

Multivitamin use and telomere length in women^{1–3}

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ABSTRACT

Background: Telomere length may be a marker of biological aging. Multivitamin supplements represent a major source of micronutrients, which may affect telomere length by modulating oxidative stress and chronic inflammation.

Objective: The objective was to examine whether multivitamin use is associated with longer telomeres in women.

Design: We performed a cross-sectional analysis of data from 586 early participants (age 35–74 y) in the Sister Study. Multivitamin use and nutrient intakes were assessed with a 146-item food-frequency questionnaire, and relative telomere length of leukocyte DNA was measured by quantitative polymerase chain reaction.

Results: After age and other potential confounders were adjusted for, multivitamin use was associated with longer telomeres. Compared with nonusers, the relative telomere length of leukocyte DNA was on average 5.1% longer among daily multivitamin users (P for trend = 0.002). In the analysis of micronutrients, higher intakes of vitamins C and E from foods were each associated with longer telomeres, even after adjustment for multivitamin use. Furthermore, intakes of both nutrients were associated with telomere length among women who did not take multivitamins.

Conclusion: This study provides the first epidemiologic evidence that multivitamin use is associated with longer telomere length among women. *Am J Clin Nutr* 2009;89:1857–63.

SUBJECTS AND METHODS

Study population

The Sister Study (<http://www.sisterstudy.org/>) is an ongoing risk-enriched prospective cohort of healthy sisters (age 35–74 y) of breast cancer patients (17). Recruitment began in 2004 and is expected to be completed in 2009. The enrollment includes a home visit for blood and urine collection, a 90-min computer-assisted telephone interview, and several self-administered questionnaires, including a detailed food-frequency questionnaire (FFQ). Details of this telomere project and sampling procedures were described elsewhere (18). Briefly, a total of 740 women were selected for telomere measurement from the first 2086 Sister Study participants by oversampling smokers, nonwhite women, and women with high perceived stress and by randomly sampling the rest. Exclusion criteria included missing or ineligible biological specimens, missing race or smoking data, major dental procedure or surgery in the past week, working on rotating shifts, recent chemotherapy or radiation treatment of cancer, or a diagnosis of breast cancer before the first annual follow-up. The sample selection criteria and sample size reflect the requirements for a US Department of Defense-funded study on stress and telomere length. We limited our analyses to 586 women with valid dietary information and duplicate laboratory assays on telomere length. The Sister Study was approved by the Institutional Review Board of the National Institute of Environmental Health Science, National Institutes of Health.

INTRODUCTION

Telomeres, the TTAGGG tandem repeat sequence, and their binding proteins at the ends of chromosomes prevent chromosomes from detrimental recombination and degradation (1). In somatic cells, the length of telomeres decreases with each cell division, which may eventually lead to cell senescence or apoptosis. Therefore, telomere length has been proposed as a marker of “biological ageing” (2). Consistent with this hypothesis, preliminary epidemiologic studies have related shorter telomeres to higher mortality (3) and higher risk of some age-related chronic diseases (4–10). Experimental evidence suggests that oxidative stress and chronic inflammation contribute to the attrition of telomeres (2, 11). Several micronutrients, such as antioxidant vitamins and minerals, can modulate the states of oxidative stress and chronic inflammation and therefore may affect telomere length (12–15). Multivitamin supplements contain large amounts of many vitamins and minerals and therefore represent a major source of micronutrient intake (16). We therefore examined whether multivitamin use was associated with longer telomeres among 586 women from the Sister Study.

Telomere length measurement

A whole blood sample was collected at enrollment and stored at -80°C until DNA extraction. Total leukocyte DNA was then used as a template for polymerase chain reaction (PCR)-based

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measurement of relative telomere length according to previously published protocols (19). The assay used 100–200 ng template DNA in 1- μ L aliquots for triplicate PCR amplifications per sample per plate. Cycle threshold was transformed into nanograms DNA based on a standard curve. The quantitative assay determines the amount of telomeric DNA (T) relative to the amount of single-copy control gene (human β -globin) DNA (S) and then calculates a T/S ratio. This PCR-based telomere assay was found to be highly correlated with Southern blot analysis (19). We further estimated the relative length of telomeres in base pairs (bp) by multiplying the T/S ratio with a constant of 4270 (19). Whereas this constant was derived and validated in another study, the assays here were performed in the same laboratory by using the same genetic controls. Of the 740 specimens submitted for telomere assays, 647 were run on duplicate plates including 3 internal controls (one each at a high, medium, and low T/S ratio) to further account for variation over time and plates. The CV across averaged adjusted replicates was 8.5%. These average plate-adjusted values were used for the present analyses.

Exposure assessment

The Sister Study dietary survey was based on a modified Block 1998 FFQ with additional questions and changes (20, 21). The FFQ asked for the portion size and frequency of consumption of 146 food items in the past 12 mo. Intake of individual nutrients was then calculated with software from the Block Dietary Data Systems (Berkeley, CA). Participants were asked whether they had taken any vitamins or minerals regularly (at least once per month) during the past 12 mo. For those who answered “yes,” the FFQ further elicited details about the use of 3 types of multivitamins [regular once-a-day, Centrum (Wyeth Consumer Healthcare, Madison, NJ), or Thera type; stress-tabs or B-complex type; and antioxidant combination type] and 16 individual supplements of vitamins or minerals. For each supplement, participants were asked the frequency of use (ranging from “did not take” to “every day”) and, for users, the duration of use (ranging from “less than 1 year” to “10+ years”). We further calculated an overall frequency variable for multivitamin use by combining the use of all 3 types of multivitamins. The Sister Study also collected information on age, race, education, smoking status, perceived stress level, self-reported health status, adult-onset diabetes, and cardiovascular diseases (ie, heart attack, bypass surgery, angioplasty, congestive heart failure, cardiac arrhythmic, medicated angina, and stroke/transient ischemic attack). Body weight and height were measured during a home visit for blood collection, and body mass index (BMI) was calculated by dividing weight in kilograms by height squared in meters.

Statistical analyses

For multivitamins, the frequency and duration of use were defined categorically. The use of most individual vitamin and mineral supplements was infrequent; therefore, the participants were classified as users or nonusers. We compared the population characteristics by multivitamin use status using Student's *t* test for continuous variables and a chi-square test for categorical variables. Dietary intake of micronutrients was categorized into quartiles ($\approx 25\%$ of the participants in each quartile) after adjustment for energy intake with the residual method (22). The

least-square means and SEM of relative telomere length for each exposure category were calculated with generalized linear regression models, adjusted for age (continuous), race (non-Hispanic white and others), BMI (continuous), education (\leq high school, some college, associate degree/technical training, college graduate, and graduate degrees), cigarette smoking (never, former, and current smokers), presence of diabetes or cardiovascular diseases (yes, no), energy intake (continuous), perceived stress level (very low, low, moderate, high, and very high), self-reported health status (excellent, very good, good, and fair or poor), and physical activity (metabolic equivalent hours in quartiles). The statistical significance of a linear trend was tested by including the median of each category as a continuous variable in the regression model and interactions by including a multiplicative term between supplement use frequency and the stratifying variable.

To further explore the relation between multivitamin use and telomere length, we conducted stratified analyses according to median age (median: $<$ and ≥ 53 y), smoking status (never and ever), BMI ($<$ and ≥ 30), and the presence of diabetes or cardiovascular diseases (yes and no). In addition, we conducted a sensitivity analysis by excluding women who reported “fair or poor” health status.

Finally, we examined whether intakes of any important micronutrients that were commonly found in multivitamins were related to telomere length in the study population. Among multivitamin users, a substantial proportion of the micronutrient intake was from multivitamins. This was particularly true for vitamins C, D, and E and most B vitamins. We therefore fit the regression models with and without adjusting for multivitamin use and conducted an analysis among women who did not take multivitamins to evaluate the independent relation between dietary micronutrient intake and telomere length. All statistical analyses were conducted by using SAS software (version 9.1; SAS Institute, Cary, NC), and the significance tests were 2-tailed with $\alpha = 0.05$.

RESULTS

Study sample characteristics are presented in **Table 1**. Compared with women who did not take multivitamins, regular users were older, more likely to be non-Hispanic white and never smokers, and had higher education level. However, multivitamin users and nonusers were not significantly based on other population characteristics.

Sixty-five percent of the women used multivitamins at least once per month, and most users (74%) took multivitamins on a daily basis. About 89% of the users took once-a-day type multivitamins, 21% took antioxidant combination, and 17% took stress-tabs or B-complex vitamins. Among users, multivitamins represented a major source of total vitamin and mineral intakes, contributing $>50\%$ of the total intake for vitamins C, E, D, B-6, B-12, folate, iron, and zinc and 30–50% for vitamin A, β -carotene, and calcium.

In general, the use of multivitamin supplements was associated with longer telomere length (**Figure 1**). Compared with nonusers, daily users had on average 5.1% longer telomeres (P for trend = 0.002). This difference (273 bp) corresponds to ≈ 9.8 y of age-related telomere loss since each year of age was associated with a 28-bp shorter telomere in our sample. Significant



TABLE 1
Characteristics of the study participants¹

	Multivitamin supplement ²			P value ³
	All (n = 586)	Nonusers (n = 203)	Users (n = 378)	
Age (y)	53.6 ± 9.6 ⁴	51.6 ± 9.1	54.6 ± 9.7	0.0005
Non-Hispanic whites [n(%)]	493 (84.1)	162 (79.8)	327 (86.5)	0.03
Education [n(%)]				0.005
≤High school	90 (15.4)	40 (19.7)	49 (13.0)	
Some college	152 (25.9)	63 (31.0)	88 (23.3)	
Associate degrees/technical training	84 (14.3)	31 (15.3)	53 (14.0)	
College graduate	148 (25.3)	37 (18.2)	110 (29.1)	
Graduate degree	112 (19.1)	32 (15.8)	78 (20.6)	
Smoking [n(%)]				0.009
Former	173 (29.5)	54 (26.6)	116 (30.7)	
Current	136 (23.2)	62 (30.5)	73 (19.3)	
BMI (kg/m ²)	27.5 ± 6.2	28.2 ± 6.6	27.2 ± 5.9	0.09
Physical activity (MET hours)	47.2 ± 30.0	44.2 ± 27.4	48.6 ± 31.0	0.07
Self-reported health [n(%)]				0.5
Excellent	185 (31.6)	62 (30.5)	121 (32.0)	
Very good	211 (36.0)	67 (33.0)	142 (37.6)	
Good	145 (24.7)	56 (27.6)	88 (23.3)	
Fair or poor	45 (7.7)	18 (8.9)	27 (7.1)	
Perceived stress level [n(%)]				0.08
Very low	109 (18.6)	35 (17.2)	74 (19.6)	
Low	144 (24.6)	43 (21.2)	99 (26.2)	
Moderate	108 (18.4)	44 (21.7)	63 (16.7)	
High	128 (21.8)	39 (19.2)	89 (23.5)	
Very high	97 (16.6)	42 (20.7)	53 (14.0)	
Self-reported diabetes or cardiovascular diseases [n(%)]	112 (19.1)	37 (18.2)	74 (19.6)	0.7
Total energy (kcal)	1590 ± 535	1590 ± 597	1594 ± 501	0.9

¹ MET, metabolic equivalent task.

² Five women were missing data on multivitamin use.

³ A Student's *t* test was used for continuous variables, and a chi-square test was used for categorical variables.

⁴ Mean ± SD (all such values).

associations were also obtained for the once-a-day or the antioxidant combination type, but not for the stress-tab or B-complex type. Excluding women who reported fair or poor health did not change the results: the relative telomere length was 5398 bp for nonusers and 5645 bp for daily users (4.6% difference; *P* for trend = 0.009). Analysis of the duration of individual multi-

tamin use showed similar results. Compared with nonusers, the adjusted telomere length of those who took multivitamins for >5 y was ≈3% longer for once-a-day type multivitamins (*P* for trend = 0.09) and 8% for antioxidant combination type (*P* for trend = 0.02). The duration of stress-tabs or B complex use was not related to telomere length. Multivitamin use was also

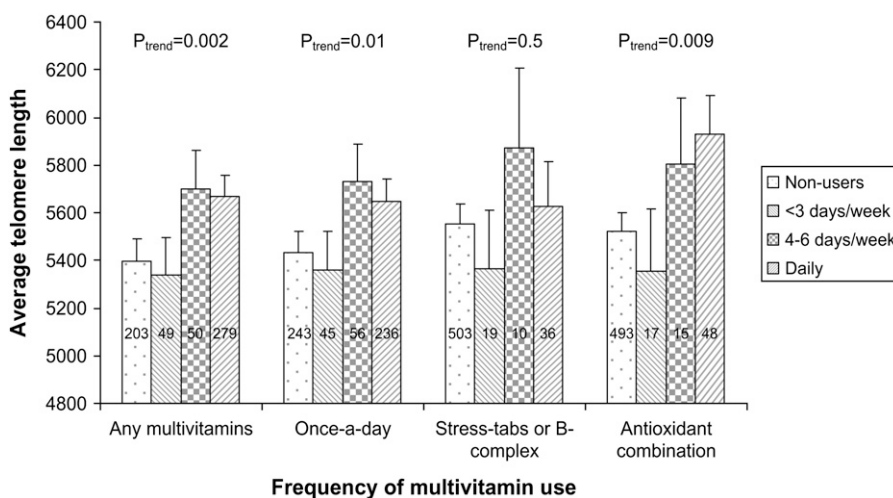


FIGURE 1. Least-squares mean (±SE) telomere length according to the frequency of multivitamin use. Generalized linear models were used in the analysis, adjusted for age, race, BMI, education, cigarette smoking, presence of diabetes or cardiovascular diseases, energy intake, perceived stress level, self-reported health status, and physical activity. Numbers within the bars represents the sample size for each exposure group.

associated with longer telomere length in most of the subgroup analyses by age, sex, and smoking status (Table 2), although not all associations were significant. Use of individual micronutrient supplements was less common in this study sample, and, in general, they were not associated with telomere length after multivitamin use was accounted for (data not shown). The only exceptions were vitamin B-12 and iron: vitamin B-12 supplement users ($n = 52$) had a longer telomere length than did nonusers ($n = 518$): 5850 ± 159 compared with 5505 ± 89 bp (5.9% difference; $P = 0.03$), and iron users ($n = 41$) had a shorter telomere length than nonusers ($n = 527$): 5121 ± 183 compared with 5583 ± 87 bp (-9.0% difference; $P = 0.007$).

The total intake of most micronutrients was positively associated with telomere length (Table 3); however, these associations became statistically nonsignificant after multivitamin use was adjusted for. Micronutrient intake from foods was generally not related to telomere length, except for vitamins C and E (Table 4). Higher dietary intake of these 2 antioxidants was associated with longer telomere length in a dose-response manner even after multivitamin use was adjusted for. Among women who did not use multivitamins ($n = 203$), higher dietary intakes of β -carotene, folate, magnesium, and vitamins C, E, and A were each associated with longer telomere length (Table 4).

TABLE 2
Average telomere length according to frequency of multivitamin use in subgroups¹

	Users				<i>P</i> for trend ²	<i>P</i> for interaction ³
	Nonusers	<3 d/wk	4–6 d/wk	Daily		
Age						0.6
<53 y						
No. of subjects	122	27	35	99		
Lsmean \pm SE	5586 ± 141	5429 ± 233	5893 ± 213	5824 ± 146	0.05	
≥ 53 y						
No. of subjects	81	22	15	180		
Lsmean \pm SE	5168 ± 138	5291 ± 237	5373 ± 286	5494 ± 126	0.02	
BMI						0.5
<30 kg/m ²						
No. of subjects	137	34	37	190		
Lsmean \pm SE	5434 ± 126	5483 ± 194	5705 ± 197	5624 ± 122	0.08	
≥ 30 kg/m ²						
No. of subjects	66	15	13	89		
Lsmean \pm SE	5230 ± 162	5009 ± 314	5618 ± 336	5708 ± 158	0.008	
Smoking						0.6
Never						
No. of subjects	87	24	26	139		
Lsmean \pm SE	5483 ± 139	5307 ± 234	5717 ± 233	5628 ± 123	0.2	
Ever						
No. of subjects	116	25	24	140		
Lsmean \pm SE	5368 ± 131	5450 ± 230	5765 ± 230	5788 ± 135	0.002	
Diabetes or cardiovascular disease						0.4
No						
No. of subjects	166	39	47	218		
Lsmean \pm SE	5587 ± 104	5508 ± 180	5817 ± 168	5796 ± 103	0.03	
Yes						
No. of subjects	37	10	3	61		
Lsmean \pm SE	4908 ± 219	4966 ± 390	5813 ± 667	5590 ± 187	0.004	

¹ Least-squares mean (Lsmean) \pm SE values were derived from generalized linear regression models, adjusted for age, race, BMI, education level, cigarette smoking, presence of diabetes or cardiovascular diseases, energy intake, perceived stress level, self-reported health status, and physical activity. Stratified variables were also adjusted for in the subgroup analysis when possible.

² Tested by including the median of each category as a continuous variable in the regression model.

³ Tested by including a multiplicative term in the regression model.

DISCUSSION

In this cross-sectional analysis, multivitamin use was related to longer telomere length in women aged 35–74 y. Nutrient analysis suggests that one or more dietary antioxidant vitamins may contribute to this relation.

Telomeres typically shorten by a few dozen to a couple hundred bps per cell division (23); therefore, telomere length has been proposed as a marker of biological aging. Furthermore, because telomere attrition may eventually lead to chromosomal instability and cell death, excessive telomere shortening may play an important role in the development of some chronic diseases (1, 23). In recent epidemiologic studies, shorter leukocyte telomeres have been linked to higher mortality (3–5), accelerated aging (1), and higher risk of a variety of chronic diseases (4, 5, 24).

Telomere attrition in human somatic cells is likely the result of multiple forces, including the “end-replication” problem and low telomerase activity (2). Compared with these factors, oxidative stress is probably a more important contributor to telomere attrition (2). Telomeres are particularly vulnerable to oxidative damages, which often cannot be efficiently repaired (23). Furthermore, inflammatory reactions induce oxidative stress, and tumor necrosis factor- α significantly decreases telomerase activity and reduces telomere length in leukemic cells (25).

TABLE 3

Average telomere length according to total intake of selected micronutrients with and without adjustment for multivitamin use¹

Nutrient	Model 1 ² (n = 586)			Model 2 ³ (n = 581)		
	Quartile 1 ⁴	Quartile 4 ⁴	P for trend ⁵	Quartile 1 ⁴	Quartile 4 ⁴	P for trend ⁵
Vitamin A (IU)	4953 ⁶	17,547		4927	17,547	
Lsmean ± SE	5293 ± 106	5633 ± 108	0.02	5356 ± 124	5571 ± 124	0.3
β-carotene (μg)	1621	7226		1613	7206	
Lsmean ± SE	5336 ± 107	5649 ± 106	0.03	5409 ± 120	5605 ± 120	0.2
Vitamin C (mg)	63	785		63	794	
Lsmean ± SE	5289 ± 107	5670 ± 106	0.02	5318 ± 132	5620 ± 121	0.2
Vitamin E (α-TE)	7.6	328		7.6	328	
Lsmean ± SE	5324 ± 104	5640 ± 111	0.1	5346 ± 158	5590 ± 127	0.5
Vitamin B-6 (mg)	1.2	4.9		1.2	4.9	
Lsmean ± SE	5424 ± 107	5737 ± 107	0.005	5604 ± 149	5619 ± 140	0.5
Vitamin B-12 (μg)	2.3	17		2.3	17	
Lsmean ± SE	5355 ± 104	5737 ± 106	0.002	5442 ± 142	5641 ± 127	0.1
Folate (μg)	236	867		236	867	
Lsmean ± SE	5336 ± 108	5689 ± 107	0.01	5452 ± 146	5516 ± 146	0.8
Vitamin D (IU)	75	639		75	637	
Lsmean ± SE	5393 ± 103	5564 ± 108	0.009	5430 ± 134	5500 ± 142	0.7
Calcium (mg)	444	1906		445	1906	
Lsmean ± SE	5407 ± 106	5578 ± 111	0.04	5454 ± 120	5501 ± 126	0.4
Selenium (μg)	61	113		61	113	
Lsmean ± SE	5369 ± 103	5663 ± 106	0.02	5446 ± 115	5602 ± 123	0.3
Iron (mg)	8.5	33		8.5	33	
Lsmean ± SE	5391 ± 107	5568 ± 110	0.03	5459 ± 131	5459 ± 135	0.97
Magnesium (mg)	184	419		184	419	
Lsmean ± SE	5314 ± 105	5659 ± 109	0.007	5364 ± 126	5589 ± 129	0.3
Zinc (mg)	7.1	31		7.1	30	
Lsmean ± SE	5332 ± 102	5630 ± 106	0.008	5350 ± 134	5560 ± 132	0.4

¹ Least-squares mean (Lsmean) ± SE values were derived from generalized linear regression models. TE, tocopherol equivalents.

² Adjusted for age, race, BMI, education level, cigarette smoking, presence of diabetes or cardiovascular diseases, energy intake, perceived stress level, self-reported health status, and physical activity.

³ Adjusted as for model 1 and for multivitamin use. Five women with missing data on multivitamin use were not included in model 2.

⁴ Each quartile represents ≈25% of the study participants: quartile 1, the lowest 25%, and quartile 4, the highest 25%.

⁵ Tested by including the median of each category as a continuous variable in the regression model.

⁶ Median (all such values).

Therefore, oxidative stress and chronic inflammation may be among the major mechanisms of telomere attrition. On the other hand, many micronutrients, such as dietary antioxidants, B-vitamins, and certain minerals, can modulate oxidative stress and inflammatory reactions (26–28) and therefore can contribute to the maintenance or attrition of telomeres. Few studies to date have investigated the role of these micronutrients in telomere maintenance. Earlier *in vitro* experiments showed that ascorbic acid or its derivatives (12, 29, 30) or α-tocopherol (13) slowed telomere shortening and increased the life span of certain somatic cells. In rats, iron overload significantly increased telomerase activity in liver cells but caused no change in telomere length (31). Recently, 2 population-based cross-sectional analyses examined dietary biomarkers in relation to telomere length (14, 15). In the first study, higher plasma vitamin D was associated with longer leukocyte telomere length among women, probably via antiinflammatory actions of vitamin D (14). In the other study, higher plasma homocysteine was associated with shorter telomere length, whereas higher folate was related to longer telomeres (15).

Containing most key vitamins and minerals 100% of the recommended daily intake, multivitamins are major sources of micronutrients in the US diet. According to the recent National Health and Nutrition Examination Survey, 35% of US adults took one or more types of multivitamins, the majority of whom were elderly white women (16). To our knowledge, this was the first epidemiologic study of multivitamin use and telomere length. Regular multivitamin users tend to follow a healthy lifestyle and have a higher intake of micronutrients, which sometimes makes it difficult to interpret epidemiologic observations on multivitamin use (16). In this study, we took extra caution in the data analyses by adjusting for and stratifying by important factors that may affect telomere length, including age, smoking status, and BMI (32, 33). Furthermore, we controlled for several indicators of socioeconomic status or lifestyle choice in all analyses. Women with less optimal health or chronic diseases may be more likely to use vitamin supplements; however, the exclusion of these women from the analysis did not alter the results. Previous epidemiologic results on multivitamin use and risk of chronic diseases vary, depending on the nutrient

TABLE 4
Average telomere length according to intake of selected micronutrients from foods¹

	All women ² (n = 581)			Multivitamin nonusers ³ (n = 203)		
	Quartile 1 ⁴	Quartile 4 ⁴	P for trend ⁵	Quartile 1 ⁴	Quartile 4 ⁴	P for trend ⁵
Vitamin A (IU)	3845 ⁶	13,160		3508	11,273	
Lsmean ± SE	5399 ± 115	5527 ± 117	0.3	5230 ± 171	5674 ± 167	0.008
β-carotene (μg)	1168	5323		1052	4708	
Lsmean ± SE	5463 ± 115	5549 ± 117	0.4	5120 ± 174	5484 ± 164	0.045
Vitamin C (mg)	42	153		38	134	
Lsmean ± SE	5340 ± 115	5683 ± 113	0.03	4945 ± 173	5701 ± 153	0.002
Vitamin E (α-TE)	6.1	12		5.7	12	
Lsmean ± SE	5311 ± 115	5672 ± 118	0.004	5134 ± 161	5533 ± 173	0.03
Vitamin B-6 (mg)	1.0	1.8		0.9	1.7	
Lsmean ± SE	5434 ± 112	5625 ± 117	0.3	5308 ± 159	5300 ± 164	0.8
Vitamin B-12 (μg)	1.8	4.6		1.7	4.6	
Lsmean ± SE	5574 ± 108	5611 ± 116	0.8	5317 ± 159	5414 ± 171	0.6
Folate (μg)	201	401		181	388	
Lsmean ± SE	5371 ± 115	5530 ± 117	0.4	5091 ± 175	5586 ± 162	0.02
Vitamin D (IU)	50	250		42	215	
Lsmean ± SE	5606 ± 110	5591 ± 114	0.8	5389 ± 157	5537 ± 164	0.3
Calcium (mg)	366	931		331	824	
Lsmean ± SE	5549 ± 110	5520 ± 116	0.9	5187 ± 165	5583 ± 163	0.06
Selenium (μg)	54	92		51	90	
Lsmean ± SE	5518 ± 110	5574 ± 116	0.8	5264 ± 163	5563 ± 163	0.1
Iron (mg)	7.6	14		6.9	14	
Lsmean ± SE	5511 ± 111	5605 ± 119	0.7	5365 ± 176	5367 ± 162	0.9
Magnesium (mg)	165	304		154	273	
Lsmean ± SE	5382 ± 113	5572 ± 118	0.1	5183 ± 167	5603 ± 172	0.04
Zinc (mg)	6.4	12		5.7	12	
Lsmean ± SE	5504 ± 114	5576 ± 116	0.6	5292 ± 162	5480 ± 168	0.2

¹ Least-squares mean (Lsmean) ± SE values were derived from generalized linear regression models. TE, tocopherol equivalents.

² The analysis among all women was adjusted for age, race, BMI, education level, cigarette smoking, presence of diabetes or cardiovascular diseases, energy intake, perceived stress level, self-reported health status, physical activity, and multivitamin use.

³ The analysis among multivitamin nonusers was adjusted as for all women, except for multivitamin use.

⁴ Each quartile represents ≈25% of the study participants: quartile 1, the lowest 25%, and quartile 4, the highest 25%.

⁵ Tested by including the median of each category as a continuous variable in the regression model.

⁶ Median (all such values).

composition, the disease type, and the design of the study (34–36). Further investigations would be needed to understand the role of multivitamin use and telomere length and its implication in the etiology of chronic diseases.

Sorting out which micronutrients underlie our findings is difficult because multivitamins contain various vitamins and minerals and contribute in large amounts to daily micronutrient intakes. Nevertheless, higher intakes of the antioxidant vitamins C and E consistently showed associations with longer telomeres in different analyses. In multivitamin users, ≈63% of vitamin C and 84% of vitamin E were from supplemental sources. Whereas the evidence is not sufficient to conclude that these 2 dietary antioxidants mediated the observed relation, the results are consistent with experimental findings that vitamins C and E protect telomeres *in vitro* (12, 13, 29, 30).

This study had several limitations. The quantitative PCR-based assay measures the average telomere length across all leukocytes in the peripheral blood. We therefore could not exclude the possibility that multivitamin use might have shifted the composition of leukocyte subpopulations in a way that favored cells with longer telomeres as an alternative explanation for our

finding. Furthermore, as the first study on this topic, our analysis was explanatory in nature; this was particularly true for subgroup analyses that included a smaller number of participants. Finally, although it is unlikely that telomere length affects multivitamin use, this analysis was cross-sectional and we were unable to make a direct causal inference. It is advisable to follow-up these findings in future large longitudinal studies.

Compared with national data, more women in our population took multivitamins (16). This may be a characteristic of women who volunteer for cohort studies and itself does not necessarily affect the validity of our study. Residual confounding is always a concern in epidemiologic research on health behaviors, such as multivitamin use. In this study, we adjusted for and stratified by a variety of potential confounders and conducted a sensitivity analysis by excluding women with fair or poor health. Finally, our dietary data relied on an FFQ, which was subject to measurement errors. However, the Block FFQ is widely used for dietary surveys and has been consistently updated and validated in various populations (20, 21, 37).

In summary, our study provides preliminary evidence linking multivitamin use to longer leukocyte telomeres. This finding



should be further evaluated in future epidemiologic studies and its implications concerning aging and the etiology of chronic diseases should be carefully evaluated.

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