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Vitamin D, diet, and lifestyle in a chronic SCI population

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Abstract

Study Design—Cross-sectional study.

Objectives—Determine dietary, lifestyle, and clinical factors associated with plasma 25-hydroxyvitamin D [25(OH)D] levels in persons with chronic spinal cord injury (SCI).

Setting—Veterans Affairs Medical Center in Boston, MA.

Methods—174 participants completed food frequency and health questionnaires, provided a blood sample, and underwent dual x-ray absorptiometry (DXA) to assess %-total body fat. Linear regression models were used to assess cross-sectional associations of personal, lifestyle, and nutritional factors with plasma 25(OH)D.

Results—Independent factors positively associated with higher plasma 25(OH)D included vitamin D intake, age, hours of planned exercise, female sex, white race, wine consumption, and if a never or former smoker. The most important predictor of 25(OH)D was supplement intake. The

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Conflicts of Interest

All authors report no conflicts of interest.

Authors' Contributions

EG, AL, CGT contributed to study conception and design. EG, NEK, RLG, PW, and MMP contributed to data analysis. The draft manuscript was prepared by NEK, EG, RLG, and JEH. NEK, EG, RLG, and JEH edited and revised the manuscript. All authors read and approved the final manuscript.

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majority of subjects (98.9% for vitamin D and 74.1% for calcium) did not meet the recommended daily allowance for adults from their diet alone. Level and completeness of SCI, injury duration, mobility, %-total body fat, time outside, and comorbid diseases were not associated with plasma 25(OH)D.

Conclusions—Plasma 25(OH)D level in chronic SCI is not associated with clinical factors specific to SCI such as injury level and completeness, injury duration, and mobility mode, but related to supplement intake and other lifestyle factors.

Introduction

Vitamin D has been linked to positive effects on bone health, cardiovascular health, and pulmonary health, among other health benefits [1-7]. Vitamin D exists in two physiologically significant forms, D_3 (cholecalciferol) and D_2 (ergocalciferol). Less is known, however, about the determinants of vitamin D levels in humans beyond sun exposure, from which vitamin D_3 is synthesized [1]. Studies conducted in populations without spinal cord injury (SCI) suggest that a number of lifestyle and personal factors may positively influence vitamin D levels, including greater physical activity, never-smoking status, and normal weight [1,8,13].

The SCI population, in particular, is known to be at risk to have lower vitamin D levels [7,14–18]. Due to muscular weakness and paralysis that varies based on the severity of injury, persons with SCI are predisposed to a sedentary lifestyle. It is unclear, however, whether level and severity of SCI is associated with vitamin D status or if other factors related to diet or lifestyle are important. In addition, the importance of dietary intake and reliance on vitamin D supplements has not been examined in this group. Vitamin D sufficiency can be assessed by measuring plasma 25-hydroxyvitamin D [25(OH)D], which has been found to be a strong indicator of physiologically important vitamin D levels [1]. In this report, we assess the associations of a variety of clinical, dietary, and lifestyle factors with plasma 25(OH)D levels in persons participating in an SCI-health study.

Methods

Subjects

Between 5/31/2013 and 8/8/2016, 183 participants with chronic SCI taking part in a study to assess factors associated with respiratory and bone health at VA Boston were asked to complete a food frequency and dietary supplement questionnaire. Participants were recruited from persons receiving care at VA Boston, by direct mail to persons who had received care at Spaulding Rehabilitation Hospital, members of the National Spinal Cord Injury Association, subscribers to New Mobility Magazine, and from the greater Boston area through advertisement. Persons were eligible if they were 22 years of age or older, one or more years after injury, were not ventilator dependent, did not have a tracheostomy, and had no other neuromuscular disease. The Institutional Review Boards at VA Boston Healthcare System and Harvard Medical School approved the protocol and informed consent was obtained.

Food frequency and health questionnaires [19]

A validated semi-quantitative food frequency questionnaire (FFQ) developed at the Harvard University T.H. Chan School of Public Health [20–22] was mailed out to participants before their scheduled study visit to complete on their own. This standardized questionnaire asked the subject to report on the current use of vitamins, including vitamin D and calcium, and to recall average dietary intake of a wide range of foods and beverages over the past year by answering a series of questions about various serving sizes. The answers to the items on the questionnaire are converted to daily intake of various nutrients and partitions total vitamin D intake into supplements and dietary intake by computerized processing at the Harvard T.H. Chan School of Public Health Nutrition Department (https://regepi.bwh.harvard.edu/health/nutrition.html). Participants completed the FFQ before the visit on a separate day and brought it in with them. If the participant left the questionnaire at home or forgot to fill it out, an additional FFQ was provided if necessary and was mailed back in shortly after the visit.

A health questionnaire based on the American Thoracic Society Questionnaire [23] with additional questions was used to obtain a standardized history that included personal characteristics. These included tobacco smoking, beer, wine, and liquor consumption in the past year (frequency and quantity), usual mobility mode (motorized wheelchair, wheelchair, use of cane/walker, or ability to walk unassisted more than half the time), and physiciandiagnosed diseases such as asthma, chronic obstructive pulmonary disease (COPD), emphysema or chronic bronchitis, heart disease treated in the last 10 years, diabetes, and hypertension [24]. Persons were also asked how much time they spent in a typical week doing a planned exercise program and then asked to list the activity and how many minutes per week they participated. Examples of planned exercise included physical therapy, stretching and range of motion exercises, weight training, swimming, wheeling, walking, running, and sports. A sun exposure questionnaire based on recommendations from McCarty (2008) [25] was used to evaluate how many hours participants spent outdoors in a typical weekday and weekend day over the past month and how often they used sunscreen. Respondents were asked to report time spent outside between sunrise and sunset on a typical weekday and on a typical weekend day in the past month (no time outside, < 15 minutes/ day, 15 to 30 minutes/day, > 30 minutes to 1 hour/day, > 1 hour to 2 hours/day, > 2 hours to 4 hours/day, > 4 hours to 6 hours/day, > 6 hours to 8 hours/day, > 8 or hours/day) and sunscreen use (usually (50%), sometimes, and never).

Neurological exam, stature, and weight

Motor level and completeness of injury were assessed based on the American Spinal Injury Association Impairment Scale (AIS) by record review and by exam [26–27]. Motor incomplete SCI included AIS C (most key muscles below the neurological level grade < 3/5) or AIS D (most muscles below the neurological level grade 3/5). For analysis, participants were further grouped into cervical motor complete (AIS A or B) and cervical AIS C, others with motor complete (AIS A or B) and AIS C, and all others AIS D. Height was obtained by measuring the body length from top of the head to the heel of the subject in a supine position. If required, wheelchairs were weighed with and without the participant, and weight was subtracted to determine the participant's weight. Self-reported height and weight were

used if severe contracture or bracing hindered measurement (n=2). Body mass index was calculated from height and weight.

Body composition analysis

We used a 5th generation GE Healthcare iDXA scanner to assess total body composition, including total % body fat, using Encore 13.60 software. Scans were performed utilizing the same densitometry technician and analyzed by a certified clinical densitometrist (AAL). Daily quality assurance was performed utilizing a phantom.

Vitamin D analysis

EDTA plasma was drawn and immediately delivered to the core blood research laboratory at our facility. Samples were centrifuged for 15 minutes at 2,600 rpm (1459 \times g) at 4°C and stored at -80°C. Analysis of 25(OH)D was done at the Clinical & Epidemiologic Research Laboratory, Department of Laboratory Medicine at Children's Hospital in Boston by high performance liquid chromatography tandem mass spectrometry (HPLC-MS/MS). The assay is linear up to 100 ng/mL, and sensitive to 1 ng/mL. Day-to-day precision (%CV) at various levels of 25(OH)D ranged from 5.6% to 8.5%.

Analyses

We excluded 1 person with sarcoidosis, 6 persons whose plasma 25(OH)D was unavailable, and 2 persons whose plasma 25(OH)D levels were considered to be outliers (values of 71.9 and 86.3 ng/mL) leaving a sample size of 174 participants. General linear models (PROC GLM, SAS version 9.4) were used to assess both continuous and categorical determinants of 25(OH)D. Residual plots were examined to assess goodness of fit. Variables significant at the 0.100 level in univariable models were assessed in multivariable models and adjusted for laboratory batch. A sensitivity analysis was conducted excluding 18 persons whose responses on the FFQ conflicted with their responses to supplement use on the health questionnaire. The Tukey method was used to determine the significance between multiple (>2) groups in categorical variables. Monthly alcohol intake of beer, wine, and other liquor was estimated by multiplying usual daily or weekly intake by the frequency of drinking. Since wine consumption was the only alcohol type associated with plasma 25(OH)D, beer and liquor consumptions were grouped together to represent all non-wine consumption. Usual time outdoors over a week was estimated by multiplying the median of each weekday category by 5 and the median of each weekend day category by 2 (using 9 hours for the highest category), and summing to obtain the total hours. Season was assigned based on calendar month of study visit. Trends across wine consumption categories were assessed using the median values and for smoking categories by assigning ordinal numbers. To examine the patterns of predictors with the outcome, we used the 2011 Endocrine Society's Practice guidelines (vitamin D deficiency: 25(OH)D < 20 ng/mL; insufficiency: 20 to < 30 ng/mL; sufficiency: 30 ng/mL) [28].

Results

Personal, lifestyle, and nutritional characteristics of the population are shown in Table 1. The mean (SD) age was 57.6 ± 12.4 years, the majority of the population was male (81.0%) and

white (85.1%), and the mean BMI was $27.2 \pm 6.1 \text{ kg/m}^2$. The average injury duration of was 20.9 ± 12.9 years and 23.0% of participants had motor complete and AIS C SCI, 37.3% had lower levels of motor complete and AIS C SCI, and 39.7% had AIS D SCI. Most subjects used either motorized (20.7%) or hand-propelled wheelchairs (46.0%). Mean plasma 25(OH)D level was 31.2 ± 10.3 (range 10.1-56.2) ng/mL. Slightly more than half of the participants were vitamin D sufficient (53.4%), about a third were vitamin D insufficient (32.2%), and the rest were vitamin D deficient (14.4%). Participants consumed a median (25th, 75th percentile) 164 (109, 255) IU of vitamin D daily from their diet and a median 561 (57, 2000) IU from supplements. All but 2 participants (98.9%) did not meet the Institute of Medicine (IOM) recommended dietary allowance (RDA) of 600 IU of vitamin D for adults up to 70 and 800 IU for ages above 70 from their diet alone [29]. Similar to vitamin D, the majority of participants (n=129, 74.1%) did not reach IOM recommendations for calcium intake based on diet alone (1,000 mg of calcium for ages 31-70 and 1,200 mg for persons over the age of 70) with a median dietary calcium intake of 748 (534, 1029) mg. Most participants also did not drink beer or liquor (51.7%) or wine in the past year (69.0%), but, of those who drank, the median (25th, 75th percentile) number of beers or drinks of liquor was 12 (4, 36) drinks per month and glasses of wine was 12 (4, 24). In addition, almost half of the participants were former smokers (49.4%) and few were current smokers (9.2%). Persons who spent time outdoors between sunrise and sunset spent an average of 12.9 ± 12.5 hours during a typical week in the previous month, and those who exercised (61.5%) participated in 6.5 ± 5.6 hours of planned exercise per week. Among the 107 persons (61.5%) who reported planned exercise, 47 persons reported time participating in stretching, range of motion, or physical therapy, accounting for 19.9% of all exercise time, and 93 persons reported time spent in other exercise activities (80.1% of all exercise time).

In analyses assessing individual factors associated with 25(OH)D (Table 2), there was a statistically significant positive association between 25(OH)D with total vitamin D intake (Figure 1), age, years of education, total calcium intake, wine consumption, total hours of planned exercise, female sex, white race, and among never smokers, and users of sunscreen. Planned exercise reported as participating in stretching, range of motion, or physical therapy was not associated with 25(OH)D, whereas as other planned exercise was associated with 25(OH)D. There was no significant relationship between 25(OH)D and SCI level and completeness of injury, BMI, %-total body fat, mobility mode, comorbid medical conditions, time outside between sunrise and sunset, or season.

In a multivariable model (Table 3), age, total vitamin D intake, total hours of planned exercise, sex, race, wine use, and smoking status remained statistically significantly associated with 25(OH)D. Race and total vitamin D intake were the most statistically significant (p < 0.001) predictors. The regression coefficient for total vitamin D intake (Table 3) remained significant but was reduced by approximately 21% in the final multivariable model compared to the univariable model (Table 2). A multivariable model with the effects of vitamin D intake divided into dietary intake and supplement use was assessed, adjusting for age, planned exercise, sex, race, wine use, and smoking status. When comparing the impacts of vitamin D from diet and from supplements, supplement intake was statistically significantly associated with increased 25(OH)D (p < 0.001), but dietary intake was not (p = 0.305) (Table 3). We also assessed stretching, range of motion, or physical

therapy and other planned exercise as separate covariates. As in the univariable model, stretching, range of motion, or physical therapy was not associated with higher 25(OH)D level and other planned exercise was associated with higher 25(OH)D levels. In the multivariable model, several variables were no longer statistically significant, specifically sunscreen use, calcium intake from diet and supplements and education level. These variables were excluded from the final model.

As noted, the FFQ was not completed during the study visit when 25(OH)D levels were assessed, so responses regarding vitamin D intake, including from multivitamins, obtained at each study visit were compared to the information provided on the food frequency questionnaire. There were 17 persons whose responses on vitamin D or multivitamin supplement usage on the FFQ were discordant with their responses at the study visit. We repeated the multivariable models excluding these persons and noted no difference in the results. Laboratory batch was also not significantly associated with 25(OH)D in the final model. There were 32 persons who completed a visit between the study dates but who did not return a FFQ. There was no significant difference in level and completeness of injury or in available mean 25(OH)D levels compared to the responders.

Discussion

In this cross-sectional analysis of persons with chronic SCI, SCI level and completeness of injury, mobility, and duration of injury were not found to be determinants of 25(OH)D. Instead, other factors based on personal, nutritional, and lifestyle characteristics were found to be statistically significant predictors of 25(OH)D. Total vitamin D intake was found to be the most closely associated with plasma 25(OH)D levels.

Nutritional characteristics of the population appeared to be the most important determinant of vitamin D status. Higher levels of total vitamin D intake, which came from dietary and supplemental sources, were closely associated with higher levels of plasma 25(OH)D. A number of studies have assessed daily vitamin D intake in chronic SCI, in which they found intake levels to be inadequate [16–18,30]. Like these studies, we also found that the median vitamin D dietary intake for our population was below the IOM recommendation. Even though diet provides an important source of vitamin D, based on our population's dietary habits, supplemental usage was a more effective determinant of plasma 25(OH)D status than dietary vitamin D. Participants who were vitamin D sufficient were consuming considerably more supplemental vitamin D than participants who were vitamin D deficient. This is in contrast to persons across all levels of vitamin D sufficiency consuming similar amounts of vitamin D from their diet. As with vitamin D intake, calcium dietary intake also did not meet national recommendations for most individuals and supplements were necessary for the population to attain the recommended intake levels of calcium.

Certain lifestyle habits such as tobacco use and exercise had an association with 25(OH)D levels. Exercise has been shown to have a number of health benefits in SCI [31–33]. In our study, planned exercise also had a positive association with plasma 25(OH)D that appeared attributable to exercise other than physical therapy, stretching, and range of motion exercises that are applicable to patients with SCI. In addition to exercise, smoking status was

negatively associated with 25(OH)D, with the lowest levels in current smokers. Although the mechanism is uncertain, previous studies have similarly shown the negative effects of smoking on 25(OH)D levels in individuals without SCI [9,10,34].

In addition to diet and smoking status, the effects of wine drinking were also significantly associated with circulating 25(OH)D levels. Although wine contains no vitamin D, wine consumption displayed a positive association with plasma 25(OH)D specifically between non-drinkers and drinkers consuming greater than 12 drinks per month. A Swiss study in 2012 similarly reported greater levels of 25(OH)D amongst heavy wine drinkers [12]. It is unclear how wine may influence 25(OH)D levels, though it is suggested that certain phytoestrogens found in red wine may enhance the levels of 25(OH)D [35].

Personal characteristics such as age, sex, and race were also found to be closely associated with plasma 25(OH)D levels. It is unclear how greater age or female sex may be positively associated with vitamin D status, but it is believed that race performed as a proxy for skin tone in this study, which influences the amount of sunlight that is absorbed into the skin and catalyzes vitamin D_3 synthesis [1]. Previous studies have shown that persons with fairer skin complexions generally have higher levels of plasma 25(OH)D [36–37].

We found no association with BMI or %-total body fat even though lesser levels of plasma 25(OH)D have been associated with a greater BMI and total body fat in some studies [15,18]. It is possible that with a larger sample size we may have observed an association since participants who were obese had slightly lower levels of 25(OH)D. Although direct sun exposure was not measured in our study, the use of sunscreen was positively associated with 25(OH)D levels in the univariable model, possibly as a marker of greater sun exposure. Sunscreen use was no longer significant, however, in the multivariable model. Additionally, while our population reported spending an average of almost 2 hours per day outside between sunrise and sunset, time outside had no association with plasma 25(OH)D status. These results may be reflective of the participants all living in the New England area where direct sun exposure does not occur during the winter. We examined time outside between sunrise and sunset in only the months of April through September to concentrate on relevant sun exposure, but still found no significant association with 25(OH)D level. Moreover, there was no association found in 25(OH)D levels with season.

A strength of this study is the use of a validated food frequency questionnaire to assess the vitamin D and calcium intake. Moreover, our comprehensive assessment of participants provided us with a wide range of detailed information on multiple SCI related and personal factors that were used to find associations with 25(OH)D.

It is possible that questions in our health questionnaire could have better assessed direct sun exposure instead of time spent outside between sunrise and sunset. A well-defined assessment of direct sunlight exposure may have shown an association between higher levels of 25(OH)D and greater sun exposure like other studies have suggested. However, associations between measures of sunlight exposure and plasma 25(OH)D have not been consistently noted as described in a report that assessed ultraviolet radiation exposure based on residential address in three large non-SCI large cohort studies [38]. Additionally,

completion of the FFQ on the same date as the study visit or a detailed personal interview may have more accurately represented dietary intake at the time when the blood draw was collected and therefore could have increased the accuracy of the associations between plasma 25(OH)D and the vitamin D and calcium intake variables. Finally, subjects in the study had to be healthy enough to travel to the VA, which could have potentially limited our ability to study persons with greater levels of injury and completeness of SCI.

To summarize, plasma 25(OH)D levels were significantly associated with dietary intake in this chronic SCI cohort and were unrelated to injury level and completeness of SCI. Usual dietary intake was considerably less than the recommended daily intake and supplement intake was most closely associated with adequate levels. Other factors related to lifestyle including planned exercise, wine consumption, and smoking status were also determinants of 25(OH)D status. Our study suggests that clinicians may want to consider recommending vitamin D supplement use in SCI patients and to provide information to patients on personal lifestyle choices that may impact 25(OH)D levels.

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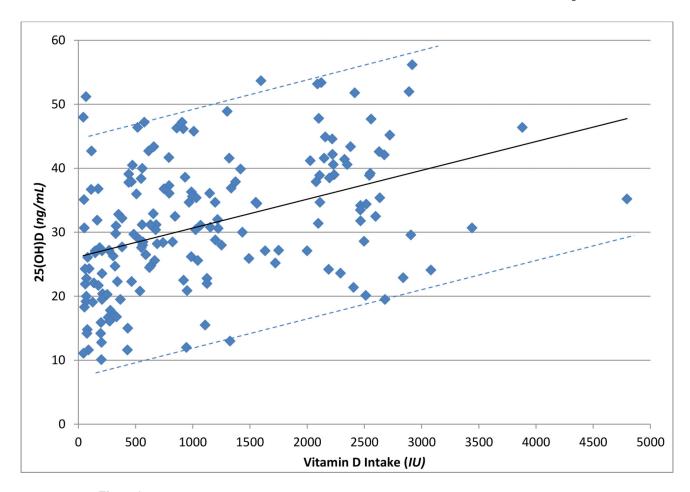


Figure 1. Plasma 25(OH)D levels and daily vitamin D intake with the fit line and 95% confidence intervals.

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Table 1

Characteristics of the population (n = 174)

	Plasma 25(OH)D Levels			Total (n) or
	Sufficient (n) [30 ng/mL]	Insufficient (n) [30 > n 20 ng/mL]	Deficient (n) [< 20 ng/mL]	Mean ± SD or Median (25% ^{ile} , 75% ^{ile})
(n)	93 (53.4%)	56 (32.2%)	25 (14.4%)	174
		Personal		
Sex				
Male	70 (40.2%)	49 (28.2%)	22 (12.6%)	141 (81.0%)
Female	23 (13.2%)	7 (4.0%)	3 (1.7%)	33 (19.0%)
Race/ethnicity				
White	88 (50.6%)	45 (25.9%)	15 (8.6%)	148 (85.1%)
Other*	5 (2.9%)	11 (6.3%)	10 (5.7%)	26 (14.9%)
Age (years)	58.6 ± 11.3	58.2 ± 14.1	52.8 ± 12.0	57.6 ± 12.4
BMI (kg/m²)	26.4 ± 6.0	28.2 ± 5.9	27.8 ± 6.4	27.2 ± 6.1
Obese (30)	20 (11.5%)	18 (10.3%)	7 (4.0%)	45 (25.9%)
Overweight (25 – < 30)	32 (18.4%)	21 (12.1%)	11 (6.3%)	64 (36.8%)
Normal/Underweight (< 25)	41 (23.6%)	17 (9.8%)	7 (4.0%)	65 (37.4%)
Total body fat (%)	36.6 ± 9.8	38.5 ± 8.6	37.7 ± 9.8	37.4 ± 9.4
Education (years)	15.4 ± 2.7	14.7 ± 3.8	13.9 ± 2.5	14.9 ± 3.1
Season of visit				
Winter	22 (12.6%)	12 (6.9%)	4 (2.3%)	38 (21.8%)
Spring	19 (10.9%)	10 (5.8%)	7 (4.0%)	36 (20.7%)
Summer	28 (16.1%)	16 (9.2%)	8 (4.6%)	52 (29.9%)
Autumn	24 (13.8%)	18 (10.3%)	6 (3.5%)	48 (27.6%)
SCI level and completeness				
Cervical motor complete & AIS C	20 (11.5%)	15 (8.6%)	5 (2.9%)	40 (23.0%)
Other motor complete & AIS C	38 (21.8%)	20 (11.5%)	7 (4.0%)	65 (37.4%)
All AIS D	35 (20.1%)	21 (12.1%)	13 (7.5%)	69 (39.7%)
Mobility mode				
Motorized wheelchair	19 (10.9%)	10 (5.8%)	7 (4.0%)	36 (20.7%)
Hand-propelled wheelchair	47 (27.0%)	26 (14.9%)	7 (4.0%)	80 (46.0%)
Walk with aid	9 (5.2%)	10 (5.8%)	4 (2.3%)	23 (13.2%)
Walk without assistance	18 (10.3%)	10 (5.8%)	7 (4.0%)	35 (20.1%)
Injury duration (years)	22.2 ± 14.3	18.6 ± 10.4	21.2 ± 11.9	20.9 ± 12.9

Comorbid diseases

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< 500

Plasma 25(OH)D Levels Total (n) or $Mean \pm SD$ or Insufficient (n) Deficient (n) Sufficient (n) Median (25%ile, 75%ile) [30 > n][< 20 ng/mL] 30 ng/mL] 20 ng/mL] 11 (6.3%) 5 (2.9%) 21 (12.1%) Asthma 5 (2.9%) COPD 12 (6.9%) 7 (4.0%) 1 (0.6%) 20 (11.5%) 11 (6.3%) OLD ** (asthma or COPD) 21 (12.1%) 5 (2.9%) 37 (21.3%) Diabetes 17 (9.8%) 6 (3.5%) 5 (2.9%) 28 (16.1%) Hypertension 36 (20.7%) 18 (10.3%) 9 (5.2%) 63 (36.2%) Heart disease, Last 10 Years 16 (9.2%) 5 (2.9%) 1 (0.6%) 22 (12.6%) Daily Intake, Median (25th, 75th percentile) Total vitamin D intake (IU) 1217 (655, 2220) 577 (231, 1224) 260 (125, 367) 834 (327, 2077) 2000 - 479636 (20.7%) 8 (4.6%) 1 (0.6%) 45 (25.9%) 1000 - < 200018 (10.3%) 10 (5.8%) 2 (1.2%) 30 (17.2%) 500 - < 100024 (13.8%) 15 (8.6%) 1 (0.6%) 40 (23.0%) < 500 15 (8.6%) 23 (13.2%) 21 (12.1%) 59 (33.9%) Vitamin D intake from diet (IU) 177 (109, 258) 150 (116, 229) 196 (88, 273) 164 (109, 255) 300 - 69220 (11.5%) 5 (2.9%) 4 (2.3%) 29 (16.7%) 200 - < 300 20 (11.5%) 12 (6.9%) 7 (4.0%) 37 (21.3%) 100 - < 2007 (4.0%) 68 (39.1%) 35 (20.1%) 26 (14.9%) < 100 20 (11.5%) 13 (7.5%) 7 (4.0%) 40 (23.0%) Vitamin D from supplements (IU) 1000 (400, 2000) 400 (50, 1064) 0 (0, 57) 561 (57, 2000) 2000 - 475030 (17.2%) 8 (4.6%) 1 (0.6%) 39 (22.4%) 1000 - < 2000 18 (10.3%) 8 (4.6%) 2 (1.2%) 28 (16.1%) < 1000 35 (20.1%) 27 (15.52%) 5 (2.9%) 67 (38.5%) None 10 (5.8%) 13 (7.5%) 17 (19.8%) 40 (23.0%) Total calcium intake (mg) 1396 (959, 1718) 1160 (633, 1427) 957 (504, 1123) 1206 (694, 1610) 2000 - 394317 (9.8%) 3 (1.7%) 20 (11.5%) 1000 - < 2000 51 (29.3%) 28 (16.1%) 12 (6.9%) 91 (52.3%) 500 - < 100017 (9.8%) 16 (9.2%) 7 (4.0%) 40 (23.0%) < 500 8 (4.6%) 9 (5.2%) 6 (3.5%) 23 (13.2%) Calcium intake from diet (mg) 863 (573, 1165) 702 (512, 949) 624 (504, 1057) 748 (534, 1029) 1000 - 276231 (17.8%) 10 (5.8%) 49 (28.2%) 8 (4.6%) 750 - < 1000 22 (12.6%) 13 (7.5%) 2 (1.2%) 37 (21.3%) 500 - < 750 25 (14.4%) 21 (12.1%) 9 (5.2%) 55 (31.6%) < 500 15 (8.6%) 12 (6.9%) 6 (3.5%) 33 (19.0%) Calcium from supplements (mg) 500 (0, 818) 200 (0, 607) 0 (0, 29) 200 (0, 618) 1000 - 222019 (10.9%) 5 (2.9%) 24 (13.8%) 500 - < 1000 29 (16.7%) 16 (9.2%) 2 (1.2%) 47 (27.0%)

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12 (6.9%)

5 (2.9%)

33 (19.0%)

16 (9.2%)

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	Plasma 25(OH)D Levels			Total (n) or
	Sufficient (n) [30 ng/mL]	Insufficient (n) [30 > n 20 ng/mL]	Deficient (n) [< 20 ng/mL]	Mean \pm SD or Median (25% ^{ile} , 75% ^{ile})
None	29 (16.7%)	23 (13.2%)	18 (10.3%)	70 (40.2%)
Beer or liquor use (drinks/month)***	14 (4, 36)	12 (4, 28)	18 (6, 56)	12 (4, 36)
Above median (> 12)	27 (15.5%)	8 (4.6%)	6 (3.5%)	41 (23.6%)
Below median (12)	27 (15.5%)	10 (5.8%)	6 (3.5%)	43 (24.7%)
None	39 (22.4%)	38 (21.8%)	13 (7.5%)	90 (51.7%)
Wine use (drinks/month)***	12 (4, 24)	12 (8, 24)	8 (2, 12)	12 (4, 24)
Above median (> 12)	18 (10.3%)	6 (3.5%)	-	24 (13.8%)
Below median (12)	20 (11.5%)	7 (4.0%)	3 (1.7%)	30 (17.2%)
None	55 (31.6%)	43 (24.7%)	22 (12.6%)	120 (69.0%)
		Lifestyle		
Smoking status				
Current	5 (2.9%)	7 (4.0%)	4 (2.3%)	16 (9.2%)
Former	45 (25.9%)	26 (14.9%)	15 (8.6%)	86 (49.4%)
Never	43 (24.7%)	23 (13.2%)	6 (3.5%)	72 (41.4%)
Planned exercise (hrs/week)	$7.2 \pm 6.1 (n=59)$	$6.0 \pm 5.3 \ (n=36)$	5.0 ± 3.5 $(n=12)$	$6.5 \pm 5.6 \; (n=107)$
Stretching/Range of Motion (ROM)/Physical Therapy (hrs/week)	$2.7 \pm 2.7 (n=24)$	$3.5 \pm 3.7 (n=17)$	$2.3 \pm 1.4 (n=6)$	$3.0 \pm 3.0 \; (n=47)$
Other Exercise (hrs/week)	6.8 ± 6.1 (n=53)	$5.0 \pm 4.4 (n=31)$	5.1 ± 4.0 (n=9)	$6.0 \pm 5.4 (n=93)$
Time outside between sunrise and sunset (hrs/week)	12.2 ± 12.1 (n=87)	14.6 ± 14.6 (n=52)	11.9 ± 8.0 $(n=24)$	12.9 ± 12.4 (n=163)
Time outside between sunrise and sunset, visits April through September (hrs/week)	14.5 ± 13.2 (n=52)	15.9 ± 14.6 (n=29)	11.6 ± 8.9 (n=16)	14.4 ± 13.0 (n=97)
Sunscreen use when outdoors				
Usually (50%)	15 (9.0%)	3 (1.8%)	-	18 (10.8%)
Sometimes	13 (7.8%)	12 (7.2%)	1 (0.6%)	26 (15.6%)
Never	62 (37.1%)	40 (24.0%)	21 (12.6%)	123 (73.7%)

^{*} Includes 19 Black, 4 Hispanic, 2 Asian, 1 Native American

^{**}Obstructive lung disease

^{***} Excludes non-drinkers

Table 2

Univariable associations with plasma vitamin D

	$\beta \pm SE$	P
Personal		
Age (Years)	0.115 ± 0.06	0.068
BMI (kg/m²)	-0.158 ± 0.13	0.225
Total body fat (%)	-0.031 ± 0.08	0.715
Education (years)	0.490 ± 0.25	0.052
Duration of injury (years)	0.010 ± 0.06	0.876
Daily Intake		
Total vitamin D intake (per 100 IU)	0.451 ± 0.07	< 0.001
Vitamin D intake from diet <i>(per 100 IU)</i>	0.695 ± 0.61	0.259
Vitamin D from supplements (per 100 IU)	0.444 ± 0.08	< 0.001
Total calcium intake <i>(per 100 mg)</i>	0.417 ± 0.11	< 0.001
Calcium intake from diet <i>(per 100 mg)</i>	0.315 ± 0.18	0.082
Calcium from supplements (per 100 mg)	0.501 ± 0.15	0.001
Lifestyle		
Planned exercise (hrs/week)	0.317 ± 0.14	0.028
Stretching/Range of Motion /Physical Therapy (hrs/week)	-0.232 ± 0.39	0.554
Other Exercise (hrs/week)	0.418 ± 0.16	0.008
Time outside, sunrise and sunset (hrs/week)	-0.011 ± 0.06	0.857
Time outside, sunrise and sunset, April – September (hr/week)	0.062 ± 0.08	0.436
	Mean $(ng/mL) \pm SE$	P
Personal		
Sex		0.021
Male	30.4 ± 0.9	
Female	35.0 ± 1.8	
Race		< 0.001
White	32.8 ± 0.8	
Other	22.3 ± 1.9	
BMI		0.262

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	β±SE	P
Obese (30)	29.8 ± 1.5	
Overweight (25 – < 30)	30.6 ± 1.3	
Normal/Underweight (< 25)	32.8 ± 1.3	
Season of Visit		0.737
Winter	32.5 ± 1.7	
Spring	30.1 ± 1.7	
Summer	30.6 ± 1.4	
Autumn	31.7 ± 1.5	
SCI level and completeness		0.319
Motor complete cervical & AIS C	29.8 ± 1.6	
Other motor complete & AIS C	32.7 ± 1.3	
All AIS D	30.6 ± 1.2	
Mobility mode		0.476
Motorized wheelchair	31.1 ± 1.7	
Hand-propelled wheelchair	32.4 ± 1.2	
Walk with aid	29.2 ± 2.2	
Walk without assistance	29.9 ± 1.8	
Comorbid Diseases		
Asthma	32.6 ± 2.3	0.507
No Asthma	31.0 ± 0.8	
COPD	32.3 ± 2.3	0.610
No COPD	31.1 ± 0.8	
OLD *(asthma or COPD)	32.8 ± 1.7	0.295
No OLD	30.8 ± 0.9	
Diabetes	30.7 ± 2.0	0.790
No diabetes	31.3 ± 0.9	
Hypertension	32.4 ± 1.3	0.261
No hypertension	30.6 ± 1.0	
Heart disease, last 10 years	34.4 ± 2.2	0.127
No heart disease	30.8 ± 0.8	
Intai	ke	
Beer or liquor use (drinks/month)		0.411
Above median (> 12)	31.0 ± 1.6	
Below median (12)	33.0 ± 1.6	
None	30.5 ± 1.1	
Wine use (drinks/month)		< 0.001
		P trend < 0.00
Above median (> 12)	37.0 ± 2.0	

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	$\beta \pm SE$	P
Below median (12)	34.0 ± 1.8	
None	29.4 ± 0.9	
L	ifestyle	
Smoking status		0.057
		P trend 0.020
Current	26.4 ± 2.6	
Former	30.7 ± 1.1	
Never	33.0 ± 1.2	
Sunscreen use when outdoors		0.003
Usually (50%) **	38.5 ± 2.3	
Sometimes	32.6 ± 1.9	
Never	30.2 ± 0.9	

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^{*} Obstructive lung disease

 $^{^{**}}$ P=0.127 compared to sometimes and P=0.004 compared to never

Table 3

Final multivariable model

Continuous Variables	β (95% CI)	P
Age (years)	0.142 (0.032, 0.251)	0.012
Total daily vitamin D intake (per 100 IU)	0.355 (0.219, 0.490)	< 0.001
Planned exercise (hours per week)	0.267 (0.029, 0.505)	0.028
Categorical Variables	$Mean (ng/mL) \pm SE$	
Sex		0.011
Male	28.6 ± 1.3	
Female	32.9 ± 1.8	
Race		< 0.001
White	34.3 ± 1.1	
Other	27.2 ± 2.0	
Wine use (drinks per month)		0.030
		P trend 0.015
Above median (> 12)	33.0 ± 2.1	
Below median (12)	30.9 ± 1.9	
None	28.3 ± 1.2	
Smoking status		0.010
		P trend 0.003
Current	28.5 ± 2.4	
Former	29.9 ± 1.3	
Never	33.8 ± 1.5	
Vitamin D by Diet and Supplement *	β (95% CI)	P
Vitamin D intake from diet (per 100 IU)	0.471 (-0.528, 1.470)	0.353
Vitamin D intake from supplements (per 100 IU)	0.353 (0.216, 0.489)	< 0.001
Planned Exercise**	β (95% (CI) P

^{*}Adjusting for age, planned exercise, sex, race, wine use, and smoking status

Other Exercise (hrs/week)

0.297 (0.034, 0.559)

0.027

^{**} Adjusting for age, total daily VitD intake, sex, race, wine use, and smoking status