

Vitamin D Concentrations and Obstructive Sleep Apnea in a Multicenter Cohort of Older Males

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Abstract

Rationale: Seasonal nadirs in 25-hydroxyvitamin D (25[OH]D) concentrations overlap with increased incidence and severity of obstructive sleep apnea (OSA) in winter. We hypothesized that, because lower 25(OH)D concentrations might lead to upper airway muscle dysfunction, low 25(OH)D would be associated with higher apnea-hypopnea index (AHI), a measure of OSA severity.

Objectives: To determine if lower 25(OH)D concentration is associated with greater prevalence and increased severity of OSA, independent of established OSA risk factors.

Methods: Using unconditional logistic regression, we performed a cross-sectional analysis in the Outcomes of Sleep Disorders in Older Men study, which included in-home overnight polysomnography, serum 25(OH)D measurement, and collection of demographic and comorbidity data. The primary outcome was severe sleep apnea, as defined by AHI of 30/h or more.

Measurements and Main Results: Among 2,827 community-dwelling, largely white (92.2%), elderly (aged 76.4 ± 5.5 yr [mean ± SD]) males, mean 25(OH)D concentration was 28.8 (± 8.8) ng/ml. Subjects within the lowest quartile of 25(OH)D (6–23 ng/ml) had greater odds of severe sleep apnea in unadjusted analyses (odds ratio = 1.45; 95% confidence interval = 1.02–2.07) when compared with the highest 25(OH)D quartile (35–84 ng/ml). However, further adjustment for established OSA risk factors strongly attenuated this association (multivariable adjusted odds ratio = 1.05; 95% confidence interval = 0.72–1.52), with body mass index and neck circumference as the main confounders. There was also no evidence of an independent association between lower 25(OH)D levels and increased odds of mild (AHI = 5.0–14.9/h) or moderate (AHI = 15.0–29.9/h) sleep apnea.

Conclusions: Among community-dwelling older men, the association between lower 25(OH)D and sleep apnea was largely explained by confounding by larger body mass index and neck circumference.

Keywords: obstructive sleep apnea; vitamin D; obesity; cross-sectional study

(Received in original form July 16, 2015; accepted in final form December 12, 2015)

Supported by National Institutes of Health (NIH) funding (to the Osteoporotic Fractures in Men Study [MrOS]), and by the National Institute of Arthritis and Musculoskeletal and Skin Diseases, the National Institute on Aging, the National Center for Research Resources, and NIH Roadmap for Medical Research grants U01 AR45580, U01 AR45614, U01 AR45632, U01 AR45647, U01 AR45654, U01 AR45583, U01 AG18197, U01 AG027810, and UL1 TR000128; the National Heart, Lung, and Blood Institute provides funding for the MrOS Sleep ancillary study "Outcomes of Sleep Disorders in Older Men" under grants R01 HL071194, R01 HL070848, R01 HL070847, R01 HL070842, R01 HL070841, R01 HL070837, R01 HL070838, and R01 HL070839, and funding for the Vitamin D assays was provided under grant R01 AG030089; the Veterans Health Administration Office of Research and Development also provided protected research time in support of this study.

Some of these data were previously presented as a poster abstract at the American Thoracic Society International Conference, San Diego, California, May 2014.

The views expressed in this article are those of the authors and do not reflect the views of the U.S. Government, the Department of Veterans Affairs, or any of the authors' affiliated institutions.

Author Contributions: U.G. contributed to the study design, data collection, data analysis, manuscript writing, and final editing, had full access to all the data in the study, and takes responsibility for the integrity and accuracy of the data analysis; K.E.E. contributed to the study design, data analysis, manuscript writing, and final editing; M.L.P. contributed to the study design, data collection, data analysis, manuscript writing, and final editing; S.R. contributed to the study design, manuscript writing, and final editing; E.S.S. contributed to the study design, manuscript writing, and final editing; J.M.S. contributed to the study design, manuscript writing, and final editing; K.L.S. contributed to the study design, manuscript writing, and final editing; K.M.K. contributed to the study design, data collection, data analysis, manuscript writing, and final editing, had full access to all the data in the study, and takes responsibility for the integrity and accuracy of the data analysis.

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Ann Am Thorac Soc Vol 13, No 5, pp 712–718, May 2016

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DOI: 10.1513/AnnalsATS.201507-440OC

Internet address: www.atsjournals.org

Obstructive sleep apnea (OSA) is a disease of recurrent, partial, or complete upper airway closure during sleep. The pathogenesis of OSA is complex (1), and a recent study reported increased OSA incidence and severity during winter months (2). Although the magnitude of the effect was modest (median difference in apnea-hypopnea index [AHI] of 3.3 events/h), the authors hypothesized that winter-related fat redistribution, medication use, fluid displacement to the neck, and/or air pollution were the reasons for their observation. However, an alternative explanation of the seasonal variation in OSA may be the coinciding winter nadir of vitamin D (3).

Low circulating 25-hydroxyvitamin D (25[OH]D) concentrations are associated with poor musculoskeletal function (4, 5). Control of upper airway muscle tone is thought to be a major contributor to OSA (1), and, therefore, patients with low 25(OH)D concentrations might have an increased risk of OSA due to worse function of the skeletal muscle supporting upper airway patency during sleep. Low 25(OH)D concentrations are also associated with airway inflammation, chronic rhinitis, and repeated upper airway infections, leading to tonsillar enlargement (6–10), which may additionally contribute to OSA incidence and severity. Low 25(OH)D concentrations are also associated with type 2 diabetes mellitus, metabolic syndrome, and obesity, all of which are frequently found in patients with OSA (11, 12). Only a few studies have studied the association between lower 25(OH)D levels and OSA, and these have reported inconsistent results (13–15).

To better address this knowledge gap, we analyzed data from a large, multicenter, community-based cohort study and tested the hypothesis that lower blood concentrations of 25(OH)D are associated with greater prevalence and increased severity of OSA, independent of classic OSA risk factors.

Methods

Study Participants

The Osteoporotic Fractures in Men Study (MrOS) enrolled 5,995 community-dwelling men aged 65 years and older during the baseline examination between 2000 and 2002 (16, 17). To be eligible for the study,

men had to be able to walk without assistance and not have had a bilateral hip replacement. Participants were recruited at six clinical centers (Birmingham, AL; Minneapolis, MN; Palo Alto, CA; Monongahela Valley near Pittsburgh, PA; Portland, OR; and San Diego, CA). The Outcomes of Sleep Disorders in Older Men Study (MrOS Sleep) visit occurred on average 3.4 (± 0.5) years (range, 1.9–4.9 yr) after the baseline examination, between December 2003 and March 2005. Ethics approval was obtained from the institutional review board at each site, and the Coordinating Center and Reading Center. Written informed consent for participation in the MrOS Sleep Study was obtained for all individuals.

The MrOS Sleep Study was an ancillary study with a target recruitment number of 3,000 men from the parent MrOS Study. Exclusions for the MrOS Sleep Study included use of nocturnal positive airway pressure or oral appliance devices, use of supplemental oxygen, and presence of an open tracheostomy. Of the 5,995 MrOS participants, 3,135 participated in the MrOS Sleep Study, whereas 2,860 participants in the main cohort did not participate in the MrOS Sleep Study. Of the nonparticipants in the MrOS Sleep Study, 1,997 refused participation, and, compared with those who participated in the MrOS Sleep Study, those who refused participation were older by 1 year (74.01 ± 5.88 yr vs. 73.05 ± 5.55 yr; $P < 0.001$) and not significantly different with respect to body mass index (BMI; 27.20 ± 3.72 kg/m² vs. 27.38 ± 3.72 kg/m²; $P = 0.10$) (18).

Of 3,135 MrOS Sleep Study participants, 2,911 (92.8%) had at least 4 hours of technically adequate sleep study data for analysis, and, of these, 2,827 (90.2%) had 25(OH)D concentrations measured. These 2,827 men constituted the analytical cohort for the current study.

Polysomnography and Other Sleep-Related Measures

In-home sleep studies using unattended polysomnography (Safiro, Compumedics, Inc., Melbourne, VIC, Australia) were performed. The recording montage consisted of C3/A2 and C4/A1 electroencephalograms, bilateral electrooculograms, a bipolar submental electromyogram, thoracic and abdominal respiratory inductance plethysmography, airflow (using nasal–oral thermocouple and

nasal pressure cannula), finger pulse oximetry, electrocardiogram, body position (mercury switch sensor), and bilateral leg movements (piezoelectric sensors).

Trained, certified staff members performed home visits for setup of the sleep study units. After sensors were placed and calibrated, signal quality and impedance were checked, and sensors were repositioned as needed to improve signal quality, replacing electrodes if impedances were greater than 5 k Ω , using approaches similar to those in the Sleep Heart Health Study (19). After studies were downloaded, they were transferred to the Case Reading Center (Cleveland, OH) for centralized scoring by a trained technician. Polysomnography data quality was excellent, with a failure rate of less than 4%, and more than 70% of studies graded as being of excellent or outstanding quality. Quality codes for signals and studies were graded using previously described approaches, which included coding the duration of artifact-free data per channel and overall study quality (reflecting the combination of grades for each channel). We note that central apnea events were included in the overall AHI, but such events were rare (only 7% of study participants had a central apnea index ≥ 5 /h).

Vitamin D Analysis

Serum for vitamin D analysis was collected at baseline of the MrOS Sleep Study and immediately frozen at -70°C . Concentrations of 25(OH)D₂ and 25(OH)D₃ were measured by liquid chromatography–tandem mass spectrometry (ThermoFisher Scientific, Franklin, MA and Applied Biosystems–MDS Sciex, Foster City, CA) at the Mayo Clinic Reference Laboratories (R. J. Singh, Ph.D.; Mayo Clinic Laboratory, Rochester, MN), using fasting samples collected at the sleep visit. Deuterated stable isotope (d_3 -25[OH]D) was added to a 0.2-ml serum sample as internal standard. 25(OH)D₂, 25(OH)D₃, and the internal standard were extracted using acetonitrile precipitation.

The extracts were then further purified online and analyzed by liquid chromatography–tandem mass spectrometry using multiple-reaction monitoring. Using three different target markers as quality controls for each assay, interassay coefficients of variation (CVs) for 25(OH)D₃ were 9.7% at 9.0 IU, 7.5% at

29 IUs, and 5.8% at 76 IUs. For 25(OH)D₂, CVs were 11.2% at 11 IUs, 8.5% at 28 IUs, and 7.7% at 74 IUs. The minimum detectable limit for 25(OH)D₂ was 4 ng/ml and for 25(OH)D₃ was 2 ng/ml. Total 25(OH)D was calculated by adding 25(OH)D₂ and 25(OH)D₃.

Clinical Data

All covariate data were collected at the time of the sleep study visit. All participants completed questionnaire data, which included questions about medical history, smoking, and alcohol intake. Hypertension was defined as a positive response to the question, "Has a doctor or other healthcare provider told you that you have hypertension or high blood pressure?" Race was based on self-report and categorized as white, African American, Asian, or Hispanic/other. BMI was calculated as weight (kg)/height (m²), and obesity was defined as a BMI greater than 30 kg/m². During the home or clinic visits, body weight was measured using a standard balance beam scale and height using a wall-mounted Harpenden stadiometer (Holtain, UK). Neck and waist circumference were also measured using standard methods. Snoring was assessed according to self-report.

The MacArthur Subjective Status Scale (range, 1–10) was used to assess perceived social status, with higher scores representing higher perceived social status. Participants were asked to bring in all current medications used within the preceding 30 days. All prescription and nonprescription medications were entered into an electronic database, and each medication was matched to its ingredient(s) based on the Iowa Drug Information Service Drug Vocabulary (College of Pharmacy, University of Iowa, Iowa City, IA). A variable for season (January–March = winter; April–June = spring; July–September = summer; October–December = fall) was calculated using the date of the participant's clinic exam.

Statistical Analysis

In primary analyses, we expressed 25(OH)D concentrations in quartiles and compared baseline characteristics across quartiles using ANOVA or chi-square testing for continuous or categorical variables, respectively. We also used the following fixed 25(OH)D categories in our analyses: less than 20 ng/ml; 20–29.9 ng/ml; and 30 ng/ml or greater. However, because results were similar, we present quartiles as our primary 25(OH)D variable.

We used unconditional logistic regression models to calculate odds ratios (ORs) and 95% confidence intervals (CIs) for 25(OH)D quartiles (referent group quartile 4) for the primary dichotomous outcome of severe sleep apnea as defined by AHI of 30/h or greater and a secondary dichotomous outcome of at least moderate sleep apnea defined by AHI of 15/h or greater. We created four logistic regression models, including an unadjusted model, a model adjusted for established OSA risk factors (age, BMI, neck circumference, and hypertension) (18, 20–22), a model further adjusted for season of blood draw (23), and a fully adjusted model that included the above variables along with medications that may affect upper airway patency (opiates, benzodiazepines and alcohol) (20, 24, 25), race, smoking, and clinic site. We conducted a sensitivity analysis to evaluate the effect of selected variables (BMI, neck circumference, hypertension) on the outcome by adding one variable at a time to the crude model. We also conducted two secondary analyses expressing 25(OH)D using clinical cutpoints (<20 ng/ml, 20–29.9 ng/ml, and ≥30 ng/ml) (26), and a polytomous regression model comparing outcomes of AHI of 0–4.9/h (no OSA) to standard OSA severity categories of mild

Table 1. Characteristics of 2,827 participants by 25-hydroxyvitamin D quartiles

Variable	Overall Cohort (n = 2,827)	Serum 25(OH) D Quartiles				P Value
		Q1 6–23 ng/ml (n = 767)	Q2 24–28 ng/ml (n = 686)	Q3 29–34 ng/ml (n = 678)	Q4 35–84 ng/ml (n = 696)	
Age, yr	76.4 ± 5.5	76.8 ± 5.6	76.4 ± 5.5	76.1 ± 5.4	76.3 ± 5.6	0.10
White, %	92.2	88.7	92.4	93.7	94.4	0.0002
BMI, kg/m ²	27.2 ± 3.8	28.1 ± 4.2	27.5 ± 4.0	26.7 ± 3.4	26.3 ± 3.1	<0.0001
Neck circumference, cm	39.4 ± 2.8	40.0 ± 2.9	39.6 ± 2.9	39.2 ± 2.6	38.9 ± 2.6	<0.0001
Systolic blood pressure, mm Hg	126.8 ± 16.2	126.8 ± 16.9	126.7 ± 16.1	126.8 ± 16.3	126.9 ± 15.7	0.99
Current smoker, %	2.0	2.4	2.2	1.2	2.2	0.68
Alcohol intake, >14 drinks/wk, %	5.4	3.9	6.0	3.6	8.2	<0.0001
Benzodiazepine use, %	4.63	4.82	4.52	4.28	4.89	0.95
Opiate use, %	3.30	5.22	3.64	3.10	3.30	0.137
Winter (January–March), %	33.4	42.6	32.1	31.0	26.7	<0.0001
Spring (April–June), %	25.3	26.6	27.1	26.3	21.0	<0.0001
Summer (July–September), %	22.6	15.8	24.1	23.0	28.2	<0.0001
Fall (October–December), %	18.8	15.0	16.8	19.8	24.1	<0.0001
AHI 0–4.9/h, %	39.1	37.2	39.7	39.1	40.5	0.26
AHI 5–14.9/h, %	34.6	34.9	31.3	36.3	35.6	0.26
AHI 15–29.9/h, %	16.6	16.8	17.8	15.9	16.0	0.26
AHI ≥30/h, %	9.8	11.1	11.2	8.7	7.9	0.26
AHI, events/h	11.8 ± 12.9	12.4 ± 13.3	12.4 ± 13.2	11.4 ± 12.7	10.9 ± 12.5	0.08
25(OH)D, ng/ml	28.8 ± 8.8	18.5 ± 4.0	26.0 ± 1.4	31.4 ± 1.7	40.37 ± 5.5	<0.0001

Definition of abbreviations: 25(OH)D = 25-hydroxyvitamin D; AHI = apnea-hypopnea index; BMI = body mass index; Q = quartile.

(AHI, 5–14.9/h), moderate (AHI, 15–29.9/h) and severe (AHI, ≥ 30 /h) OSA.

In exploratory analyses, to examine effect modification by BMI, we evaluated the association between 25(OH)D and OSA excluding men with BMI of 30 kg/m² or greater, hypothesizing that the relationship between 25(OH)D and OSA would be stronger in nonobese men. We also tested for an interaction between 25(OH)D and BMI for the prediction of severe OSA, using the log-likelihood ratio test.

Results

The study cohort consisted of largely white (92.2%) older men (aged 76.4 \pm 5.5 yr [mean \pm SD]), with a mean BMI of 27.2 (\pm 3.8) kg/m². The demographic and comorbid disease characteristics of the study participants are provided in Table 1.

The mean 25(OH)D concentration was 28.8 (\pm 8.8) ng/ml, with concentrations 30 ng/ml or greater (widely considered to represent replete vitamin D status) in 1,247 (44.1%), between 20 and 29.9 ng/ml (widely

considered vitamin D insufficient) in 1,205 (42.6%), and less than 20 ng/ml (widely considered vitamin D deficient) in 375 (13.3%). The distribution of AHI categories 0–4.9, 5–14.9, 15–29.9, and 30 or greater events/h was 1,105 (39%), 977 (34.6%), 470 (16.6%), and 276 (9.8%), respectively. Among the 276 participants with AHI of 30/h or greater, 50 (18.1%) were 25(OH)D deficient (<20 ng/ml), compared with 134 (12.1%) who were 25(OH)D deficient among those with AHI less than 5/h ($P = 0.06$).

Participants in the lowest quartile of 25(OH) D (6–23 ng/ml) had greater odds of AHI of 30/h or greater (crude OR = 1.45; 95% CI = 1.02–2.07) compared with participants in the highest 25(OH)D quartile (35–84 ng/ml) (Table 2). After adjustment for traditional OSA risk factors, this association was no longer evident (adjusted OR = 1.05; 95% CI = 0.72–1.52). Further adjustment for season and other covariates did not alter these results. Findings were similar when AHI of 15/h or greater was substituted for AHI 30/h or greater, when 25(OH)D was expressed using

clinical cutpoints of less than 20 ng/ml, 20–29.9 ng/ml, and 30 ng/ml or greater, and in polytomous regression evaluating the association between 25(OH)D quartiles and odds of no OSA versus mild/moderate/severe OSA (Table 3).

Sensitivity analysis suggested that the association between lower 25(OH)D concentrations and higher odds of OSA was largely explained by greater BMI and larger neck circumference among those men with lower 25(OH)D concentrations (Table 4).

In exploratory analysis restricted to those with BMI less than 30 kg/m² ($n = 2,255$; 79.8% of overall cohort), the modest association between low 25(OH)D and higher odds of severe sleep apnea did not reach significance (fully adjusted OR [quartile 1 vs. quartile 4] of AHI ≥ 30 /h = 1.27; 95% CI = 0.82–1.97; fully adjusted OR [25(OH)D <20 ng/ml vs. ≥ 30 ng/ml] of AHI ≥ 30 /h = 1.44; 95% CI = 0.90–2.28). There was no evidence of an interaction between obesity (BMI <30 vs. ≥ 30 kg/m²) and 25(OH)D for the prediction of AHI of 30/h or greater, when 25(OH)D was categorized by quartiles ($P = 0.197$) or by clinical cutpoints ($P = 0.562$).

Table 2. Logistic regression for odds of obstructive sleep apnea

25(OH)D Quartiles	OR (95% CI) of AHI ≥ 15 /h	OR (95% CI) of AHI ≥ 30 /h
Crude		
Q1 vs. Q4	1.24 (0.98–1.56)	1.45 (1.02–2.07)
Q2 vs. Q4	1.31 (1.03–1.66)	1.47 (1.02–2.12)
Q3 vs. Q4	1.04 (0.82–1.34)	1.11 (0.76–1.63)
<i>P</i> value for trend	0.027	0.016
Adjusted for typical OSA risk factors*		
Q1 vs. Q4	0.96 (0.75–1.23)	1.05 (0.72–1.52)
Q2 vs. Q4	1.11 (0.87–1.43)	1.18 (0.81–1.72)
Q3 vs. Q4	0.98 (0.76–1.27)	1.06 (0.72–1.57)
<i>P</i> value for trend	0.940	0.737
Adjusted for typical OSA risk factors* and 25(OH)D modifiers[†]		
Q1 vs. Q4	0.94 (0.72–1.22)	0.97 (0.65–1.44)
Q2 vs. Q4	1.13 (0.87–1.46)	1.15 (0.78–1.70)
Q3 vs. Q4	1.00 (0.77–1.29)	1.05 (0.71–1.56)
<i>P</i> value for trend	0.827	0.931
Fully adjusted[‡]		
Q1 vs. Q4	0.94 (0.72–1.23)	0.96 (0.65–1.43)
Q2 vs. Q4	1.13 (0.87–1.47)	1.15 (0.78–1.70)
Q3 vs. Q4	0.98 (0.75–1.27)	0.99 (0.66–1.48)
<i>P</i> value for trend	0.884	0.993

Definition of abbreviations: 25(OH)D = 25-hydroxyvitamin D; AHI = apnea–hypopnea index; CI = confidence interval; OR = odds ratio; OSA = obstructive sleep apnea; Q = quartile.

*“Typical OSA risk factors” covariates: age, body mass index, neck circumference, hypertension.

[†]Winter season: January–March.

[‡]Fully adjusted covariates: age, body mass index, neck circumference, hypertension, winter season, clinic site, race, alcohol consumption, smoking, benzodiazepine use, opioid use.

Discussion

Despite plausible mechanisms potentially connecting vitamin D deficiency to OSA pathogenesis and severity, we found no evidence of an independent association between 25(OH)D concentration and OSA in our analyses of this cohort of older community-dwelling men.

Our findings suggest that the association between lower 25(OH)D levels and a higher odds of OSA was due, in large part, to greater BMI and larger neck circumference among participants with lower 25(OH)D levels. These results indicate that low 25(OH)D may simply be a marker of larger BMI and neck circumference, rather than directly contributing to OSA pathogenesis.

Several lines of evidence link obesity with lower 25(OH)D concentrations. Obese subjects are more likely to be restricted in physical activity, thus limiting their exposure to sunlight and resulting in lower 25(OH)D concentrations (27). In addition, inflammatory cytokines up-regulated in adiposity are known to inversely affect 25(OH)D bioavailability and increase its

Table 3. Polytomous logistic regression for odds of obstructive sleep apnea

25(OH)D Quartiles	OR (95% CI)		
	AHI 5–14.9/h vs. AHI 0–4.9/h	AHI 15–29.9/h vs. AHI 0–4.9/h	AHI ≥30/h vs. AHI 0–4.9/h
Crude			
Q1 vs. Q4	1.07 (0.84–1.36)	1.15 (0.85–1.56)	1.53 (1.05–2.23)
Q2 vs. Q4	0.90 (0.70–1.15)	1.14 (0.84–1.55)	1.45 (0.99–2.13)
Q3 vs. Q4	1.06 (0.83–1.35)	1.04 (0.76–1.42)	1.14 (0.76–1.71)
P value for trend		0.073	
Adjusted for typical OSA risk factors*			
Q1 vs. Q4	0.89 (0.69–1.14)	0.87 (0.64–1.19)	0.97 (0.65–1.45)
Q2 vs. Q4	0.80 (0.62–1.03)	0.96 (0.70–1.32)	1.07 (0.72–1.60)
Q3 vs. Q4	1.00 (0.78–1.28)	0.95 (0.69–1.31)	1.05 (0.69–1.59)
P value for trend		0.534	
Adjusted for typical OSA risk factors* and 25(OH)D modifiers†			
Q1 vs. Q4	0.91 (0.70–1.18)	0.88 (0.63–1.24)	0.91 (0.59–1.39)
Q2 vs. Q4	0.82 (0.63–1.06)	0.99 (0.72–1.38)	1.06 (0.70–1.61)
Q3 vs. Q4	1.01 (0.79–1.30)	0.98 (0.70–1.36)	1.05 (0.69–1.60)
P value for trend		0.720	
Fully adjusted‡			
Q1 vs. Q4	0.91 (0.70–1.19)	0.89 (0.63–1.25)	0.90 (0.59–1.38)
Q2 vs. Q4	0.81 (0.62–1.06)	0.99 (0.71–1.38)	1.06 (0.70–1.61)
Q3 vs. Q4	0.99 (0.7–1.28)	0.98 (0.70–1.36)	0.98 (0.64–1.50)
P value for trend		0.739	

Definition of abbreviations: 25(OH)D = 25-hydroxyvitamin D; AHI = apnea–hypopnea index; CI = confidence interval; OR = odds ratio; OSA = obstructive sleep apnea; Q = quartile.

*“Typical OSA risk factors” covariates: age, body mass index, neck circumference, hypertension.

†Winter season: January–March.

‡Fully adjusted covariates: age, body mass index, neck circumference, hypertension, winter season, clinic site, race, alcohol consumption, smoking, benzodiazepine use, opioid use.

metabolic clearance (28). Finally, poor dietary habits leading to obesity often provide a poor source of oral vitamin D intake.

We explored the possibility that 25(OH)D effects on OSA pathogenesis might be most pronounced in nonobese participants, because mechanisms other than soft tissue accumulation and external airway pressure might account for loss of upper airway patency during sleep in nonobese individuals. In these analyses, we observed similar nonsignificant findings, although the adjusted OR point estimates for low 25(OH)D and AHI of 30/h or greater were higher than in analyses that included both obese and nonobese individuals. There was no evidence of an interaction between obesity and 25(OH)D on OSA, but further studies with larger sample sizes are needed to more definitively evaluate the potential role of vitamin D deficiency in the pathogenesis of OSA in nonobese individuals.

Despite reasons to believe that vitamin D deficiency may play a role in the pathogenesis of OSA, few studies to date

have explored this potential association, and these have reported mixed results. In a selected population of 150 adult patients with OSA (50 patients each with mild, moderate, and severe OSA) and control subjects matched on BMI, sex, and age ($n = 32$), Mete and colleagues (13) noted no significant difference in 25(OH)D concentrations between patients with OSA and control subjects ($17.9 \pm 9.3 \mu\text{g/dl}$ vs. $19.2 \pm 7.2 \mu\text{g/dl}$; $P = 0.468$). However, the authors reported that, among those with AHI greater than 30/h, 78% were 25(OH)D deficient ($<20 \mu\text{g/dl}$) compared with 50% among control subjects with AHI less than 5/h ($P = 0.02$). We found similar results in our population of older men, although our observed difference in prevalence of 25(OH)D deficiency was smaller in magnitude (18.1% among those with AHI ≥ 30 /h, and 12.1% among those with AHI < 5 /h).

However, the study by Mete and colleagues notably preselected obese patients in their study (mean BMI of 32 kg/m^2), and they were therefore not able to adjust for the effect of BMI on the

relationship between 25(OH)D and AHI. We have addressed this in our study by enrolling an unselected population of community-dwelling older men with a wide range of BMI (and AHI) and adjusting our analysis for BMI and several other potential confounders.

In contrast to the results of Mete and colleagues and our study, Kheirandish-Gozal and colleagues (14) found a statistically significant correlation between 25(OH)D concentrations and AHI ($r = -0.285$; $P < 0.001$) in pediatric patients with OSA (mean age, 6.5–7.2 yr). The relevance of these results to adult OSA is not clear, because pediatric OSA has a very different pathogenesis than adult OSA. In children, upper airway inflammation, in the form of adenotonsillar hypertrophy, is believed to be the major contributor to OSA (29) as opposed to that in adults, where fat redistribution, upper airway muscle dysfunction, and age-related changes in upper airway anatomy are believed to be the major contributors (1).

More recently, Bertisch and colleagues (15) studied the association between

Table 4. Sensitivity analysis: logistic regression with adjustment for only one typical obstructive sleep apnea risk factor (age, body mass index, neck circumference, hypertension)

25(OH)D Quartiles	OR (95% CI)	
	AHI $\geq 15/h$ vs. $<15/h$	AHI $\geq 30/h$ vs. $<30/h$
Adjusted for continuous age		
Q1 vs. Q4	1.22 (0.97–1.55)	1.43 (1.00–2.04)
Q2 vs. Q4	1.30 (1.02–1.65)	1.47 (1.02–2.11)
Q3 vs. Q4	1.05 (0.82–1.34)	1.12 (0.77–1.65)
P value for trend	0.035	0.024
Adjusted for continuous BMI		
Q1 vs. Q4	0.99 (0.78–1.27)	1.12 (0.77–1.61)
Q2 vs. Q4	1.14 (0.89–1.45)	1.23 (0.85–1.78)
Q3 vs. Q4	0.99 (0.77–1.27)	1.04 (0.70–1.53)
P value for trend	0.806	0.429
Adjusted for neck circumference		
Q1 vs. Q4	1.07 (0.84–1.36)	1.22 (0.85–1.75)
Q2 vs. Q4	1.18 (0.92–1.50)	1.29 (0.89–1.87)
Q3 vs. Q4	0.99 (0.77–1.28)	1.07 (0.73–1.57)
P value for trend	0.354	0.201
Adjusted for hypertension		
Q1 vs. Q4	1.23 (0.97–1.56)	1.45 (1.01–2.06)
Q2 vs. Q4	1.32 (1.04–1.68)	1.49 (1.03–2.14)
Q3 vs. Q4	1.04 (0.81–1.33)	1.11 (0.75–1.63)
P value for trend	0.028	0.016

Definition of abbreviations: 25(OH)D = 25-hydroxyvitamin D; AHI = apnea–hypopnea index; BMI = body mass index; CI = confidence interval; OR = odds ratio; OSA = obstructive sleep apnea; Q = quartile.

25(OH)D concentrations and various sleep measures, including AHI in a multicenter, multiethnic cohort of 1,721 adults with a mean age of 68.2 years. The authors reported that those with 25(OH)D under 20 ng/ml had a statistically higher median AHI (2.1 events/h higher) than those with 25(OH)D over 29 ng/ml. Similar to our findings, those with lower 25(OH)D concentrations were also more obese. Also similar to our findings, the AHI difference was no longer significant after adjusting for covariates, such as age, sex, and waist circumference.

In exploratory analyses, the authors found that AHI was higher by 7.1 events/h in Chinese American participants, ($n = 205$) with low 25(OH)D compared with higher 25(OH)D, but this was not found in the white, African American, or Hispanic American participants. The authors importantly noted that 25(OH)D concentrations were measured an average of 10.3 years before the collection of polysomnography data. Whether or not 10-year-old 25(OH)D measures reflect 25(OH)D status at the time of polysomnography is a major potential limitation. Our 25(OH)D and

polysomnography data were concurrent and therefore truly cross-sectional. Nevertheless, the data from their cohort and our cohort would appear to be consistent with an overall lack of association between 25(OH)D concentrations and sleep apnea.

The main strengths of our study are the large sample size and the community-based sample of older men. Our participants were not preselected for presence of any condition, particularly OSA, vitamin D deficiency, or obesity, thus minimizing selection biases. Despite the community-based sampling design, we had a wide distribution of AHI, 25(OH)D concentrations, and BMI, thus allowing us to analyze such relationships across a wide range of these variables seen in clinical practice. Additional strengths of our study include performance of 25(OH)D assays at an experienced, high-quality reference laboratory, careful review and cleaning of sleep study data, and accurate measurement of potential confounders of 25(OH)D and OSA.

Our study has some inherent limitations as well. Our study participants were generally healthy, largely white,

elderly males, so we cannot generalize these findings to nonwhite individuals, females, and younger patients. This was also a cross-sectional analysis with a single night's measurement of sleep and a single 25(OH)D measurement. Although our data suggest that low 25(OH)D is not likely to predict future development of OSA, a longitudinal study design would be required to specifically test such a hypothesis. We also note that, although measures of OSA typically display high night-to-night reliability (30), 25(OH)D levels vary by season, latitude, skin tone, sunscreen use, and time spent outdoors. We adjusted for season in our regression analysis, but we had no measures of within-participant seasonal variation in 25(OH)D concentrations or other factors that may have varied over time. Therefore, our single 25(OH)D measure may not fully reflect seasonal variations throughout the year.

An aim of our study was also to assess vitamin D status, irrespective of vitamin D source (outdoor sunlight exposure, diet, or oral supplement use). Our analyses were adjusted for season of blood draw and geographic region, but we acknowledge that different sources of vitamin D may have differential impacts on factors, such as seasonal variations in 25(OH)D concentrations and prevalence of individuals with vitamin D deficiency. Finally, we note that some data suggest that myopathy related to vitamin D deficiency may be most pronounced at very low 25(OH)D concentrations, such as less than 12 ng/ml (31). Because our community-dwelling cohort had very few persons with such low levels (only 2.1% had concentrations <12 ng/ml), our study was not able to adequately address whether profound vitamin D deficiency could be independently associated with OSA.

Conclusions

Among community-dwelling older men, the association between lower serum 25(OH)D concentrations and higher odds of sleep apnea was explained by greater BMI and larger neck circumference among those with lower 25(OH)D levels. ■

Author disclosures are available with the text of this article at www.atsjournals.org.

Acknowledgment: The authors thank the participants in the Osteoporotic Fractures in Men Sleep Study.

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