25-Hydroxyvitamin D and Parathyroid Hormone Levels Do Not Predict Changes in Carotid Arterial Stiffness

The Multi-Ethnic Study of Atherosclerosis

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Objective—To evaluate the impact of vitamin D and parathyroid hormone (PTH) on longitudinal changes in arterial stiffness.

Approach and Results—Distensibility coefficient and Young's elastic modulus of the right common carotid artery were evaluated at baseline and after a mean (SD) of 9.4 (0.5) years in 2580 Multi-Ethnic Study of Atherosclerosis (MESA) participants. Cross-sectional and longitudinal associations were evaluated using multivariable linear regression and analysis of covariance. At baseline, participants were 60.1 (9.4) years old (54% female; 26% black, 20% Hispanic, 14% Chinese). Mean annualized 25(OH)D was <20 ng/dL in 816 participants, and PTH was >65 pg/dL in 285 participants. In cross-sectional analyses, low 25(OH)D (<20 ng/mL) was not associated with stiffer arteries after adjustment for cardiovascular disease risk factors (P>0.4). PTH >65 pg/mL was associated with stiffer arteries after adjustment for cardiovascular disease risk factors, other than systolic blood pressure (distensibility coefficient: $\beta=-2.4\times10^{-4}$ mm Hg^{-1}, P=0.003; Young's elastic modulus: β=166 mm Hg, P=0.01); however, after adjustment for systolic blood pressure, these associations no longer were statistically significant. Longitudinal arterial stiffening was associated with older age (P<0.0001), higher systolic blood pressure (P<0.008), and use of antihypertensive medications (P<0.006), but not with 25(OH)D or PTH (both P>0.1).

Conclusions—Carotid arterial stiffness is not associated with low 25(OH)D concentrations. Cross-sectional associations between arterial stiffness and high PTH were attenuated by systolic blood pressure. After nearly a decade of follow-up, neither baseline PTH nor 25(OH)D concentrations were associated with progression of carotid arterial stiffness. (***Arterioscler Thromb Vasc Biol*. 2014;34:1102-1109.)**

Key Words: cardiovascular diseases ■ carotid arteries ■ parathyroid hormone ■ vascular stiffness ■ vitamin D

Vitamin D deficiency and hyperparathyroidism are associated with cardiovascular disease (CVD) risk.[1–9] Low circulating concentrations of 25-hydroxyvitamin D (25[OH]D) and elevated parathyroid hormone (PTH) have been linked to hypertension, insulin resistance, metabolic syndrome, coronary heart disease, congestive heart failure, CVD, and death.[1,7–14]

Increased arterial stiffness is associated with aging, fragmentation of elastin fibers, and a decrease in the elastin-to-collagen ratio in arterial walls.[15] This process may underlie the development of hypertension, CVD, cerebral dysfunction, and stroke[16–20] because a rigid arterial tree is less able to accommodate the large pulsatile output from the heart. Increased vascular stiffness accelerates atherogenesis and is associated with an increase in cardiac morbidity and mortality.[21] Vitamin D and PTH are closely linked and may affect vascular smooth muscle tone through the renin–angiotensin–aldosterone axis[22] and may promote vascular endothelial growth factor.[23] Additionally, lymphocyte and monocyte/phagocyte differentiations are modulated by vitamin D, thereby affecting the release of inflammatory cytokines that promote arterial plaque formation[24] because heightened vascular smooth muscle tone, endothelial dysfunction, and plaque formation are directly linked to hypertension, coronary artery disease, and stroke. Increased vascular stiffness is a plausible mechanism through which 25(OH)D and PTH may affect CVD risk.[2,16,17,21,25,26]

Structural and functional alterations in the arterial bed, such as circumferential widening of large arteries and wall thickening, lead to changes in carotid artery distensibility and elasticity, measured with distensibility coefficient (DC) and Young's elastic modulus (YEM), respectively. These are validated, noninvasive measures of arterial function, which characterize arterial stiffness[15,27] and can identify individuals at increased CVD risk.[21] Both measure the ability of an artery to expand and contract with each cardiac pulsation;

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Nonstandard Abbreviations and Acronyms	
CVD	cardiovascular disease
DC	distensibility coefficient
PTH	parathyroid hormone
SBP	systolic blood pressure
YEM	Young's elastic modulus

however, the major difference between these stiffness parameters is that YEM accounts for carotid artery wall thickness in an attempt to separate whether arterial stiffening is solely related to pressure differences or intrinsic changes in the arterial wall.[15,18,27]

A limited number of studies have evaluated the associations of elevated PTH and low 25(OH)D with increased arterial stiffness; however, these studies are limited by their small sample size and their cross-sectional design.[2,28–30] The aim of this study was to explore the relationship between markers of bone-mineral metabolism and changes in arterial stiffness in an ethnically diverse cohort without clinically evident CVD.

Materials and Methods

Materials and Methods are available in the online-only Supplement.

Results

Baseline Characteristics

Baseline characteristics are shown in Table 1. Participants were a mean (SD) of 60.1 (9.4) years old, 54% were female, 39.5% were white, 25.5% were black, 20.5% were Hispanic, and 14.5% were Chinese. The mean annualized 25(OH)D was 26.3 (11.5) ng/mL and was <20 ng/dL in 816 (30%) participants and 20 to 30 ng/mL in 973 (36%) participants. The mean PTH was 43.5 (18.8) pg/dL and was >65 pg/dL in 285 (11%) participants; 86% of subjects graduated from high school and 44% earned <$40000. The average physical activity score was 1665 MET-min/wk. At baseline, the mean DC was 3.1 (1.3)×10^{-3} mm Hg^{-1}, and the mean YEM was 1591 (938) mm Hg.

Cross-Sectional Associations With Arterial Stiffness Measurements

In cross-sectional analyses, continuous 25(OH)D was not associated with stiffness parameters before or after adjustment for CVD risk factors (P>0.1; Tables 2 and 3). When grouped by category of 25(OH)D concentrations, no significant trend toward increasing stiffness with lower 25(OH)D was observed after adjustment for traditional CVD risk factors (P>0.3; Figure). The strongest association with increased stiffness at examination 1 was seen in participants with 25(OH)D concentrations <20 ng/mL (lower DC, β=−1.6×10^{-4} mm Hg^{-1}, P=0.01; higher YEM, β=107.2 mm Hg, P=0.03); however, these associations disappeared after adjustment for traditional CVD risk factors (P>0.4). As a continuous variable, 25(OH)D concentration was not associated with arterial stiffness (DC, β=−2.4×10^{-7} mm Hg^{-1}, P=0.91; YEM, β=0.2 mm Hg, P=0.92; Tables 2 and 3, cross-sectional model 3).

At baseline, higher PTH concentrations were associated with greater stiffness demonstrated by lower DC (β=−2.5×10^{-6} mm Hg^{-1}, P=0.04) and higher YEM (β=1.98 mm Hg, P=0.06; Figure). This relationship seemed to be nonlinear, with overtly elevated PTH concentrations (≥65 pg/mL) being most strongly associated with differences in DC and YEM. Adjusting for CVD risk factors other than blood pressure, PTH >65 pg/mL was associated with lower DC (β=−2.4×10^{-4} mm Hg^{-1}, P=0.003) and higher YEM (β=166 mm Hg, P=0.01; Tables 4 and 5). However, these associations no longer were statistically significant when baseline systolic blood pressure (SBP) was included in the model (DC: β=−1.4×10^{-4} mm Hg^{-1}, P=0.08; YEM: β=118 mm Hg, P=0.07).

Within race/ethnicity groups, there were no significant associations between baseline 25(OH)D and YEM or DC (all P>0.05). The associations of PTH with DC and YEM seemed strongest for Hispanic participants (DC: β=−3.6×10^{-4} mm Hg^{-1}, P=0.02; YEM: β=275 mm Hg, P=0.04), but the P values for the interaction of race/ethnicity with PTH were not statistically significant for DC (P=0.15) or YEM (P=0.08).

Longitudinal Associations With Arterial Stiffness Measurements

DC decreased from 3.1 (1.3)×10^{-3} mm Hg^{-1} at examination 1 to 2.7 (1.2)×10^{-3} mm Hg^{-1} at examination 5, and YEM increased from 1591 (938) mm Hg at examination 1 to 1754 (1340) mm Hg at examination 5, both indicating progression of arterial stiffness during the follow-up period. Longitudinal changes in DC and YEM were associated with older age (DC: β=−2.0×10^{-5} mm Hg^{-1}, per year, P<0.0001; YEM: β=13.4 mm Hg, per year, P<0.0001) and higher SBP (DC: β=−2.9×10^{-6} mm Hg,[1] P=0.007) and use of antihypertensive medication (YEM: β=157.4 mm Hg, P=0.006). No associations or even trends were observed between baseline 25(OH)D or PTH and carotid stiffness (all P>0.3) with or without adjustment for baseline DC and YEM. Additionally, those with baseline PTH >65 pg/mL or 25(OH)D <20 also were not associated with a significant change in DC or change in YEM after nearly 10 years of follow-up (Tables 2 and 3) compared with the reference groups.

Within race/ethnicity groups, no significant associations between 25(OH)D and PTH with longitudinal changes in YEM or DC were observed (all P>0.05), and the P values for the interaction of race/ethnicity with PTH were not statistically significant for changes in DC (P=0.15) or YEM (P=0.96).

Discussion

In the current analysis, we observed a cross-sectional association of higher PTH concentrations with increased arterial stiffness that was independent of CVD risk factors except baseline SBP. No associations were present for 25-OHD. After nearly a decade of aging, neither baseline PTH nor 25-OHD concentrations were associated with changes in arterial stiffening.

Potentially deleterious effects of vitamin D deficiency on CVD risk have been described and have even led some clinicians to promote vitamin D supplementation for CVD risk reduction.[1,5–7,31] A relationship between low vitamin D concentrations and increased arterial stiffness has been described in

Table 1. Baseline Participant Characteristics

		Annualized 25(OH) Vitamin D, ng/mL			Parathyroid Hormone, pg/mL			
	All Subjects	<20.0	20.0–29.9	≥30.0	Tertile 1 <32.8	Tertile 2 32.8–43.7	Tertile 3 43.8–65.0	>65.0
Number of subjects	2707	816	973	918	806	808	808	285
Age, y	60.1 (9.4)	58.6 (9.2)	60.3 (9.4)	61.3 (9.3)	58.8 (9.2)	59.6 (9.5)	61.3 (9.3)	61.9 (9.2)
Female sex, %	1449 (53.5)	459 (56.3)	481 (49.4)	509 (55.5)	429 (53.2)	415 (51.4)	429 (53.1)	176 (61.8)
Ethnicity, %								
White	1070 (39.5)	145 (17.8)	388 (39.9)	537 (58.5)	395 (49.0)	328 (40.6)	276 (34.2)	71 (24.9)
Black	691 (25.5)	406 (49.8)	194 (19.9)	91 (9.9)	127 (15.8)	189 (23.4)	259 (32.1)	116 (40.7)
Chinese	392 (14.5)	93 (11.4)	176 (18.1)	123 (13.4)	155 (19.2)	138 (17.1)	86 (10.6)	13 (4.6)
Hispanic	554 (20.5)	172 (21.1)	215 (22.1)	167 (18.2)	129 (16.0)	153 (18.9)	187 (23.1)	85 (29.8)
Blood pressure parameters, mm Hg								
SBP	123.7 (20.1)	125.8 (20.9)	123.5 (20.0)	122.0 (19.4)	119.6 (19.0)	121.7 (18.9)	127.0 (20.5)	131.3 (22.0)
DBP	71.7 (10.1)	73.1 (10.2)	71.7 (10.0)	70.5 (9.9)	70.6 (9.6)	71.3 (10.0)	72.7 (10.2)	73.3 (10.8)
Hypertension, %	1160 (42.9)	376 (46.1)	426 (43.8)	358 (39.0)	277 (34.4)	323 (40.0)	398 (49.3)	162 (56.8)
HTN meds, %	896 (33.1)	295 (36.2)	330 (33.9)	271 (29.5)	212 (26.3)	258 (31.9)	304 (37.7)	122 (42.8)
Diabetes mellitus status, %								
IFG	329 (12.2)	120 (14.7)	122 (12.6)	87 (9.5)	86 (10.7)	95 (11.8)	109 (13.5)	39 (13.7)
Untreated	43 (1.6)	14 (1.7)	25 (2.6)	4 (0.4)	12 (1.5)	8 (1.0)	18 (2.2)	5 (1.8)
Treated	200 (7.4)	76 (9.3)	83 (8.5)	41 (4.5)	68 (8.5)	53 (6.6)	54 (6.7)	25 (8.8)
Lipid levels, mg/dL								
Total cholesterol	194.1 (34.8)	193.4 (36.6)	192.8 (34.3)	196.0 (33.6)	195.1 (34.9)	193.1 (34.7)	195.2 (34.4)	190.5 (35.7)
LDL-C	117.1 (30.4)	118.7 (32.5)	116.5 (29.9)	116.3 (29.1)	117.3 (29.3)	116.8 (30.8)	118.3 (30.5)	114.1 (32.4)
HDL-C	51.6 (15.3)	50.3 (14.7)	50.1 (14.7)	54.3 (16.0)	51.8 (14.6)	50.8 (15.0)	51.7 (15.9)	52.8 (16.1)
Triglycerides	127.7 (81.2)	122.3 (88.9)	132.0 (78.0)	128.1 (77.2)	130.1 (78.7)	128.3 (90.2)	128.6 (79.7)	117.1 (63.3)
Lipid-lowering meds, %	421 (15.6)	109 (13.4)	157 (16.1)	155 (16.9)	112 (13.9)	118 (14.6)	135 (16.7)	56 (19.7)
BMI, kg/m^2	27.8 (5.0)	29.2 (5.3)	27.7 (4.8)	26.5 (4.4)	26.5 (4.5)	27.6 (4.8)	28.5 (5.0)	29.7 (5.7)
Waist, cm	96.4 (13.7)	99.4 (14.0)	96.7 (13.7)	93.3 (12.6)	93.0 (12.9)	95.8 (13.4)	98.5 (13.1)	101.5 (15.4)
Smoking status, %								
Former	965 (35.7)	283 (34.7)	343 (35.3)	339 (36.9)	289 (35.9)	280 (34.7)	295 (36.5)	101 (35.4)
Current	312 (11.5)	121 (14.9)	105 (10.8)	86 (9.4)	103 (12.8)	100 (12.4)	84 (10.4)	25 (8.8)
Creatinine, mg/dL	0.9 (0.2)	0.9 (0.2)	0.9 (0.2)	0.9 (0.2)	0.9 (0.2)	0.9 (0.2)	0.9 (0.2)	1.0 (0.3)
Calcium, mg/dL	9.6 (0.4)	9.6 (0.4)	9.6 (0.4)	9.7 (0.4)	9.6 (0.4)	9.7 (0.4)	9.6 (0.4)	9.6 (0.4)
Phosphorous, mg/dL	3.7 (0.5)	3.7 (0.5)	3.6 (0.5)	3.7 (0.5)	3.8 (0.5)	3.7 (0.5)	3.6 (0.5)	3.5 (0.5)
CRP, mg/L	3.3 (4.9)	3.8 (4.8)	3.1 (4.9)	3.1 (5.0)	2.7 (4.0)	3.3 (5.0)	3.6 (4.7)	4.3 (6.9)
25(OH)D, ng/mL	26.3 (11.5)	14.3 (3.8)	25.0 (2.9)	38.4 (9.7)	30.2 (11.1)	27.5 (11.8)	23.6 (10.6)	19.8 (9.4)
Parathyroid hormone, pg/mL	43.5 (18.8)	51.4 (22.7)	42.7 (16.7)	37.2 (14.0)	26.1 (4.8)	38.0 (3.3)	52.6 (5.9)	82.2 (22.2)
Carotid wall thickness, cm	0.148 (0.031)	0.150 (0.032)	0.148 (0.031)	0.147 (0.030)	0.142 (0.029)	0.147 (0.030)	0.153 (0.033)	0.153 (0.032)
PSI diameter, cm	0.628 (0.075)	0.630 (0.075)	0.631 (0.074)	0.623 (0.075)	0.620 (0.072)	0.628 (0.074)	0.633 (0.074)	0.636 (0.081)
EDI diameter, cm	0.582 (0.071)	0.584 (0.071)	0.584 (0.070)	0.576 (0.071)	0.573 (0.068)	0.582 (0.070)	0.586 (0.071)	0.591 (0.076)
YEM, mm Hg	1591 (938)	1652 (1011)	1599 (904)	1528 (902)	1493 (799)	1574 (915)	1626 (1044)	1815 (1006)
DC, 10^{-3} mm Hg^{-1}	3.1 (1.3)	3.0 (1.2)	3.1 (1.3)	3.2 (1.2)	3.3 (1.3)	3.2 (1.3)	3.0 (1.2)	2.7 (1.1)

All values are mean (SD) unless noted otherwise. 25(OH)D indicates serum 25-hyroxyvitamin D; BMI, body mass index; CRP, c-reactive protein; DBP, diastolic blood pressure; DC, distensibility coefficient; EDI, end-diastolic internal; HDL-C, high-density lipoprotein–cholesterol; HTN, hypertension; IFG, impaired fasting glucose; LDL-C, low-density lipoprotein–cholesterol; meds, medication; NA, not applicable; PSI, peak systolic internal; SBP, systolic blood pressure; and YEM, Young's elastic modulus.

cross-sectional observational studies[2,25,29]; however, the effects of vitamin D status on longitudinal changes in arterial stiffness are less clear. Our results are in accordance with small randomized controlled trials of vitamin D supplementation, which failed to demonstrate improvements in arterial stiffness with vitamin D supplementation,[28,32] although the longest of these trials only followed subjects for 3 years.[28] Relationships between vitamin D concentrations and CVD end points have been mixed. For example, low vitamin D concentrations have been associated with increased risk of incident coronary heart disease[33] and presence of coronary artery calcium,[34,35] but not with congestive heart failure or carotid intima-media

Table 2. Cross-Sectional and Longitudinal Associations of Serum 25(OH)D Concentrations and Distensibility Coefficient

	Cross-Sectional Analyses					Longitudinal Analyses*				
	Baseline Distensibility Coefficient, mm Hg^{-1}×10^{-4} (95% Confidence Interval)					Change in Distensibility Coefficient, mm Hg^{-1}×10^{-4}† (95% Confidence Interval)				
25(OH)D, ng/mL		Beta Parameter					Beta Parameter			
	n	Model 1‡	Model 2‡	Model 3§	Model 4§	n	Model 1‖	Model 2‖	Model 3¶	Model 4¶
≥30.0	918	Ref	Ref	ref	ref	872	ref	ref	ref	Ref
20.0–29.9	973	–0.5 (–1.5, 0.6)	0.1 (–1.0, 1.1)	0.2 (–0.8, 1.3)	0.3 (–0.7, 1.3)	935	–0.1 (–1.0, 0.8)	0.1 (–0.8, 1.0)	0.2 (–0.7, 1.1)	0.2 (–0.7, 1.1)
<20	816	–1.6# (–2.8, –0.4)	–0.7 (–1.9, 0.5)	–0.6 (–1.8, 0.6)	–0.3 (–1.5, 0.8)	773	0.3 (–0.7, 1.3)	0.5 (–0.5, 1.6)	0.6 (–0.5, 1.6)	0.6 (–0.4, 1.6)
P trend	-	0.01	0.33	0.46	0.69	-	0.66	0.35	0.30	0.28

Model 1 was adjusted for age, sex, race, study field center, education, and income. Model 2 was Model 1 plus physical activity, waist circumference, smoking status, and body mass index. Model 3 was Model 2 plus diabetes mellitus status, antihypertensive medication use, log[c-reactive protein], total cholesterol, high-density lipoprotein–cholesterol, lipid-lowering therapy, and creatinine. Model 4 was Model 3 plus systolic blood pressure.

*Longitudinal analyses shown with adjustment for baseline stiffness measures.

†Between the 2 carotid ultrasounds (9.4 y).

‡Sixty-seven participants missing data on covariates.

§Eighty-three participants missing data on covariates.

‖Sixty-two participants missing data on covariates.

¶Seventy-seven participants missing data on covariates.

#*P*<0.01.

thickness.[36] Associations between low circulating vitamin D concentration and CVD risk may also be partly confounded by CVD risk factors such as obesity and inactivity.[37] The only large, long-term randomized controlled trial of vitamin D supplementation showed no change in CVD events during a 7-year period.[38]

High PTH concentrations have been associated with poor CVD outcomes[9] in observational studies.[29,30] In previous cross-sectional analysis among the Multi-Ethnic Study of Atherosclerosis (MESA) participants, higher PTH concentrations were associated with increased blood pressure, higher central aortic pressure, and lower large artery elasticity.[39] PTH levels seem to be more strongly associated with congestive heart failure events than coronary heart disease events.[9] The results of the present study agree with previously reported studies describing cross-sectional associations between elevated PTH and increased carotid stiffness measures; however, the results were blunted when SBP was included in the model. This suggests that the cross-sectional associations between arterial stiffness and PTH may be mediated through blood pressure. It also is possible that the baseline SBP is more collinear with DC and YEM because pulse pressure, which takes blood pressure into account, is a part of the formulae used to calculate these outcome measures.

Table 3. Cross-Sectional and Longitudinal Associations of Serum 25(OH)D Concentrations and Young's Elastic Modulus

	Cross-Sectional Analyses					Longitudinal Analyses*				
	Baseline Young's Elastic Modulus, mm Hg (95% Confidence Interval)					Change in Young's Elastic Modulus, mm Hg† (95% Confidence Interval)				
25(OH)D, ng/mL		Beta Parameter					Beta Parameter			
	n	Model 1‡	Model 2‡	Model 3§	Model 4§	n	Model 1‖	Model 2‖	Model 3¶	Model 4¶
≥30.0	918	ref	Ref	ref	Ref	872	ref	ref	ref	ref
20.0–29.9	973	50.2 (–34.1, 134.6)	24.0 (–60.7, 108.7)	13.1 (–72.0, 98.2)	12.3 (–71.8, 96.3)	935	–75.6 (–188.3, 37.2)	–92.7 (–206.4, 21.1)	–91.0 (–205.4, 23.4)	–91.2 (–205.5, 23.2)
<20	816	107.2# (11.0, 203.3)	59.5 (–38.5, 157.5)	53.5 (–45.1, 152.2)	42.2 (–55.1, 139.6)	773	–87.5 (–216.1, 41.2)	–109.5 (–241.1, 22.1)	–100.3 (–232.6, 32.1)	–102.2 (–234.6, 30.1)
P trend	-	0.03	0.24	0.32	0.42	-	0.15	0.08	0.11	0.10

Model 1 was adjusted for age, sex, race, study field center, education, and income. Model 2 was Model 1 plus physical activity, waist circumference, smoking status, and body mass index. Model 3 was Model 2 plus diabetes mellitus status, antihypertensive medication use, log[c-reactive protein], total cholesterol, high-density lipoprotein–cholesterol, lipid-lowering therapy, and creatinine. Model 4 was Model 3 plus systolic blood pressure.

*Longitudinal analyses shown with adjustment for baseline stiffness measures.

†Between the 2 carotid ultrasounds (9.4 y).

‡Sixty-seven participants missing data on covariates.

§Eighty-three participants missing data on covariates.

‖Sixty-two participants missing data on covariates.

¶Seventy-seven participants missing data on covariates.

#*P*<0.05.

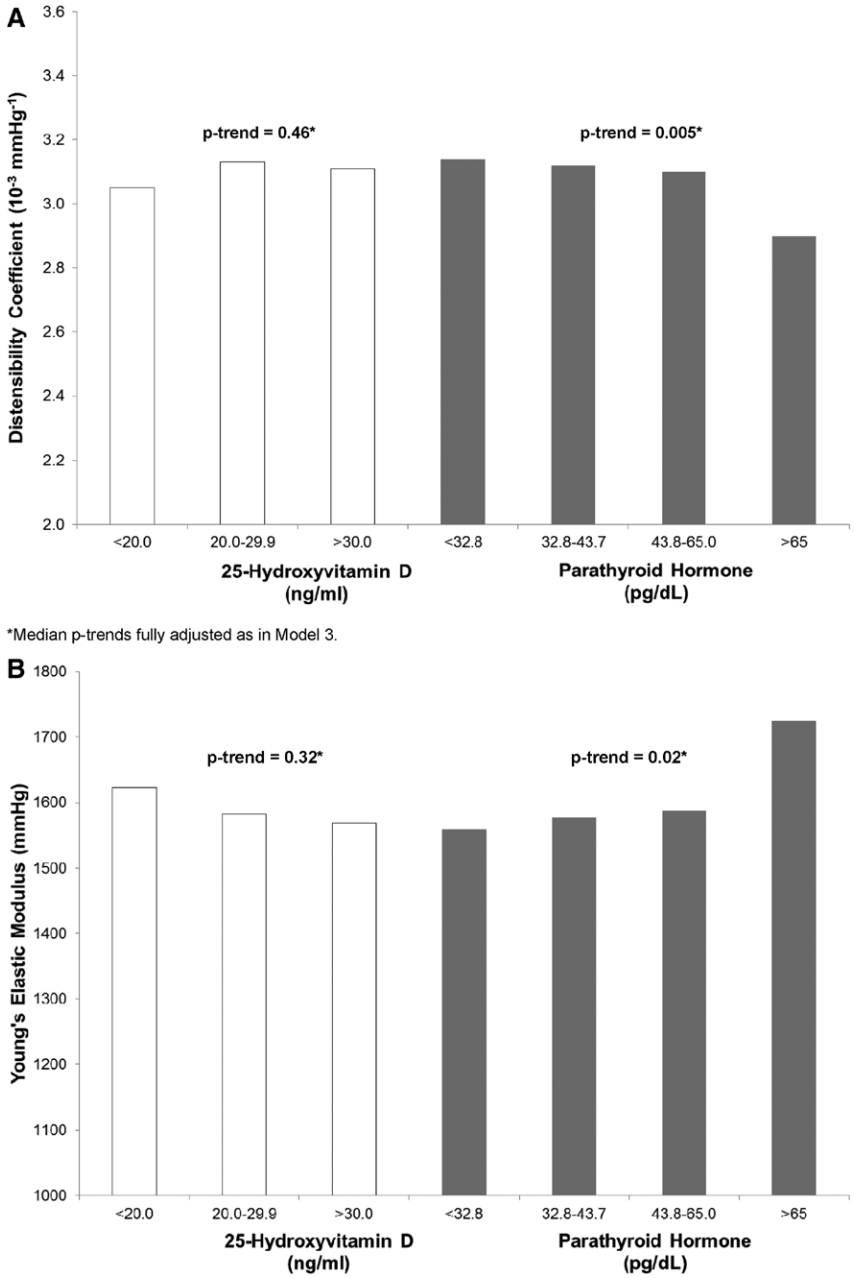

Figure. A, Baseline distensibility coefficient by 25-hydroxyvitamin D and parathyroid hormone concentrations. **B**, Baseline Young's elastic modulus by 25-hydroxyvitamin D and parathyroid hormone concentrations.

It may be expected that higher PTH concentrations at baseline would lead to more rapid progression of arterial stiffness during a decade of aging; however, we did not observe a longitudinal association between baseline PTH concentrations and progressive arterial stiffening. Because those with the highest PTH levels also were found to have stiffer arteries at baseline, acceleration of stiffness over time may be blunted because there could be less physiological room for progression of the carotid stiffness parameters (ceiling effect). However, when baseline stiffness parameters were included in the models to attempt to account for this discrepancy, still no associations or even consistent trends were observed. Although PTH and vitamin D were not longitudinally associated with changes in YEM and DC, acceleration of stiffness parameters was observed as expected with traditional CVD risk factors, such as advancing age and hypertension.[40] Alternatively, although 25(OH)D has a relatively long circulating half-life (≈3 weeks) and is considered a good biomarker, a single measurement may not fully capture cumulative vitamin D exposure during a 10-year period, resulting in misclassification. It is similar for PTH, which comparatively has a much shorter half-life (2–4 minutes), making it even more subject to misclassification. A single measurement seems adequate for cross-sectional associations but is less useful during a decade of follow-up.

Several limitations of the current study should be considered. First, we imaged the carotid arteries but measured brachial artery blood pressure. Brachial artery pressure is considered to be a surrogate for central aortic pressure, but may overestimate stiffness measurements because brachial measurements can overestimate central pressures, although the

Table 4. Cross-Sectional and Longitudinal Associations Between Parathyroid Hormone and Distensibility Coefficient

	Cross-Sectional Analyses					Longitudinal Analyses*				
	Baseline Distensibility Coefficient, mm Hg^{-1}×10^{-4}					Change in Distensibility Coefficient, mm Hg^{-1}×10^{-4}†				
	(95% Confidence Interval)					(95% Confidence Interval)				
PTH, pg/mL	n	Beta Parameter				n	Beta Parameter			
		Model 1‡	Model 2‡	Model 3§	Model 4§		Model 1‖	Model 2‖	Model 3¶	Model 4¶
<32.8	806	Ref	Ref	ref	ref	765	ref	ref	ref	Ref
32.8–43.7	808	–0.6 (–1.7, 0.5)	–0.2 (–1.3, 1.0)	–0.2 (–1.3, 0.9)	–0.2 (–1.3, 0.9)	772	1.1# (0.1, 2.0)	1.2# (0.2, 2.1)	1.0 (0.0, 1.9)	1.0 (0.0, 1.9)
43.8–65.0	808	–1.2# (–2.3, –0.0)	–0.4 (–1.6, 0.7)	–0.5 (–1.6, 0.7)	0.0 (–1.1, 1.1)	777	0.4 (–0.6, 1.4)	0.6 (–0.4, 1.5)	0.5 (–0.5, 1.4)	0.5 (–0.4, 1.5)
>65	285	–3.4** (–5.0, –1.8)	–2.2** (–3.8, –0.5)	–2.4** (–4.0, –0.8)	–1.4 (–3.0, 0.1)	266	–0.3 (–1.6, 1.1)	0.0 (–1.4, 1.4)	–0.2 (–1.6, 1.2)	–0.1 (–1.4, 1.3)
P trend	-	<0.001	0.01	0.005	0.16	-	0.61	0.93	0.73	0.91

Model 1 was adjusted for age, sex, race, study field center, education, and income. Model 2 was Model 1 plus physical activity, waist circumference, smoking status, and body mass index. Model 3 was Model 2 plus diabetes mellitus status, antihypertensive medication use, log[c-reactive protein], total cholesterol, high-density lipoprotein–cholesterol, lipid-lowering therapy, and creatinine. Model 4 was Model 3 plus systolic blood pressure.

*Longitudinal analyses shown with adjustment for baseline stiffness measures.

†Between the 2 carotid ultrasounds (9.4 y).

‡Sixty-seven participants missing data on covariates.

§Eighty-three participants missing data on covariates.

‖Sixty-two participants missing data on covariates.

¶Seventy-seven participants missing data on covariates.

#*P*<0.05.

*******P*<0.01.

2 measures are highly correlated, especially in older adults.[41] Vitamin D and PTH status were defined by baseline concentrations, and the absence of follow-up levels of either hormone poses a challenge to the interpretation of the longitudinal analyses. This potentially is more likely to be an issue with vitamin D because supplementation is common in the general population, and we did not have information concerning vitamin D supplementation during the follow-up period. Also, the race/ethnicity subgroup analyses may be limited by small sample size. Because all participants had ultrasound studies at

Table 5. Cross-Sectional and Longitudinal Associations Between Parathyroid Hormone and Young's Elastic Modulus

	Cross-Sectional Analyses					Longitudinal Analyses*				
	Baseline Young's Elastic Modulus, mm Hg					Change in Young's Elastic Modulus, mm Hg†				
	(95% Confidence Interval)					(95% Confidence Interval)				
PTH, pg/mL	n	Beta Parameter				n	Beta Parameter			
		Model 1‡	Model 2‡	Model 3§	Model 4§		Model 1‖	Model 2‖	Model 3¶	Model 4¶
<32.8	806	ref.	ref.	ref.	ref.	765	ref.	ref.	ref.	ref.
32.8–43.7	808	39.2 (–50.9, 129.4)	20.1 (–70.0, 110.1)	17.9 (–72.3, 108.1)	18.3 (–70.8, 107.3)	772	–63.5 (–184.3, 57.2)	–67.9 (–189.0, 53.2)	–56.8 –178.1, 64.5)	–56.5 (–177.8, 64.8)
43.8–65.0	808	63.4 (–28.7, 155.5)	28.4 (–64.2, 120.9)	28.6 (–64.2, 121.5)	5.6 (–86.2, 97.5)	777	–59.0 (–182.1. 64.2)	–73.2 (–197.6, 51.2)	–57.5 (–182.2, 67.3)	–62.7 (–187.7, 62.2)
>65	285	217.0# (88.1, 345.9)	159.0** (28.8, 289.2)	166.2** (35.6, 296.7)	117.6 (–11.8, 247.0)	266	110.3 (–63.9, 284.5)	88.1 (–88.3, 264.6)	80.5 (–96.5, 257.5)	70.9 (–106.5, 248.4)
P trend	-	0.002	0.03	0.02	0.14	-	0.37	0.54	0.55	0.64

Model 1 was adjusted for age, sex, race, study field center, education, and income. Model 2 was Model 1 plus physical activity, waist circumference, smoking status, and body mass index. Model 3 was Model 2 plus diabetes mellitus status, antihypertensive medication use, log[c-reactive protein], total cholesterol, high-density lipoprotein–cholesterol, lipid-lowering therapy, and creatinine. Model 4 was Model 3 plus systolic blood pressure.

*Longitudinal analyses shown with adjustment for baseline stiffness measures.

†Between the 2 carotid ultrasounds (9.4 y).

‡Sixty-seven participants missing data on covariates.

§Eighty-three participants missing data on covariates.

‖Sixty-two participants missing data on covariates.

¶Seventy-seven participants missing data on covariates.

#*P*<0.01.

*******P*<0.05.

exams 1 and 5, there may be a survivorship bias. Participants who were followed up to examination 5 were healthier and less likely to have a nonfatal CVD event than the complete MESA cohort, which would likely result in a null bias.

Conclusions

In cross-sectional analyses, we did not observe any independent associations between arterial stiffness measures and vitamin D status. Carotid arterial stiffness was associated with PTH concentrations >65 mg/dL; however, the associations were attenuated by adjustment for SBP. In longitudinal analyses, advancing age and hypertension were associated with progression of arterial stiffness; however, neither baseline PTH nor 25(OH)D was associated with changes in arterial stiffness measures after nearly a decade of follow-up. Elevated PTH is associated with carotid stiffness; however, the causal and temporal inter-relationships of PTH, blood pressure, and carotid stiffness are not entirely clear and warrant further study.

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Significance

Low vitamin D and high parathyroid hormone concentrations have been associated with heart disease and hypertension, but much less is known about their long-term effects on arterial stiffening, which has been linked to the development of heart failure, strokes, and heart attacks. In this study, carotid artery stiffness was associated with high parathyroid hormone levels, although this finding was attenuated by systolic blood pressure. Vitamin D concentration was not associated with baseline arterial stiffness. Neither baseline parathyroid hormone nor vitamin D concentrations were associated with changes in arterial stiffening during nearly a decade of follow-up. These findings suggest that parathyroid hormone may impact the development of arterial stiffness; however, the causal and temporal inter-relationships of parathyroid hormone, blood pressure, and carotid stiffness are not entirely clear and warrant further study.