Cellular Aging and Restorative Processes: Subjective Sleep Quality and Duration Moderate the Association between Age and Telomere Length in a Sample of Middle-Aged and Older Adults

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Study Objectives: To examine whether subjective sleep quality and sleep duration moderate the association between age and telomere length (TL).

Design: Participants completed a demographic and sleep quality questionnaire, followed by a blood draw.

Setting: Social Neuroscience Laboratory.

Participants: One hundred fifty-four middle-aged to older adults (age 45-77 y) participated. Participants were excluded if they were on immunosuppressive treatment and/or had a disease with a clear immunologic (e.g., cancer) component.

Interventions: N/A.

Measurements and Results: Subjective sleep quality and sleep duration were assessed using the Pittsburgh Sleep Quality Index (PSQI) and TL was determined using peripheral blood mononuclear cells (PBMCs). There was a significant first-order negative association between age and TL. Age was also negatively associated with the self-reported sleep quality item and sleep duration component of the PSQI. A significant age × self-reported sleep quality interaction revealed that age was more strongly related to TL among poor sleepers, and that good sleep quality attenuated the association between age and TL. Moreover, adequate subjective sleep duration among older adults (i.e. greater than 7 h per night) was associated with TL comparable to that in middle-aged adults, whereas sleep duration was unrelated to TL for the middle-aged adults in our study.

Conclusions: The current study provides evidence for an association between sleep quality, sleep duration, and cellular aging. Among older adults, better subjective sleep quality was associated with the extent of cellular aging, suggesting that sleep duration and sleep quality may be added to a growing list of modifiable behaviors associated with the adverse effects of aging.

Keywords: Aging, sleep quality, telomeres

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INTRODUCTION

Sleep complaints constitute one of the most common difficulties facing middle-aged and older adults.1 Indeed, epidemiological studies suggest that most community-dwelling older adults have some concerns about their ability to fall asleep or stay asleep, with fewer than 20% reporting “rarely” or “never” having any sleep complaints.2 Older adults often take longer to fall asleep, wake more frequently during the night, and experience lower sleep efficiency than their younger counterparts.3 Importantly, these objective age-related changes in sleep across the life span are often associated with self-reports of poor sleep quality.4,6

One pathway through which both objective and subjective measures of sleep may be associated with health is inflammation. Sleep is closely linked to immune system changes via the sleep-wake cycle, with immune cells at their highest blood levels in the early evening and lowest in the morning,7 and disruption of the regular sleep-wake cycle appears to significantly affect immune function.8,9 Importantly, sleep deprivation has been found to induce increases in circulating levels of inflammatory markers, such as tumor necrosis factor-α (TNF-α),10 C-reactive protein (CRP),11 and interleukin-6 (IL-6),12 with significant elevations occurring after only 1 night of sleep loss.11 Additionally, experimentally induced sleep deprivation in the laboratory has also been associated with nuclear factor kappa-B cell (NF-kB) activation, a molecular pathway through which sleep disturbance may influence leukocyte inflammatory gene expression and inflammation-related disease risk.13 This body of literature demonstrates effects of sleep loss on a range of well-known markers of the acute inflammatory system, independent of actual injury or infection to the host. The combination of high prevalence of sleep problems among older adults, as well as the effects of sleep loss on inflammatory processes, may put older adults at increased risk for many chronic conditions, because aging alone is associated with low-grade elevation of inflammatory markers.14

In addition to increased inflammation during chronological aging, inflammation also appears to play a role in cellular aging. Immune system cells are under tremendous proliferative demand, requiring telomeres to remain intact. Telomeres are repetitive structures at the end of chromosomes that help to promote cellular stability.15,16 However, with each successive replication of the cell, telomeres shorten and when a critical threshold is met, the result is cellular senescence. In all mitotic tissues aside from germline tissue, telomere length (TL)
declines with age. Telomere activity serves several critical genomic purposes, including the prevention of chromosomal fusions and unregulated cellular activity. Lymphocytes, an immune cell, are capable of upregulating the enzyme (telomerase) that elongates telomeres, prolonging cellular life span.

In addition, recent work demonstrates telomere shortening in monocytes is associated with increased synthesis of proinflammatory cytokines. Although the set point of TL appears to be genetically determined, it is not yet fully understood what drives age-associated telomere loss, although several factors have been identified (e.g., chronic stress and proliferation-induced shortening).

A small but growing body of literature now demonstrates associations between sleep and TL. Of these studies, three provide evidence that self-reported sleep quality and sleep duration are related to TL. For example, Prather and colleagues found an association between self-reported sleep quality among middle-aged women and TL, such that women who reported poorer sleep quality had shorter leukocyte TL. This finding has recently been replicated in healthy men (M_age = 63.3, standard deviation = 5.6). In addition, Liang and colleagues found that long sleep duration was significantly associated with long TL among women younger than 50 years, but not among women older than 50 years. These findings raise the question of whether associations between sleep quality, sleep duration, and TL change as we age. That is, as we get older, do sleep duration and self-reported sleep quality differentially predict TL compared to a relatively younger population? These prior studies suggest that further study examining associations between sleep duration, self-reported sleep quality, and TL is warranted.

The purpose of the current study was to examine associations among age, subjective sleep quality, sleep duration, and telomere length in a sample of middle-aged and older adults. The first aim was to replicate prior findings of an association between age and TL. A second aim was to replicate recent findings demonstrating associations between subjective sleep quality and self-reported sleep duration with TL, with the prediction that poorer sleep quality and shorter sleep duration would be associated with shorter TL. Additionally, prior research suggests a U-shaped association between sleep duration and negative health effects. A third aim of the current study was to examine a nonlinear model. Chronological age is generally associated with poorer sleep quality as well as cellular aging, and poor sleep quality is linked to shorter TL. Thus, it was predicted that increasing age would be associated with shorter TL among those reporting shorter sleep duration and poorer sleep quality.

**METHOD**

Participants

As part of a larger study, 154 relatively healthy participants (89 men and 65 women) were included in this study. An additional 23 participants were studied, but for various reasons had no telomere data (e.g., blood draw or blood separation problems). Importantly, comparing these individuals with participants who had complete data on age—the only demographic factor related to telomere length—revealed no significant group differences (mean age = 60.1 versus 60.6, P > 0.70). We recruited a middle-aged to older adult sample (45 to 77 y [M_age = 60.1, standard deviation = 6.76]) to allow for greater variability in TL. Because many older adults are on some form of medication, we only excluded individuals who were on immunosuppressive treatment (e.g., corticosteroid therapy) and/or had current cancer or human immunodeficiency virus to address concerns about potential effects on peripheral blood mononuclear cells (PBMCs), which we used for the determination of TL. Corticosteroid therapy and treatment regimen are factors known to influence immune cell function. Because there was little evidence on the extent of such influences on TL determined from these cells at the time of the study, we conservatively used them as exclusion criteria.

Procedure

The current investigation was performed in accordance with the ethical standards of the University of Utah institutional review board. All eligible participants were scheduled for an appointment at the Social Neuroscience Laboratory, and following informed consent, were rechecked against the exclusion criteria. Participants then completed a demographic questionnaire (e.g., age, sex) and the Pittsburgh Sleep Quality Index (PSQI). Following completion of the questionnaires, approximately 20 cc of blood were drawn and treated with EDTA to prevent clotting. Telomere length was determined using PBMCs. DNA from the PBMCs was extracted from fresh cells, which were separated using density gradient centrifugation with Ficoll-Hypaque (GE Healthcare). Following the blood draw, participants were debriefed and received $75.00 for their participation.

**Measures**

**Pittsburgh Sleep Quality Index**

The PSQI assesses sleep habits during the previous month. The scale is composed of 19 items, which are used to derive seven component scores: sleep quality, sleep latency, sleep duration, habitual sleep efficiency, sleep disturbances, sleep medication, and daytime dysfunction. Component scores on the PSQI are summed to produce a global measure of subjective sleep quality score, with higher scores indicating poorer sleep quality. This instrument has demonstrated good reliability (alpha = 0.83) and validity. For the current study, the sleep duration component score and the item assessing sleep quality were used.

**Telomere Length**

DNA was extracted from isolated PBMCs using the reagents and protocols contained in Qiagen’s Gentra Puregene Blood Kit, Catalog # 158389, with the exception that the red blood cell lysis step was skipped, as it was unnecessary due to prior removal of RBCs by centrifugation. Relative TLs were then determined for the PBMCs using quantitative polymerase chain reaction (PCR). For each DNA sample, we measured the factor by which the sample differed from a reference DNA sample in its ratio of telomere repeat copy number to single copy gene copy number. This ratio is proportional to the average TL (T/S ratio). These assays were run in several batches, and each
batch consisted of DNA samples from 96 participants per quantitative PCR run. The interassay geometric mean of the coefficient of variation for this assay was 3.13%, based on examining the variation in the mean T/S ratio across days. The intra-assay geometric mean of the coefficient of variation was 5.22%. This was determined by examining variations in the T/S of samples across triplicate repeat measurements of the samples, collected in the same run. Outliers were determined within triplicate batch consisted of DNA samples from 96 participants per quantitative PCR run. The interassay geometric mean of the coefficient of variation for this assay was 3.13%, based on examining the variation in the mean T/S ratio across days. The intra-assay geometric mean of the coefficient of variation was 5.22%. This was determined by examining variations in the T/S of samples across triplicate repeat measurements of the samples, collected in the same run. Outliers were determined within triplicate measurements of T/S for a given sample: when PCR amplification of either the telomere target and/or the single copy gene target failed for one of the three replicates, that replicate was removed from the analysis. When amplification of two or more of the three replicates of a sample failed, the assay was repeated in triplicate for that DNA sample. The Shapiro-Wilk test of normality, which is appropriate for this sample size, was 0.98, indicating a high degree of normality in the resulting T/S ratio data.

### Overview of Analyses

To characterize the sample with respect to age, sleep duration, self-reported sleep quality, and telomere length, means, standard deviations and zero-order correlations among these variables were calculated.

Separate regression analyses examined independent associations among age, sleep duration (PSQI component score), subjective sleep quality (PSQI item), and TL. All models included covariates known to be associated with telomere length (sex; body mass index; average weekly alcohol consumption; average weekly tobacco use; average weekly exercise; use of hormone replacement therapy; and use of hypertension, cholesterol, and diabetes medications). All variables were centered to minimize multicollinearity. In separate regression models that included the PSQI sleep duration component and the PSQI sleep quality item scores age remained a significant predictor of TL (P < 0.001), above and beyond the following covariates: sex, body mass index, average weekly alcohol consumption, average weekly tobacco use, average weekly exercise, use of hormone replacement therapy, use of hypertension, cholesterol, diabetes medications.

To examine the moderating effect of sleep duration and sleep quality on the association between age and TL, separate regression models that included centered first-order effects and the cross product terms (age \times sleep quality item; age \times sleep duration component score) were conducted. One significant interaction emerged. There was a significant interaction between age and self-reported sleep quality (b = -0.011, t = -2.42, P < 0.05, ΔR² = 0.04) above and beyond sex, body mass index, average weekly alcohol consumption, average weekly tobacco use, average weekly exercise, use of hormone replacement therapy, and use of medications for hypertension, cholesterol, and diabetes. No other main effects or interactions of these PSQI scores emerged. In order to probe the significant age \times self-reported sleep quality interaction, the regression model was restructured on high and low values (one standard deviation above and below the mean) of sleep duration and sleep quality, respectively.

Finally, because prior research suggests a U-shaped association between sleep duration and negative health effects, a nonlinear model was also examined. Significant interactions were probed by restructuring the regression model one standard deviation above and below the means of age and sleep duration.

### RESULTS

#### Primary Analyses

Means, standard deviations, and correlations among the study variables are presented in Table 1. Age was significantly inversely correlated with the sleep duration component of the PSQI and the PSQI item about self-reported sleep quality. The well-known association between age and TL was replicated (b = -0.011, t = -3.83, P = 0.0002).

Because prior research on sleep in older adults suggests that sleep duration and sleep quality are important for health, we examined the PSQI sleep duration component and self-reported sleep quality item. In all regression models, variables were centered to minimize multicollinearity. In separate regression models that included the PSQI sleep duration component and the PSQI sleep quality item scores age remained a significant predictor of TL (P < 0.001), above and beyond the following covariates: sex, body mass index, average weekly alcohol consumption, average weekly tobacco use, average weekly exercise, use of hormone replacement therapy, use of hypertension, cholesterol, diabetes medications.

To examine the moderating effect of sleep duration and sleep quality on the association between age and TL, separate regression models that included centered first-order effects and the cross product terms (age \times sleep quality item; age \times sleep duration component score) were conducted. One significant interaction emerged. There was a significant interaction between age and self-reported sleep quality (b = -0.011, t = -2.42, P < 0.05, ΔR² = 0.04) above and beyond sex, body mass index, average weekly alcohol consumption, average weekly tobacco use, average weekly exercise, use of hormone replacement therapy, and use of medications for hypertension, cholesterol, and diabetes. No other main effects or interactions of these PSQI scores emerged. In order to probe the significant age \times self-reported sleep quality interaction, the regression model was restructured on high and low values (one standard deviation above and below the mean) of self-reported sleep quality (item). As displayed in Figure 1 (which displays plotted values one standard deviation above and below the mean of the PSQI item about self-reported sleep quality), follow-up analyses...
revealed that age was more strongly negatively related to TL among those with poor self-reported sleep quality (\(b = -0.021, P \leq 0.0001\)) than those with better self-reported sleep quality (\(b = -0.005, P = 0.24\)), suggesting that sleep quality attenuated the effect of age on TL.

Finally, given prior research suggesting that both high and low sleep duration may have negative health effects (i.e., a curvilinear effect), a nonlinear association between the component score of sleep duration and TL was examined in a regression model that included sex, age, the linear sleep duration term (component score), and a quadratic sleep duration term. There was a significant association between the quadratic sleep duration term and TL (\(b = 0.05, P < 0.05\)) above and beyond sex; body mass index; average weekly alcohol consumption; average weekly tobacco use; average weekly exercise; use of hormone replacement therapy; use of hypertension, cholesterol, and diabetes medications; and chronological age. Moreover, age moderated the association between the quadratic sleep duration term and TL (\(b = 0.013, t = 3.61, P < 0.001, \Delta R^2 = 0.08\)). In order to probe the significant interaction, the regression model was restructured one standard deviation above and below the means for both age and the quadratic sleep duration term. Analysis of simple slopes revealed that sleep duration was unrelated to TL for the middle-aged adults in our sample (\(P > 0.10\)). However, adequate sleep duration (i.e., > 6.9 h/night) in older adults was positively associated with TL (\(b = 0.25, P < 0.01\)). Moreover, it appears that the benefits of adequate sleep duration begin around age 60 y (\(b = 0.06, P < 0.05\)). Although the simple slope for older adults with low sleep duration (< 5 h/night) was also significant (\(b = 0.12, P < 0.01\)), very few older adults reported sleep duration this low (n = 4) and the predicted TL in this group was well below that of the middle-aged participants. Figure 2 shows mean TL values based on the range of values for the sleep duration component on the PSQI and on values one standard deviation above and one standard deviation below the mean of age (\(M_{age} = 60.57\) y). TL for older adults with adequate sleep duration (i.e., > 6.9 h/night) is comparable to that of the middle-aged group. Thus, adequate sleep duration in older adults appears to be associated with attenuated aging, as indicated by TL.

**DISCUSSION**

In this study we examined associations between chronological age, cellular aging, sleep duration, and subjective sleep quality in a sample of middle-aged and older men and women. Consistent with prior research and our first aim, we found an inverse association between age and TL. For our second aim, we did not find the PSQI item of self-reported sleep quality to significantly predict TL on its own. However, it did moderate the association between age and TL, suggesting that among older adults, better sleep quality was significantly associated with longer TL; this pattern was not found for younger adults in this sample. Similar to findings by Jackowska and colleagues, results of the current study indicated a nonlinear association between the component of sleep duration and TL, such that both high and low sleep durations were associated with shorter TL. This nonlinear effect varied with age such that there was evidence of longer TL for older adults with adequate sleep duration; however sleep duration was unrelated to TL for the middle-aged adults in our sample.

The current study replicates and extends recent findings on the links between aging and TL. However, the current study differs from prior studies in important ways. First, the current study is the first to use both men and women in examining the association between PSQI measures of sleep quality and sleep duration with TL. Next, in contrast to findings of Liang and colleagues, results of the current study showed the sleep duration component was significantly associated with TL among older adults compared to middle-aged adults. We interpret current results to suggest that getting an adequate amount of sleep in older adulthood is potentially buffering deleterious effects of TL shortening that normally occur with age; thus, sleep is demonstrated to be a restorative process at the cellular level in older adults. Future research should examine the circumstances that shape how sleep duration influences cellular aging. These general findings are consistent with prior work showing the importance of subjective sleep quality and duration as predictors of TL.

TL, a marker of cellular aging, has been related to other psychosocial factors known to affect health, such as chronic stress, divorce, and childhood trauma. Thus, the results of this investigation may have implications for understanding the mechanisms through which daily metabolic insults such as stress affect health in older adults. Within this general framework, stress exposure, stress reactivity, stress recovery, and restoration are component processes by which stress adversely affects health, and sleep is considered to be a restorative process. Inadequate sleep is associated with adverse health consequences including impaired immune functioning, susceptibility to the common cold, metabolic syndrome, inflammatory processes, coronary artery calcification, coronary heart disease, and type 2 diabetes. Moreover, among middle-aged and older adults, sleeping less than 7 h or more than 8 h is associated with poor health outcomes and increased mortality rates. The data from this study highlight the potential positive effect of better sleep quality and longer sleep duration, especially for older adults.

As shown in Table 1, the older adults in this study reported better sleep quality (PSQI sleep quality item) and slept longer (PSQI sleep duration component) than their middle-aged counterparts. Figure 2—Mean telomere length among middle-aged and older adults by hours of sleep per night (Pittsburgh Sleep Quality Index component sleep duration).

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counterparts. These findings are inconsistent with the view that older adults report more sleep complaints and experience a decline in sleep quality over the life span. However, this sample was composed of relatively healthy middle-aged and older adults, and results of the current study are consistent with prior research indicating that sleep complaints among older adults are not the result of age per se, but are often secondary to physical health problems. Moreover, the results of the current study are consistent with findings from a recent population-based study of subjective sleep quality across the life span showing a decline in sleep quality until age 60 y, followed by a transient increase in sleep quality from age 60 y to 66 y, then a plateau in sleep quality after age 66 y. Although the finding that the older adults in this study reported better sleep quality than their middle-aged counterparts is in contrast to the view that sleep quality declines with age, these results suggest a growing need for a more differentiated description of changes in sleep quality during old age.

There are some limitations to the current study. First, the participants were a relatively healthy community sample of older adults, so generalizations to chronically ill populations and other age groups should be made cautiously. Future research should examine the association between TL and self-reported sleep quality in the context of diagnosed sleep disorders, such as insomnia and obstructive sleep apnea (OSA), especially in light of evidence that OSA in adults has been linked to shorter TL, whereas OSA in children has been associated with longer TL. Thus, more work examining connections between OSA and TL is needed before attempting to understand the role self-reported sleep quality may play in that relationship. Additionally, the data are cross-sectional in nature, limiting the ability to draw causal inference. Moreover, only 2% to 3% of the total variance in TL was explained by interactions between age and the individual item of sleep quality, indicating a fairly modest effect. Longitudinal investigations may increase our understanding of how poor sleep quality affects cellular aging throughout adulthood. Although self-reported sleep quality is important to examine and figures prominently in the development of chronic sleep disturbance, investigating these associations using objective indices of sleep (e.g., actigraphy and polysomnography) will be important. Relating indices of cellular aging to circadian and homeostatic processes may also help to clarify the mechanism through which sleep affects cellular aging.

Despite these limitations, the current study provides evidence for an association between subjective sleep quality, sleep duration, and cellular aging processes among middle-aged and older adults. These data suggest that sleep and subjective sleep quality and sleep duration can be added to a growing list of modifiable health behaviors linked to shorter TL. Thus, interventions aimed at improving modifiable aspects of sleep may help to lessen the adverse effects associated with cellular aging.

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