials have not shown an effect of vitamin D supplementation on risk of incident cancer, meta-analyses of earlier trials have shown significant reductions in cancer-related mortality with vitamin D supplementation.(21, 27) A reduction in cancer-associated mortality with 2,000 IU/day of vitamin D was also observed in the VITAL study in a post-hoc analysis that excluded cancers which developed within the first one or two years.(25)

Given inconsistent results from published trials, further study is needed to determine the effects of vitamin D supplementation on cancer incidence and cancer-related mortality, especially in higher-risk populations. The Vitamin D and Type 2 Diabetes (D2d) study is a multi-center, randomized, placebo-controlled clinical trial conducted to examine the causal

relationship between oral vitamin D supplementation and development of diabetes among overweight adults with prediabetes.(28) The D2d cancer outcomes secondary study (D2dCA) will assess the effect of 4,000 IU/day of vitamin D supplementation on incidence of cancer and precancerous lesions in the D2d population which is at higher risk for cancer than average.(29)

METHODS

Overview of trial design and oversight

D2d is a randomized, double-blind, placebo-controlled clinical trial conducted in 22 medical centers in the United States (www.d2dstudy.org/sites) to evaluate the safety and efficacy of oral administration of 4,000 IU/day of cholecalciferol (vitamin D₃) vs. placebo in people with prediabetes who are followed for incident diabetes for approximately 3 years after randomization.(28) The D2d study is primarily supported by the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK) and the Office of Dietary Supplements (ODS) of the National Institutes of Health (NIH) through a U01 cooperative agreement to Tufts Medical Center (principal investigator A.G.P.) where the D2d Coordinating Center is based. D2dCA is a secondary study to D2d to test the hypothesis that, compared to placebo, high-dose (4,000 IU/day) oral vitamin D supplementation reduces risk of incident cancer and pre-cancerous lesions in a cohort of overweight adults with prediabetes who are also at high risk for cancer.(29)

The study is approved and monitored by an independent Data and Safety Monitoring Board and the Institutional Review Board of each collaborating clinical research site.

Participants

Target participants are adults at risk for diabetes (i.e., with prediabetes), defined by the American Diabetes Association (ADA) in 2010 (30), as meeting at least 2-out-of-3 of the following: fasting plasma glucose (FPG) 100-125 mg/dL; 2-hour plasma glucose (2hPG) after a 75-gram glucose load 140-199 mg/dL; hemoglobin A1c (HbA1c) 5.7-6.4%. The other inclusion criteria are: age ≥ 30 years (≥ 25 years for American Indians, Alaska Natives, Native Hawaiians, or other Pacific Islanders) and body mass index (BMI) of 24–42 kg/m² (22.5–42 kg/m² for Asians). Key exclusion criteria include any glycemic criterion in the diabetes range (30); use of medications approved for treatment of diabetes; hyperparathyroidism; nephrolithiasis; bariatric surgery; use of supplements with vitamin D or calcium over study limitation (1000 IU/day or 600 mg/day respectively); medications or conditions that could interfere with absorption or metabolism of vitamin D; major cardiovascular disease; hypercalcemia; hypercalciuria; chronic kidney disease; and history of cancer (except for basal cell skin cancer) within the past 5 years. Treated prostate or well-differentiated thyroid cancer not expected to require any treatment over the next 4 years are not exclusions. Blood 25-hydroxyvitamin D concentration is not an eligibility criterion. For a complete list of eligibility criteria, see the Supplementary Appendix. The recruitment process relied primarily upon the use of electronic health records to identify potentially eligible adults who then had an in-person screening visit and, if qualified, a second screening

visit to determine final eligibility by measuring FPG, 2hPG, and HbA1c at the study's central laboratory.(31)

Intervention

Participants were randomized to take a single soft-gel once daily that contained 4,000 IU of vitamin D₃ or matching placebo. Randomization was blocked-stratified by site, body mass index (less than 30 or greater than or equal to 30 kg/m²) and race (White or non-White). The dose of 4,000 IU daily was chosen to balance safety, by staying within the tolerable upper limit, as defined by the Institute of Medicine,(32) and efficacy in terms of obtaining a large difference in blood 25-hydroxyvitamin D concentration between the treatment and control groups.

The vitamin D and placebo pills are prepared by Tishcon Corporation and shipped to the drug distribution center (Veterans Affairs Cooperative Studies Program, Albuquerque, NM) in bulk where they are packaged into sealed, serially numbered bottles identical in appearance and weight, and shipped to each site for storage and distribution to participants. Both the manufacturer and drug distribution center perform quality control analyses on each lot; and the drug distribution center performs periodic potency testing of each batch to ensure the amount of vitamin D stays within specifications (3,600 to 4,800 IU) throughout the 2-year recommended shelf-life. Site staff dispenses the study pills to participants twice a year at each scheduled visit using an Interactive Web Response System that ensures correct treatment assignment according to the randomization code while maintaining blinding. Unless stopped for a safety reason, study treatment continues until the last study encounter, even after participants meet the primary study outcome of diabetes. Bottles with unused pills are returned at the next visit. Adherence is assessed by pill count as the percentage of pills taken in relation to the number that should have been taken. Participants are asked to refrain from using diabetes or weight-loss medications during the study and to limit the use of outside-of-study vitamin D to 1000 IU per day from all supplements, including multivitamins. To optimize safety, the study also limits calcium supplements to 600 mg per day.

Participants have a study visit at month 3, month 6, and twice a year thereafter. Midway through the visits, an interim contact (by phone or email depending on each participant's preference) takes place to encourage adherence to the intervention and to evaluate for adverse event occurrence and changes to medical conditions, including cancer-related outcomes.

Incident Cancer and Cancer Mortality Outcomes

Throughout D2d, a new diagnosis of cancer is identified as part of a standard health questionnaire administered at each encounter (in-person visit or phone/email encounter, 4 times/year). The questionnaire includes three general questions: "Have you had any changes in your health? Have you seen a doctor or had any visit to a healthcare provider since your last visit? Have you had any medications changed or new medications added?" Further information is obtained for any positive response. Starting in July 2017, after recruitment was completed, all randomized participants are asked questions specific to cancer screening

and any diagnoses of cancer since the beginning of the study (“Have you been told you have cancer? Have you had a colonoscopy? Have you had a skin mole or lesion removed or biopsied? Have you had a breast biopsy? Have you had a prostate biopsy?”). These questions are then repeated as part of the standard health questionnaire at each subsequent encounter. A self-report of cancer, endoscopic colorectal examination, or a biopsy is entered in the electronic data capture (EDC) system; and supporting medical records (including pathology and procedure reports) are obtained and provided to the Cancer Clinical Events Reviewer for adjudication. To ensure that all potential cancers are captured, the Coordinating Center also reviews all adverse events reported by participants to identify events reported as cancer *or* those that may “harbor cancer” (e.g., pancreatic mass). The Coordinating Center then requests that sites submit such events and supporting documentation for adjudication, if not previously submitted as part of the health- or cancer-specific questionnaires.

For each cancer outcome, the Cancer Clinical Events Reviewer, a board-certified oncologist who is unaware of the treatment assignment, adjudicates and confirms the following information: whether there is sufficient information for adjudication; the type of cancer and whether it is cancer or precancer; laterality (right origin of primary, left origin of primary, not applicable); date of diagnosis; grade for breast, colon/rectum, lung (primary), and pancreas (adenocarcinoma) cancers; Gleason score for prostate cancer; hormone receptor status (estrogen receptor, progesterone receptor, human epidermal growth factor receptor-2) for breast cancer; and date of diagnosis. Because history of cancer, other than basal cell skin cancer or well-differentiated thyroid cancer or prostate cancer not expected to need treatment within the past 5 years, is an exclusion criterion, a diagnosis of cancer is considered “new” unless there is sufficient evidence to be classified as “recurrent.” If there is equivalent evidence, it is classified as “undetermined.” From prostate and breast biopsies and all endoscopic colorectal examinations, the following precancerous lesions are adjudicated: high-grade prostatic intraepithelial neoplasia for prostate; atypical ductal hyperplasia (ADH) or atypical lobular hyperplasia (ALH) for breast; and adenomatous polyp for colorectal. Table 1 describes the data sources and outcomes collected. The D2dCA study does not collect data on cancer stage because: (1) the study lacks power to draw insight from analyses by stage; (2) procedures to stage cancer accurately may have differed by site, and therefore staging might have been incomplete.

In addition to incident cancer, information on cause of all deaths in D2d is being collected. All reports of death are confirmed with medical records or death certificates, which are collected by the sites and sent to the Coordinating Center. Cause of death is being adjudicated both by the Cardiovascular Disease adjudicator and by the Cancer Clinical Events adjudicator.

Recognizing that there may be smaller than expected numbers of incident cancer and cancer-related deaths, we are currently exploring options to contact participants periodically by phone or email after study completion to continue collecting data on incident cancer and mortality for a longer period of time.

Covariates

Information on potential confounders or effect modifiers (e.g., age, gender, tobacco use) was collected for all participants by self-report at the baseline visits, which began in November 2013. A questionnaire to assess family history of cancer in first-degree relatives was administered to all participants beginning in January 2018. Current medication use is being collected throughout the study. At study initiation, only certain medications were entered in the database, and this did not originally include aspirin; however, aspirin use is being collected in source documents. In July 2017, sites were asked to retrospectively review concomitant medications in source documents and enter aspirin use (for regular regimens only, not occasional use) into the database. Subsequently, information on aspirin use is entered at every encounter. Use of non-steroidal anti-inflammatory drugs is not entered into the database because the benefit-burden ratio and reliability of data would be low. The International Physical Activity Questionnaire is administered at baseline, month 6 and yearly thereafter in all participants.(33) The Multicultural Food Frequency Questionnaire is administered at baseline in all participants and at month 12 and month 36 visits in participants free of diabetes.(34) Weight and height are measured at baseline and at every visit to calculate BMI.

Blood 25-hydroxyvitamin D level at baseline is not an eligibility criterion for the following reasons: (1) the definition of ‘optimal’ 25-hydroxyvitamin D is controversial and no consensus exists;(32, 35-42) (2) to ensure that the study is as “pragmatic” as possible and results are generalizable to clinical practice; (3) blood 25-hydroxyvitamin D level varies by season (32) and it is an acute-phase response;(43-45) (4) baseline 25-hydroxyvitamin D level is an important treatment selection marker in the subgroup analyses, and to assess its performance, participants with a wide range of 25-hydroxyvitamin D must be enrolled.(46) The study plans to measure serum 25-hydroxyvitamin D level at baseline, month 12 and month 24. If there is available funding, serum 25-hydroxyvitamin D level will also be measured at month 36 and 48.

The D2d cohort was recruited at a constant rate throughout the calendar year and cancer outcomes are captured 4 times/year; therefore, the potential for seasonal variability confounding the effect of vitamin D supplementation and cancer outcomes is low. All analyses will adjust for site, which approximates latitude at the participant’s residence.

Sample size calculations

As an event-driven trial, the parent trial will continue until the required number of diabetes events is reached.(28) Based on national registries (e.g., Surveillance Epidemiology and End Results [SEER]), and based on the observed characteristics of the D2d participants, the expected incidence for common non-basal cell cancers (e.g., breast, colon,) for the age (60-64 years) and gender (45% women) distribution for the D2d population is estimated to be 26.4 cases yearly, or approximately 1.1% per year. The incidence of cancer in the D2d overweight population with prediabetes is likely to be higher.(29) The parent study was powered based on the primary outcome of incident diabetes. Power calculations for the D2dCA study can be estimated based on predicted incidence of cancer in the placebo group (SEER data), fixed sample size of 2,423 participants, the median expected follow-up of 3

years, and different scenarios for relative risk reduction of cancer between study groups (Table 2). For example, if yearly incidence of cancer in the placebo group is 1.5%, at 3 years, the power to detect a 40% relative risk reduction in cancer incidence in the vitamin D group compared to placebo would be 0.65.

Assumptions: 2,423 participants randomized to placebo or vitamin D at 1:1 ratio; mean follow-up 36 months; 1.5% yearly incidence of composite cancer (excluding non-melanoma skin) in placebo arm; $\alpha = 0.05$

Statistical Analyses

Descriptive data will include means and standard deviations, medians, or percentages. Intention-to-treat analyses will compare groups defined by the randomization procedure and will include all participants and all events observed during the study, irrespective of adherence to assigned treatment. Follow-up time in the study will be calculated as time from randomization until the outcome of interest (cancer or precancer) is met, death, active withdrawal, or last encounter with available data. We will also conduct a “per-protocol” analysis that censors follow-up at the first time a participant did not follow the protocol; for example, stopped study pills or started out-of-study calcium or vitamin D supplements at doses above study limitations.

The main outcome for D2dCA is first diagnosis during D2d (i.e., since randomization) of any type of cancer (excluding non-melanoma skin cancers). Each participant with a new cancer will be counted only once. Two analyses for the primary outcome will be conducted: one analysis will include the entire cohort; another analysis will be within each gender (men, women) because the association between prediabetes/diabetes and cancer risk can differ by gender, and the effect of vitamin D on cancer risk may also differ by gender.(47, 48) A sensitivity analysis will be conducted after excluding cancers that were diagnosed within the first 6 months after randomization, as these cancers may be pre-existing and previously unrecognized.

Kaplan-Meier estimates will be plotted for each group. Cox proportional hazard models will be used to compare the hazard rate of incident cancer between the two groups.(49) The model will include group assignment as its main predictor variable. The variables (study site, BMI, and race) used to stratify the randomization will be included in the model. Additional Cox models will be evaluated, adjusting for other covariates of importance (e.g., history of cancer, smoking history, aspirin use). The p -value from the primary analysis will be based on the chi-square statistic from a likelihood ratio test obtained from proportional hazards models with and without the term for intervention arm. All randomized participants will be included in the analysis and no attempt will be made to impute missing data for participants with missed encounters. Additional analyses will be conducted for the most common types of cancer (e.g., breast, colorectal, prostate), though individual totals are likely to be very small.

Analyses will also be conducted for all precancers combined and for selected individual precancerous histologies (breast, colorectal adenomas, prostate) though individual totals are likely to be very small. (Table 1).

There is some evidence, while not consistent, that vitamin D supplementation may increase the risk of prostate cancer;(11, 15) therefore, a sensitivity analysis in the entire cohort will be done where the primary outcome will be defined as all cancers but will exclude prostate cancer. Similarly, inconsistent associations have been found between vitamin D and risk of pancreatic cancer;(18, 19) and so a sensitivity analysis will be conducted after excluding cases of pancreatic cancer.

Subgroup Analyses.—Variability of response to vitamin D supplementation will be assessed and defined by key baseline variables: calcium supplement intake (none vs any); serum 25-hydroxyvitamin D concentration (<12 or ≥12 ng/mL; <20 or ≥20 ng/mL; continuous), as serum levels could reflect total exposure to vitamin D sources, including sources outside of the intervention;(50) body mass index (<30 or ≥30 kg/m²); age (median value); race (white vs. Black vs. other); ethnicity (Hispanic or non-Hispanic); glycemic risk by number of prediabetes criteria met (two or three criteria); history of cancer; smoking history (never or ever). Each analysis of participant subgroups will include a test for interaction, and effect modification will be claimed only if the test for interaction reaches statistical significance. No adjustment will be made for subgroup analyses.

Two-sided *p*-values less than 0.05 will be considered statistically significant. Statistical analyses will be conducted using SAS software, Version 9.4 (SAS Institute Inc., Cary, NC).

BASELINE AND STUDY CHARACTERISTICS

From November 2013 through February 2017, 7,133 people were assessed for eligibility. Of these, 2,423 were randomly assigned to vitamin D or placebo and formed the intention-to-treat population of D2dCA.(51) The baseline characteristics for D2dCA and how this cohort compares to other large trials that have tested the effect of vitamin D supplementation on cancer risk are described in Table 3. Women (45%) and non-White participants (36%) are well-represented in D2dCA. Thirty-two percent of participants reported a family history of breast, colorectal, prostate, or lung cancer in a first-degree relative. Among participants reporting intake of outside-of-study supplements, mean vitamin D intake from supplements was 730 IU/day; and mean calcium intake from supplements was 311 mg/day. Data from the food frequency questionnaire are not currently available.

Compared to other trials testing the effect of vitamin D supplementation and cancer risk, D2dCA is unique in its participant population, comprised of overweight/obese people with prediabetes (Table 3). By contrast, the recently reported VITAL study included 13% of participants with diabetes.(52) Compared to other studies, D2dCA has greater racial and ethnic diversity.(23, 24, 52-54) Additionally, compared to other studies, D2dCA is unique in the dose of vitamin D₃ administered (i.e., higher) and in its frequent assessments of adherence and outcomes.(23, 52, 53). Participants in D2dCA were similar to the VITAL study participants in baseline use of vitamin D and calcium supplements.(24, 52)

DISCUSSION

The D2dCA provides a unique opportunity to assess the effect of vitamin D supplementation on risk of cancer and precancerous lesions in a population (overweight adults with prediabetes) at higher risk for cancer. The D2dCA has many strengths including detailed phenotyping of baseline characteristics, multiple time points of contact with participants to capture cancer events, and the higher dose of vitamin D compared to other vitamin D supplementation trials.(20-25) Although this population is at higher risk of cancer compared to other studies assessing the association between vitamin D and cancer risk, potential limitations of this study include low power to detect a significant association and the relatively short follow-up time.

Mechanisms linking prediabetes to cancer risk

Overweight/obesity and the conditions of prediabetes and diabetes have been associated with increased risk of cancer.(29, 55) However, for prostate cancer an inverse association has been described with a lower incidence of prostate cancer among people with diabetes.(47) In general, the associations between prediabetes/diabetes and cancer risk are thought to be linked through common non-modifiable and modifiable risk factors, particularly age, increased adiposity, and decreased physical activity. Additionally, there is evidence that hyperglycemia, hyperinsulinemia and adiposity (all of which are generally present in patients with prediabetes/diabetes) can promote carcinogenesis at the cellular level through a variety of mechanisms. Through direct effects or effects mediated through other molecular mediators including cytokines, adipokines, and hormonal pathways, the milieu of hyperglycemia, hyperinsulinemia and chronic inflammation can lead to DNA damage, mitogenesis, cancer cell proliferation, protection of cancer cells from apoptosis, invasion, and metastasis (Figure 1).(55, 56)

Mechanisms linking vitamin D to cancer risk

There are plausible mechanisms linking vitamin D to reduced cancer risk. Vitamin D is converted in the liver to 25-hydroxyvitamin D and then to its most biologically active metabolite, calcitriol, in the kidney by 1-alpha hydroxylase. Calcitriol has been implicated in many disease processes, including cancer, and many extra-renal organs express 1-alpha hydroxylase, which may further influence risk of carcinogenesis within these organs.(57-59) Calcitriol acts as a steroid hormone by binding to vitamin D receptors (VDR) and affecting gene expression, either promoting or inhibiting gene transcription. The calcitriol-VDR complex has also been found to exert cellular effects via non-genomic pathways.(60) Most cell types have VDRs, and the binding of calcitriol to VDRs results in a wide variety of effects that differ by cell type. With regards to the association between calcitriol and cancer, VDRs are found in most types of malignant cells, and the putative anti-neoplastic effects of calcitriol are thought to be mediated through its binding with VDRs. Through in vitro and animal models, several anti-neoplastic mechanisms have been identified, including: promotion of apoptosis or autophagy of cells, promotion of cellular differentiation, regulation of cellular proliferation, inhibition of angiogenesis, and prevention of invasion of malignant cells. Additionally, calcitriol has been found to decrease inflammatory reactions through mechanisms including inhibition of the production of inflammatory cytokines and

prostaglandins.(1-3) Vitamin D, through the anti-carcinogenic mechanisms described above, could counter-act the carcinogenic mechanisms promoted by the hyperglycemia, hyperinsulinemia and adiposity in patients with prediabetes (Figure 1). Finally, in addition to the direct genomic and non-genomic anti-carcinogenic effects of vitamin D, if vitamin D has an impact on diabetes risk through lowering glucose and/or insulin levels, it may additionally have indirect effects by reducing the carcinogenic processes which are promoted in a diabetic environment.

Review of observational studies and trials

Most human studies reporting on the association between vitamin D and cancer incidence are observational studies, (4-19) which preclude a definitive assessment of cause and effect because residual confounding or reverse causation cannot be excluded. Several large trials have reported on the effect of vitamin D supplementation with or without calcium on risk of precancers or cancers with inconclusive results.(23-25, 54, 61-65)

In the Women's Health Initiative Calcium and Vitamin D (WHI CaD) Trial (vitamin D₃ 400 IU/day and calcium 1,000 mg/day), there was no benefit of vitamin D on individual or total invasive cancer risk.(61-63) However, a reanalysis of participants within the WHI CaD study who did not take outside-of-study calcium and vitamin D supplements showed a significant reduction in total cancer risk and risk of breast cancer in the group randomized to calcium and vitamin D, as well as a non-significant reduction in risk of colorectal cancer.(66) The Vitamin D and Colon Polyp study (vitamin D 1000 IU/day with or without calcium carbonate 1200 mg/day) showed no significant benefit of vitamin D on colon polyp recurrence.(65) Two trials with combined vitamin D and calcium supplementation have been conducted on post-menopausal women in Nebraska.(23, 64) The earlier of the two trials demonstrated a reduced incidence of cancer for the participants in the calcium (1400-1500 mg/day) and vitamin D (1100 IU/day) arm compared to placebo (relative risk of 0.40; 95% CI: 0.20, 0.82); however, incident cancer was a secondary outcome.(64) The more recent trial by Lappe et al, designed with cancer as the primary outcome, provided highly encouraging but inconclusive evidence for an effect of vitamin D supplementation on cancer risk.(23) This trial randomized 2,303 women (mean age 65) to a combination of vitamin D₃ (2,000 units/day) and calcium (1,500 mg/day) or placebos and followed them for 4 years for incident cancer.(23) Risk of cancer was lower in the vitamin D/calcium vs. placebo group; however, the difference was not statistically significant (hazard ratio 0.70; 95%CI 0.45 to 1.02). Of additional interest, the achieved 25-hydroxyvitamin D level was inversely associated with cancer incidence ($p=0.03$, $\text{coef}=-0.017$).(23) Specifically, compared with 25-hydroxyvitamin D level of 30 ng/mL as baseline, the estimated hazard ratio for cancer incidence for 25-hydroxyvitamin D levels between 30 and 55 ng/ml was 0.65 (CI, 0.44, 0.97).(23) The study by Lappe et al was well conducted but there were several possible reasons for missing statistical significance (e.g., high baseline 25-hydroxyvitamin D, high intake of out-of-study vitamin D), as discussed by the authors. Secondary analyses of the earlier trial and of both trials combined, additionally combined with a volunteer cohort of similar participants, evaluated the association between vitamin D levels and cancer risk; and these analyses found lower risks of cancer with higher 25-hydroxyvitamin D levels.(67, 68)

In a post-hoc analysis of the ViDA trial from New Zealand (oral vitamin D₃ initial bolus dose of 200 000 IU and followed by 100,000 IU/month vs placebo), there was no difference between treatment arms in incidence of cancer.(24) Finally, the recently reported VITAL study evaluated the effects of vitamin D (2,000 IU/day) and omega-3 fatty acids (1g/day) on incident cancer and cardiovascular disease as co-primary outcomes.(25) The incidence in cancer overall was not different between treatment arms; however, in pre-specified subgroup analyses, there was a significant interaction between vitamin D and baseline BMI; compared to placebo, those participants taking vitamin D who were in the lower BMI group (<25 kg/m²) had a significant reduction in cancer incidence (hazard ratio: 0.76; 95% CI: 0.63, 0.90). Additionally, compared to placebo there was a non-statistically significant reduction among Black participants taking vitamin D (hazard ratio: 0.77; 95% CI: 0.59, 1.01)(*p* for interaction=.21).(25) Also, in a post-hoc analysis assessing cancer mortality, excluding the first one or two years of cancer deaths, compared to placebo, there was a significant reduction in cancer mortality among those participants taking vitamin D.(25) This reduction in cancer-related mortality has also been reported in systematic reviews of trials that tested the effect of vitamin D supplementation on cancer-related mortality.(21, 27)

Strengths

The D2dCA has several strengths in its design and conduct, as follows: a study population (overweight adults with prediabetes) at higher risk of cancer;(29, 55, 69) inclusion of both women and men; sizeable proportion of non-White participants; higher dose of vitamin D (expected to increase achieved 25-hydroxyvitamin D level) given daily (considered more physiologic than non-daily doses); (24, 25, 61) robust methods to collect events through direct contact with participants and adjudicate for cancer and precancerous lesions. Beyond the primary question of the effect of vitamin D supplementation on cancer, D2dCA will also provide data on the incidence of cancer and precancerous lesions, as detected in routine clinical practice, in a modern cohort of adults with prediabetes.

Limitations

While the D2dCA has many strengths, there are certain limitations. In D2dCA, cancer outcomes and precancerous lesions are assessed as post-hoc hypotheses with an anticipated median follow-up of approximately 3 years, which is a relatively short period for development of cancer; longer-term follow-up may be needed to uncover an effect, if present. Cancer screening procedures (e.g., colonoscopy) in study participants may vary across sites, which may introduce risk of reporting bias. To guard against this potential pitfall, analyses will adjust for study site at randomization. Finally, the D2dCA cohort includes overweight patients with prediabetes, which may limit generalizability; however, overweight and prediabetes are very prevalent (86 million Americans are estimated to have prediabetes) and this group is considered at high risk for cancer.

Conclusion

Based on highly consistent data from longitudinal observational studies, supportive evidence from mechanistic studies, and encouraging but inconclusive data from randomized trials, the D2dCA study will utilize the strengths of the D2d study to contribute important data on

causality by providing long-term clinical trial data regarding the role of vitamin D supplementation at a dose of 4,000 IU/day for prevention of cancer and precancerous lesions among participants with prediabetes who are at higher-than-average risk of cancer.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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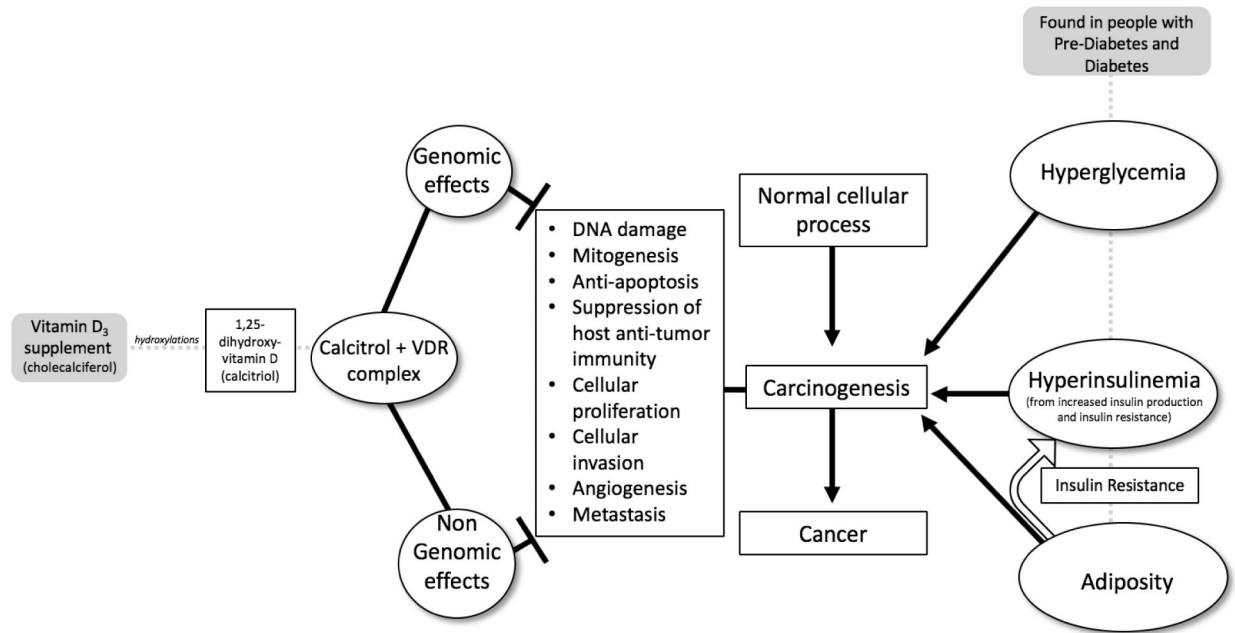


Figure 1.
Mechanisms of associations between prediabetes, vitamin D, and cancer risk

Table 1.

Primary data sources and cancer outcomes for D2dCA

Data sources
Prostate biopsies
Breast biopsies
Skin biopsies
Colonoscopy reports
Self-reported cancer or possible cancer diagnoses
Adjudicated cancer outcomes
Prostate
Breast (includes ductal carcinoma in-situ [DCIS])
Melanoma (including melanoma-in-situ)
Colon or rectum
Blood (l leukemia, and myeloma)
Lymphoma
Brain
Cervical
Endometrium
Gastric
Liver
Lung
Neuroendocrine
Ovary
Pancreas (adenocarcinoma)
Thyroid
Other cancer
Adjudicated precancers
Prostate (high-grade prostatic intraepithelial neoplasia)
Breast (atypical ductal hyperplasia [ADH], atypical lobular hyperplasia [ALH])
Colorectal adenomatous polyp

Table 2.

Power for possible scenarios of risk reduction in cancer incidence in D2dCA

Relative risk reductionfor vitamin D vs placebo (%)	Power
60%	0.964
55%	0.923
50%	0.856
45%	0.762
40%	0.647

Table 3.

Baseline characteristics and study intervention of D2dCA participants compared with other large trials of vitamin D and cancer

	D2dCA (n=2,423)	Lappe et al(23) (n=2,303)	VITAL(52) (n=25,874)	ViDA(24) (n=5,108)
Characteristic				
Vitamin D Intervention (and other components if applicable)	D ₃ , 4,000 IU/day vs. placebo	D ₃ , 2,000 IU/day and calcium 1500 mg/day vs. placebos	D ₃ , 2,000 IU/day vs. placebo (in a 2×2 factorial design with marine omega-3 fatty acid 1 g/dsupplements)	D ₃ , 200,000 IU bolus followed by 100,000 IU/month vs. placebo
Outcome definition (post-hoc or primary)	Incidence of all cancers, excluding non-melanoma skin cancers (post-hoc)	Incidence of all cancers, excluding non-melanoma skin cancers (primary)	Incidence of all cancers (primary)	Incidence of all cancers (post-hoc)
Frequency of outcome assessment (method)	4 times/year: 2 times/year in-person; 2 times/year by phone/email (questionnaire)	2 times/year, in-person (questionnaire)	Month 6, year 1, then yearly (mail-in questionnaires)	Monthly (mail-in questionnaires)
Median follow-up, years	3 (anticipated)	4	5.3	3.3
Target population	At risk for diabetes, men or women age 30 years (25 years for people of the following groups: American Indian, Alaska Native, Native Hawaiian, or other Pacific Islander)	Healthy, post-menopausal women age 55 years	Healthy men age 50 or women age 55 years	Healthy men or women, aged 50-84 years
Mean age, years	59.4	65.2	67.1	65.9
Women, no. (%)	1086 (44.8)	2303 (100)	13,081 (50.6)	2140 (41.9)
White race, no. (%)	1616 (66.7)	2291 (99.5)	18,047 (71.3)	4253 (83.3)
Mean body mass index, kg/m²	32.0	~ 30	28.1	28.5
Self-reported family history of cancer, %	Colorectal 8.6%; Breast cancer (women only), 15.4%; Prostate cancer (men only) 10.3%; Lung cancer 11.0%	Not available	Colorectal 13.4%; Breast cancer (women only) 17.7%; Prostate cancer (men only) 15.2%; Lung cancer 14.5%	Not available
Smoking, %				
Never	58.7	~ 67	51.7	51.0
Former	34.9	Not available	41.1	42.5
Current	6.5	~6	7.2	6.3
Vitamin D supplements outside-of-study allowed, IU/day	1000	800	800	600 if age 50-70 years; 800 if age 71-84years
Participants taking vitamin D supplements, %	42.6	Not available	42.6	8.0
Vitamin D intake from supplements among	730	~ 700	Not available	Not available

	D2dCA (n=2,423)	Lappe et al(23) (n=2,303)	VITAL(52) (n=25,874)	ViDA(24) (n=5,108)
participants taking vitamin D supplements, IU/day				
Participants taking calcium supplements, %	33.0	Not available	26.4	4.9
Calcium intake from supplements among participants taking calcium supplements, mg/day	311	600	Not available	Not available
Aspirin use, %	24.7	Not available	45.4	Not available