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RECEIVED 26 March 2025

ACCEPTED 26 May 2025

PUBLISHED 06 June 2025

CITATION

Qian J, Yu X and Zhang Q (2025) Association of sleep duration and sleep quality with cognitive frailty in Chinese older adults. *Front. Public Health* 13:1596965. doi: 10.3389/fpubh.2025.1596965

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Association of sleep duration and sleep quality with cognitive frailty in Chinese older adults

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Objective: To explore the relationship between sleep duration, sleep quality and the occurrence of cognitive frailty in the older adults.

Methods: A total of 9,970 participants were screened in China over the past 9 years. They were divided into cognitive frailty group and non-cognitive frailty groups, and they were evaluated for sleep duration and sleep quality and their relationship with cognitive frailty was analyzed. If interactions are found, further hierarchical analysis is conducted.

Results: One thousand six hundred eighty-four participants (16.89%) were diagnosed with cognitive frailty. Participants with cognitive frailty were more likely to be “unmarried,” live in rural areas, and were female, with no social activity in the last month. Poor sleep quality, short sleep duration, no napping, and excessive napping are at high risk of cognitive frailty. There was a significant interaction between daytime napping and sleep duration and sleep quality. Among participants with good sleep quality, those who took excessive naps had a 123% increased risk of developing cognitive frailty, with an OR of 2.23 (95% CI: 1.72, 2.86). In the subgroup with sleep duration > 9 h, participants who napped excessively had a significantly increased risk of cognitive frailty (OR = 1.62, 95% CI 1.14–2.30, $p < 0.001$).

Conclusion: Chinese older adults with poor sleep quality are at a 67% higher risk of cognitive weakness, and individuals with less than 6 h of sleep are at a 48% higher risk of cognitive weakness; No napping and excessive napping, the risk of cognitive debilitation increased by 23 and 69%, respectively. There is an additive interaction between sleep duration and quality and daytime napping on cognitive frailty in the older adults.

KEYWORDS

sleep duration, sleep quality, napping time, risk factor, cognitive frailty

1 Introduction

Population aging has become a common phenomenon worldwide, and the average life expectancy of Chinese is also increasing year by year (1, 2). Cognitive impairment and frailty have become major threats to healthy ageing and the quality of life of the older adults (3, 4), and there is an interaction between them, which accelerates the decline of physical functions (5). Therefore, the International Academy on Nutrition and Aging (IANA) and the International Association of Gerontology and Geriatrics (IAGG) formally proposed the concept of “cognitive frailty” in 2013. refers to the presence of both physical frailty and cognitive impairment and excludes dementia and other neurodegenerative diseases (6).

Cognitive frailty has been reported to increase the incidence of dementia (7) and is associated with adverse health outcomes such as decreased functioning, disability, poor quality of life (8), and increased mortality (9). It is generally accepted that the prevalence is higher in older age, females, and people with low educational attainment (10–14). Some scholars have also explored the influencing factors in terms of lifestyle behavior, dietary nutrition, and mental health status, but the conclusions of the study mainly rely on cross-sectional surveys (12, 15) (13, 15).

Sleep constitutes a significant portion of our daily routine, however, there has been considerable changes in the average sleep duration of population in recent years (16). About one-third of the adults in the United States reported obtaining less than 7 h of daily sleep (17).

Both short and long sleep durations are associated to increased risk of major health problems, including diabetes, cardiovascular disease, and mortality (18, 19). And there is growing evidence that sleep affects the risk of cardiovascular disease (20).

As older adults age, they may experience decreased nighttime sleep (21), poorer sleep efficiency and continuity (22), more frequent nocturnal awakenings (23), less time spent in slow-wave sleep and REM sleep (24), more fragmented sleep, and a faster sleep-to-wake transition (25), and these changes may also lead to changes in brain function (26). Sleep disturbance leads to the development of cognitive dysfunction (27) and accelerates the loss of nerve cells in the frontal, parietal, and temporal lobes of the brain, affecting synaptic plasticity and neuronal function, leading to cognitive impairment or dementia (28, 29). Studies have shown that changes in sleep parameters are associated with cognitive decline (30), and cross-sectional studies have found that the rate of cognitive decline in patients with sleep disorders is 2–4 times higher than that of those without sleep disorders (31). Therefore, sleep disturbance is seen as a possible potential trigger or biomarker of cognitive alteration, and monitoring sleep quality is promising as a non-invasive means to assess the risk of future Alzheimer's disease (AD) or to monitor the effects of clinical interventions (32).

The concept of sleep deprivation is often involved in studying the link with cognition. In many studies, shortened sleep is considered total sleep deprivation (TSD), a useful indicator to comprehensively investigate how shortened sleep harms cognition (33). As the number of sleep-deprived people continues to grow, it is important to understand how sleep deprivation affects human cognitive function to prevent its negative effects. In particular, people in many fields of work are affected by reduced sleep duration, which puts them at higher risk while working. For example, shortened sleep duration is a significant cause of industrial and transportation-related accidents (34). Therefore, exploring how sleep deprivation affects human cognitive function is essential to prevent its harmful effects. Previous studies have shown that TSD impairs different levels of cognitive function. For example, TSD significantly impairs attentional performance (35–38), and TSD also impairs higher cognitive processes and even social cognitive abilities (39, 40).

To sum up, we can see that sleep and cognitive frailty are closely related. Previous studies have mostly independently studied the relationship between sleep and cognitive function or frailty (41–43), but there is still a gap in the research on sleep and cognitive frailty, and the relationship between sleep duration, sleep quality and cognitive frailty in the older adults is not very clear, which also provides space for this

study to explore. This study aims to explore the relationship between sleep duration, sleep quality and cognitive frailty in the older adults.

2 Materials and methods

2.1 Study design and participants

The data for this study were obtained from the China Health and Elderly Care Longitudinal Survey (CHARLS), which was approved by the Biomedical Ethics Committee of Peking University (IRB00001052-11015) in 2011. The project is a nationally representative cohort study that comprehensively investigates information from multiple perspectives, including basic information, health status and function information, personal income, household income and expenditure. The project was evaluated every 2 years, and the latest round of data was released on 2023-11-16. This study was a nine-year longitudinal study based on 2011 data, with 17,708 participants initially recruited and 9,970 participants included in the analysis. The enrollment criteria were: (1) seniors aged 60 years and above; (2) The data items required by the research subjects are complete and accurate. The exclusion criteria were: (1) No complete information of cognitive function questionnaire and FRAIL scale; (2) Those who did not have successful follow-up. The inclusion flowchart is shown in Figure 1.

2.2 Measurements

2.2.1 Assessment of napping time

Participants were asked to describe their nap habits by answering the following question: “Are you in the habit of taking daytime naps? Those who reported “no” were classified as non-nappers, while others who reported “yes” were classified as nappers. For nappers, napping duration was categorized as ≤ 60 min/day, 61–90 min/day, and > 90 min/day.

2.2.2 Assessment of sleep quality and duration

The sleep duration was derived from self-reports by participants, collected by asking the following question regarding nocturnal sleep duration: “In the previous month, how many hours of actual sleep did you get each night?” Based on previous research (44, 45), night-time sleep duration was divided into short (≤ 6 h), medium (6–9 h), and long (> 9 h).

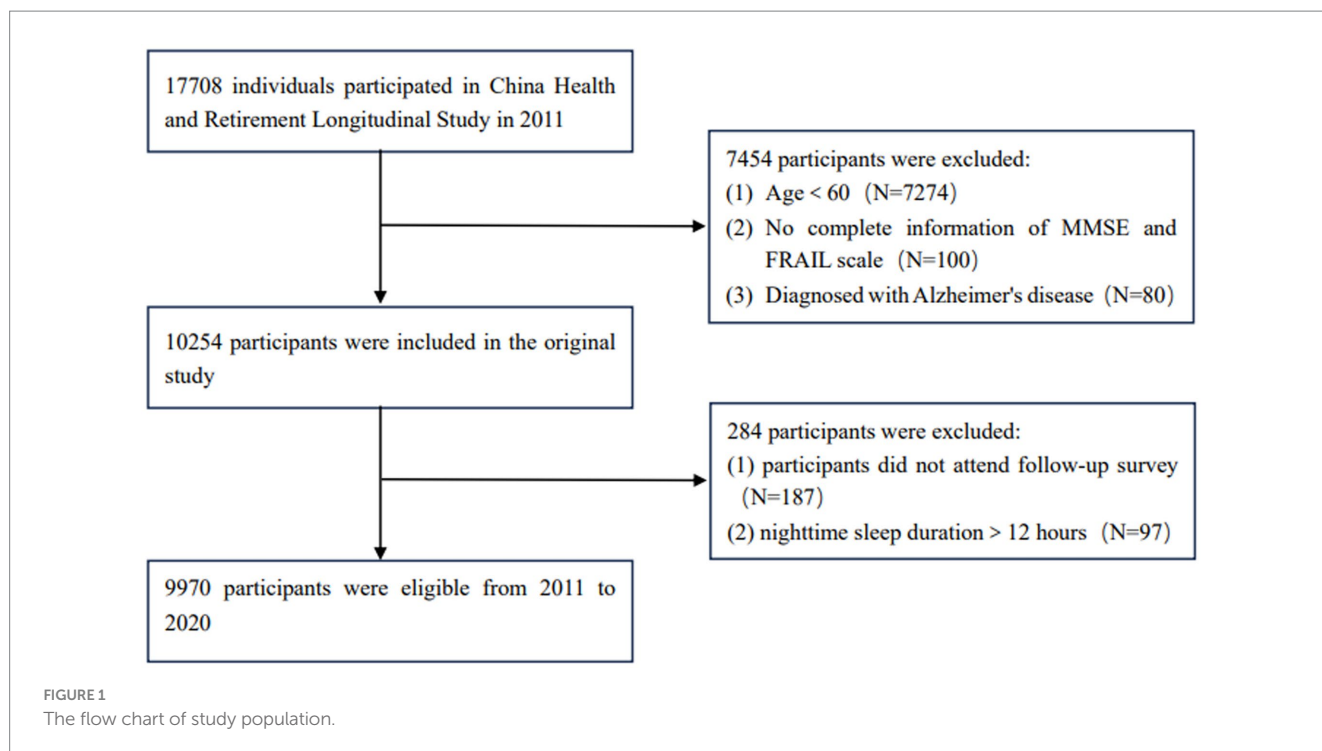
Sleep quality was also self-reported by participants, determined by their response to the question, “How many days of restless sleep did you have in the last week?” The response options encompassed four categories: (1) rarely or none of the time (< 1 d/week); (2) some or a little of the time (1–2 days/week); (3) occasionally or a moderate amount of the time (3–4 days/week); and (4) most or all of the time (5–7 days/week). We classified (1) as good sleep quality, (2) and (3) as moderate sleep quality, and (4) as poor sleep quality drawing upon prior studies (46).

The above sleep questionnaire is shown in Supplementary Table 1.

2.2.3 Assessment of cognitive frailty

2.2.3.1 Physic frail

Physical frailty was assessed using the modified version of the Fried's Frailty Phenotype (47), which has been widely used among



older PWH populations. The tool includes 5 components with self-report and objective measurements: unintentional weight loss, self-reported low physical activity, self-reported exhaustion, weak grip strength, and slow gait speed. Measurement details for each component have been published and described elsewhere. Participants who met 0 components were classified as robust, 1 or 2 components as frailty, and 3 or more components as frailty.

2.2.3.2 Cognitive function

CHARLS's assessment of cognitive function includes self-reported assessment of cognitive decline, telephone interview on cognitive status (TICS-10), word recall, and graphic drawing. The TICS scale measures an individual's cognitive function in two dimensions: memory and mental state (48). Memory is measured by testing the respondent's ability to recall words immediately (0–10 points) and delayed word recall (0–10 points), and respondents are asked to recall and repeat them, and respondents are able to repeat a word for a short period of time or after an interval of 1 point. Mental state is measured from three aspects, including orientation, visual construction, and mathematical performance. The cognitive function score is calculated by summing the scores of the above questions, with a total score ranging from 0 to 31, with higher scores indicating better cognitive function and more complete cognitive ability (49). The cutoff value was set at 18 for illiterate individuals, 21 for individuals with 1–6 years of education, and 25 for individuals with seven or more years of education, and cognitive impairment was defined as those whose score was lower than the cutoff value according to their education level.

2.2.3.3 Cognitive frailty

In line with the definition by (I. A. N. A/I. A. G. G) international consensus group. I.e. as mentioned above, cognitive frailty was

defined as the simultaneous present of both cognitive impairment and physical frailty, which has been validated in previous studies (50, 51).

2.2.4 Assessment of covariates

According to the previous literature (49, 52–57), potential confounding factors were included as covariates in this study, mainly including age, sex (male and female), level of education (elementary school, junior high school, high school, university), marital status (married and unmarried), and place of residence (rural and urban), smoking and drinking, all of which can be obtained from the Demographic Background section of CHARLS. Body mass index (BMI) is calculated by dividing your weight by the square of your height. Chronic disease refers to a person with any of the 14 diseases in the DA007 answers in the CHARLS Household Questionnaire.

2.3 Statistical analysis

Continuous variables were expressed as mean \pm standard deviation, while categorical variables were described using rates and proportions. The Kruskal–Wallis H test was applied to continuous variables, and chi-squared tests were used for categorical variables. We employed multivariate logistic regression analysis to explore the relationship between sleep and cognitive frailty. Three models were constructed for this purpose by adjusting for potential confounding factors. The results are presented as odds ratios (OR) with 95% confidence intervals (95% CI). Subgroup analyses stratified by interaction analyses were performed. Restricted cubic spline (RCS) was utilized to further explore the relationship between sleep duration and cognitive frailty risk. The significance level of the statistical tests was set as $p < 0.05$.

3 Results

3.1 Characteristics of 9,970 participants grouped by whether they were diagnosed with cognitive frailty

In this study, 1,684 participants were defined as cognitive frailty, accounting for 16.89%, with a median age of 68 years, 48.56% of participants were male and 51.44% were female. Participants with cognitive frailty were more likely to be “unmarried,” live in rural areas, and were female, with no social activity in the last month (see [Table 1](#)).

3.2 Prevalence and adjusted OR (95% CI) of cognitive frailty by groups of napping time, sleep duration and quality

[Table 2](#) presents that individuals who sleep less than 6 h have a significantly higher risk of cognitive frailty. This finding was confirmed in different models. Specifically, in Model 1, those who slept less than 6 h had a 55% increased risk of developing cognitive frailty compared to those who slept 6 to 9 h (OR = 1.55, 95% CI: 1.36, 1.77); In fully adjusted model 2, those who slept less than 6 h had a 48% increased risk of developing cognitive frailty (OR = 1.48, 95% CI: 1.25, 1.64); In Model 3, which was further adjusted for sleep quality or daytime napping, those who slept less than 6 h had a 27% increased risk of developing cognitive frailty (OR = 1.27, 95% CI: 1.13, 1.49); Conversely, no significant differences were observed in participants who slept more than 9 h across the models (model 1: OR = 1.03, 95% CI: 0.88, 1.29; model 2: OR = 0.98, 95% CI: 0.76, 1.19; model 3: OR = 1.27, 95% CI: 0.68, 1.17).

Regarding sleep quality, in Model 1, individuals with poor sleep quality had a 38% increased risk of developing cognitive frailty compared to those with good sleep quality (OR = 1.38, 95% CI: 1.14, 1.57). This trend was further confirmed by adjusting for all covariates in Model 2. In addition, after adjusting for potential confounders and napping (model 3), people with poor sleep quality had a higher chance of developing cognitive frailty [OR 1.88 (95% CI: 1.58, 2.19)].

About the nap aspect, after further adjustment for sleep quality (Model 3), participants who did not nap and who took excessive naps for > 90 min/day had high rates of cognitive frailty, with an OR (95% CI) of 1.23 (1.04 to 1.46) and 1.69 (1.38 to 2.19), respectively.

3.3 Additive interactive effect of sleep quality, sleep duration and napping time on cognitive frailty in older adults people based on multivariate logistic regression analysis

[Table 3](#) shows that Multivariate Logistic regression analysis was conducted with the older adults cognitive frailty as the dependent variable, sleep quality, Sleep duration and napping time as independent variables, and controlling for confounding factors such as age, education level, residence, and the number of chronic diseases. The results showed that there was an interaction among sleep quality, sleep duration and siesta on cognitive frailty in the older adults, and the p values of the interaction were 0.035, 0.042 and 0.005, respectively.

Specifically, the coefficient for the interaction between sleep duration and sleep quality was 0.728 (SE = 0.793, $p = 0.035$), suggesting a significant interaction between sleep duration and sleep quality on depression (95% CI: 0.631, 2.585). The coefficient of the interaction between sleep quality and napping was 0.896 (SE = 1.485, $p = 0.042$), suggesting that there was a significant interaction between sleep quality and napping on cognitive frailty (95% CI: 0.802, 2.248). The coefficient of the interaction between sleep duration and napping is 0.963 (SE = 1.129, $p = 0.005$), suggesting a significant interaction between sleep quality and napping on cognitive frailty (95% CI: 1.022, 2.748). Co-linearity diagnosis: VIF of all variables is less than 3, indicating that the multicollinearity problem is negligible. This suggests that these factors do not act independently, but act together in the effects of sleep quality, sleep duration, and napping on cognitive weakness, and it is necessary to further stratify sleep quality, sleep duration, and napping.

3.4 Joint effect of sleep duration and sleep quality on cognitive frailty

[Table 4](#) displays that due to the interaction between sleep duration and sleep quality on cognitive weakness in the older adults ($\beta = 0.728$, 95% CI: 0.631 ~ 2.583, $p = 0.035$). When the sleep quality is poor, the protective effect of prolonged sleep time on cognition is weakened. Model multicollinearity acceptable (VIF = 2.0). We observe that individuals with short sleep duration have a higher risk of developing CF, regardless of whether they sleep well or poorly. However, people with short sleep duration and poor sleep quality have a 65% increased risk of cognitive frailty. The individuals who slept for long periods of time did not show a similar increased risk.

3.5 Joint effect of daytime napping duration and sleep quality on cognitive frailty

Since we found a significant interaction between daytime napping and sleep quality ($\beta = 0.896$, 95% CI: 0.802 ~ 2.248, $p = 0.042$), a stratified analysis by sleep quality ([Table 5](#)) showed a 123% increased risk of developing cognitive frailty in participants with good sleep quality who napped for more than 90 min, with an OR of 2.23 (95% CI: 1.72 ~ 2.86). No association was found between napping and cognitive frailty in participants with poor sleep quality. The model multicollinearity is acceptable (VIF = 2.3).

3.6 Joint effect of daytime napping duration and nighttime sleep duration on cognitive frailty

The relationship between sleep duration, napping duration and cognitive frailty is shown in [Table 6](#) ($\beta = 0.963$, 95% CI: 1.022 ~ 2.874, $p = 0.005$). No significant association was found between daytime naps and cognitive impairment in the shorter and moderate nocturnal sleep groups. In the subgroup with longer sleep duration, participants who took excessive naps had a significantly increased risk of cognitive frailty (OR = 1.62, 95% CI: 1.14 ~ 2.30, $p < 0.001$), i.e., excessive

TABLE 1 Characteristics of 9,970 participants by CF group and non-CF group.

Item	Total	CF	Non-CF	<i>P</i>
N	9,970	1,684	8,286	-
Age (years), mean (SD)	67 (63,71)	68 (64,72)	66 (63,70)	<0.0001
Gender, <i>n</i> %				<0.0001
Male	4,841 (48.6)	690 (40.1)	4,151 (50.1)	
Female	5,129 (51.4)	994 (59.9)	4,135 (49.9)	
Education level				<0.0001
Elementary school	2,884 (28.9)	806 (47.9)	2,078 (25.1)	
Junior high school	4,313 (43.3)	434 (25.8)	3,879 (46.8)	
High school	1,362 (13.7)	276 (16.4)	1,086 (13.1)	
University	1,411 (14.1)	168 (9.9)	1,243 (15)	
Area of residence				<0.0001
Urban	4,279 (42.9)	624 (37.1)	3,655 (44.1)	
Rural	5,691 (57.1)	1,060 (62.9)	4,631 (55.9)	
Current marital status, <i>n</i> %				<0.0001
Not married	1,618 (16.2)	325 (19.3)	1,293 (15.6)	
Married or cohabitated	8,352 (83.8)	1,359 (80.7)	6,993 (84.4)	
BMI(kg/m ²)				<0.0001
< 18.5	4,156 (41.7)	818 (48.6)	3,338 (40.3)	
18.5–23.9	4,380 (44.0)	595 (35.3)	3,785 (45.7)	
≥ 24	1,434 (14.3)	271 (16.1)	1,163 (15.0)	
Drinking				0.053
Never	4,656 (46.7)	767 (45.5)	3,889 (46.9)	
Former	683 (6.9)	234 (13.9)	449 (7.9)	
Current	4,431 (44.4)	683 (40.6)	3,748 (45.2)	
Smoking				0.041
Never	7,886 (79.1)	1,315 (78.1)	6,571 (79.3)	
Former	1,172 (11.8)	211 (12.5)	961 (11.6)	
Current	912 (9.1)	158 (9.4)	754 (9.1)	
Family income (anu/year), %				<0.0001
< 10,000	1,254 (12.5)	310 (18.4)	944 (11.4)	
10,000–19,999	1,881 (18.9)	497 (29.5)	1,384 (16.7)	
≥ 20,000	6,835 (68.6)	877 (52.1)	5,958 (71.9)	
Social interaction				0.0005
Never	6,196 (62.1)	918 (54.5)	5,278 (63.7)	
At least once a month	3,774 (37.9)	766 (45.5)	3,008 (36.3)	
Numbers of chronic diseases				0.324
≤1	2,919 (29.3)	317 (18.8)	2,602 (31.4)	
2–3	2,235 (22.4)	495 (29.4)	1,740 (21.0)	
≥4	4,816 (48.3)	872 (51.8)	3,944 (47.6)	
Depression	1,611 (16.2)	527 (31.3)	655 (7.9)	<0.0001
Daytime napping	30 (0,60)	40 (0,60)	30 (0,60)	<0.0001
Sleep duration	6 (5,7)	6 (5,8)	6 (5,7)	0.042
Sleep quality				<0.0001
Good	2,433 (24.4)	295 (17.5)	2,138 (25.8)	
Moderate	2,382 (23.9)	443 (26.3)	1,939 (23.4)	
Poor	5,155 (51.7)	946 (56.2)	4,209 (50.8)	

CF: cognitive frailty.

TABLE 2 Prevalence and adjusted OR (95% CI) of cognitive frailty by groups of napping time, sleep duration and quality.

Variables	Model 1	Model 2	Model 3
Sleep quality			
Good	Ref	Ref	Ref
moderate	1.36 (1.18,1.56)	1.57 (1.38,1.79)	1.68 (1.48,1.89)
Poor	1.38 (1.14,1.57) ***	1.67 (1.47,1.95) ***	1.88 (1.58,2.19) ***
Napping time			
60-90 min	Ref	Ref	Ref
Non-nappers	1.28 (1.13,1.46)	1.25 (1.04,1.47)	1.23 (1.04,1.46) ***
< 60 min	1.23 (1.08,1.59)	1.21 (1.06,1.56)	1.19 (1.02,1.48)
> 90 min	1.71 (1.35,2.12)	1.68 (1.31,2.17)	1.69 (1.38,2.19) ***
Sleep duration			
6-9 h	Ref	Ref	Ref
< 6 h	1.55 (1.36,1.77) ***	1.48 (1.25,1.64) ***	1.27 (1.13,1.49) ***
> 9 h	1.03 (0.88,1.29)	0.98 (0.76,1.19)	0.96 (0.68,1.17)

Model 1: adjusting for sex, age, education level, residence, current marital status, personal income, smoking status, alcohol consumption, social interaction; Model 2: additionally adjusting for BMI, numbers of chronic diseases; Model 3: additionally adjusting for daytime napping or nighttime sleep quality. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

TABLE 3 Additive interactive effect of sleep quality, sleep duration and napping time on cognitive frailty in older adults people based on multivariate logistic regression analysis.

Variables	B	SE	Wald χ^2	p value	OR(95%CI)	VIF
Sleep quality X Sleep duration	0.728	0.793	0.048	0.035	1.535 (0.631, 2.585)	2.0
Sleep quality X Napping time	0.896	1.485	0.082	0.042	1.347 (0.802, 2.248)	2.3
Sleep duration X Napping time	0.963	1.129	0.384	0.005	1.476 (1.022, 2.874)	1.6

TABLE 4 Joint effect of sleep duration and sleep quality on cognitive frailty ($n = 1,684$).

Variables	Model 1	Model 2	Model 3
Duration quality			
6-9 h good	Ref	Ref	Ref
<6 h good	2.09 (1.65, 2.32) ***	1.85 (1.36, 2.01) ***	1.45 (1.14, 1.85) ***
<6 h poor	2.12 (1.74, 2.48) ***	1.90 (1.48, 2.23) ***	1.65 (1.34, 2.15) ***
>9 h good	1.62 (1.23, 2.34)	1.52 (0.98, 2.33)	1.25 (0.74, 1.95)
>9 h poor	1.42 (0.84, 2.88)	1.20 (0.58, 2.63)	1.05 (0.64, 2.35)

Model 1: adjusting for sex, age, education level, residence, current marital status, personal income, smoking status, alcohol consumption, social interaction; Model 2: additionally adjusting for BMI, numbers of chronic diseases; Model 3: additionally adjusting for daytime napping or nighttime sleep quality. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

napping (> 90 min) was associated with cognitive frailty in participants who slept 9 h or more at night. The model multicollinearity is acceptable (VIF = 1.6).

The study revealed a linear relationship between sleep duration and cognitive frailty, but also detected a nonlinear relationship (P-nonlinearity <0.001). Using a fully adjusted model, restrictive cubic spline regression showed a nonlinear (U-shaped) relationship between nighttime sleep duration and napping time and the risk of cognitive frailty (Figure 2). As sleep duration increased to 7.5 h per night or nap 1.2 h, the risk of cognitive impairment decreased, followed by a reversal trend above these thresholds. Therefore, our study suggests that both too short and too long sleep may increase the risk of cognitive weakness. The results for longer sleep durations in the restricted cubic spline regression differed from the non-significant results for the fully adjusted linear model. When comparing the linear model and the nonlinear model, it is shown that the linear model

performs better than the nonlinear model. Therefore, our study tends to conclude that prolonged sleep is unlikely to increase the risk of cognitive weakness. This difference may be attributed to the relatively small sample size in the long sleep duration group, which complicates the interpretation of the results.

3.7 Mediation analysis of the association of sleep disorders with cognitive frailty by depression

Based on the significant association between sleep disorder and cognitive frailty observed among older adults in the abovementioned results, a mediation analysis was further conducted to examine whether depression mediated the association between sleep disorder and cognitive frailty (Table 7). The results showed that the association

TABLE 5 Joint effect of daytime napping duration and sleep quality on cognitive frailty (*n* = 1,684).

Variables	Model 1	Model 2
Good sleep quality		
None	Ref	Ref
< 60 min	1.86 (1.36,2.25)	1.68 (1.22,2.06)
60-90 min	2.04 (1.62,2.96)**	1.89 (1.03,2.64)**
> 90 min	2.23 (1.72,2.86)***	2.10 (1.63,2.75)***
Poor sleep quality		
None	Ref	Ref
< 60 min	1.16 (0.85,1.48)	1.13 (0.76,1.43)
60-90 min	1.18 (0.79,1.65)	1.36 (0.96,1.79)
> 90 min	1.06 (0.66,1.25)	1.12 (0.71, 1.44)

Model 1: adjusting for sex, age, education level, residence, personal income, smoking status, alcohol consumption, social interaction; Model 2: additionally adjusting for BMI, numbers of chronic diseases. *P* < 0.05, ***P* < 0.01, ****P* < 0.001.

TABLE 6 Joint effect of daytime napping duration and nighttime sleep duration on cognitive frailty (*n* = 1,684).

Variables	Model 1	Model 2	Model 3
<6 h/nighttime			
None	Ref	Ref	Ref
< 60 min	0.98 (0.48, 1.95)	0.94 (0.40, 1.81)	0.95 (0.53, 2.07)
60-90 min	0.88 (0.37, 1.89)	0.75 (0.42, 1.76)	0.64 (0.45, 1.57)
>90 min	1.14 (0.38, 1.98)	1.17 (0.78, 2.06)	1.06 (0.79, 1.96)
6-9 h/nighttime			
None	Ref	Ref	Ref
< 60 min	0.98 (0.47, 1.83)	0.95 (0.34, 1.93)	0.99 (0.78, 2.17)
61-90 min	0.73 (0.38, 1.66)	0.85 (0.52, 1.86)	0.65 (0.56, 1.85)
>90 min	1.28 (0.59, 2.08)	1.25 (0.69, 1.91)	1.18 (0.91, 2.21)
>9 h/nighttime			
None	Ref	Ref	Ref
<60 min	0.78 (0.33, 1.78)	0.85 (0.36, 1.79)	0.84 (0.65, 1.94)
61-90 min	1.34 (0.56, 1.95)**	1.43 (0.86, 2.21)**	1.56 (1.21, 2.54)**
>90 min	1.46 (0.88, 1.98)***	1.55 (0.91, 2.36)***	1.62 (1.44, 2.60)***

Model 1: adjusting for sex, age, education level, residence, personal income, smoking status, alcohol consumption, social interaction; Model 2: additionally adjusting for BMI, numbers of chronic diseases; Model 3: addition-ally adjusting for daytime napping or nighttime sleep quality. *P* < 0.05, ***P* < 0.01, ****P* < 0.001.

between sleep and risk of cognitive frailty was significantly mediated by depression. After adjusting for potential confounding factors, depression completely mediated the sleep disorder–cognitive frailty association, with a significant indirect effect (OR = 0.995, 95%CI: 0.993 ~ 0.997, *p* < 0.001) but a non-significant direct effect (OR = 1.004, 95% CI: 0.998 ~ 1.012, *p* = 0.278). It can be seen from the mediation analysis revealed that the relationships of poor sleep quality and sleep duration with cognitive frailty were mediated by depression in older adults.

4 Discussion

This study indicated the prevalence of cognitive frailty was 16.89%. Participants with cognitive frailty were more likely to be “unmarried,” live in rural areas, have low levels of education, have

had no social activity in the last month, and have depression. Previously, a study of 7,338 older adults people in the community by Aliberti et al. (58). reported a 5% prevalence of cognitive frailty. In a study of 1,620 older patients over 65 years of age at the Toulouse Frailty Day Hospital, the prevalence of cognitive frailty was as high as 26.7 percent (59). The difference in the detection rate of cognitive frailty may be due to the different age ranges and urban–rural distribution of the survey subjects. In addition, it may also be related to the assessment methods and evaluation tools. We assessed the association between sleep duration, sleep quality, nap duration, and cognitive frailty after controlling for confounding factors. Participants who slept less than 6 h, no and excessive napping, and poor sleep quality had significantly increased risk of cognitive vulnerability. Previous investigators, Kong et al. (60). using a cross-sectional survey of community-dwelling older adults with type 2 diabetes, found that those who had insufficient sleep at night were at

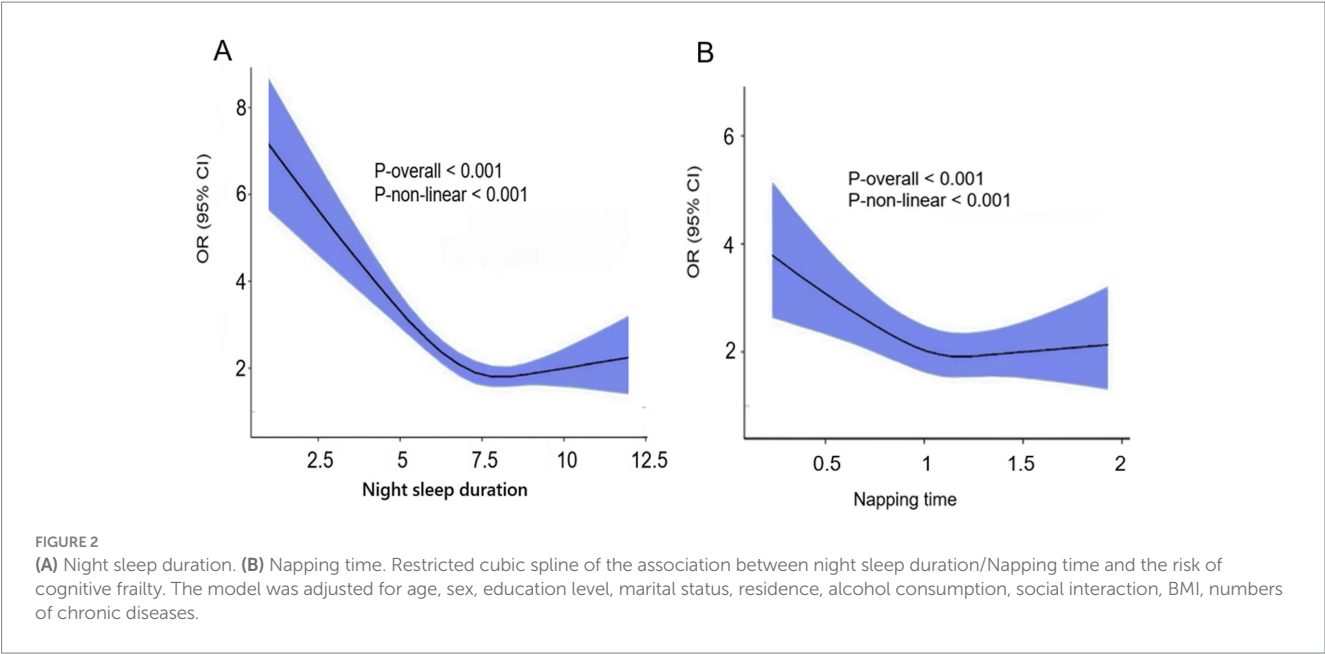


TABLE 7 Mediation analysis of the association of sleep disorders with cognitive frailty by depression.

Variables	Coefficient (95% BCa CI)	OR (95% CI)	Z	p value
Sleep quality				
Indirect effect-path ab	0.030 (0.015–0.049)	1.022 (1.009–1.035)	3.453	<0.001
Direct effect-path c'	0.021(–0.006–0.04)	1.034 (1.006–1.060)	2.341	0.159
Direct effect-path c	0.052 (0.020–0.085)	1.059 (1.028–1.093)	3.384	<0.001
Napping time				
Indirect effect-path ab	0.022 (0.011–0.034)	1.030 (1.015–1.048)	3.867	<0.001
Direct effect-path c'	0.023 (0.009–0.071)	1.025 (0.093–1.057)	1.325	0.146
Direct effect-path c	0.053 (1.08,1.59)	1.051 (1.026–1.084)	3.342	<0.001
Sleep duration				
Indirect effect-path ab	–0.004 (–0.006 to –0.002)	0.995 (0.993–0.997)	–4.187	<0.001
Direct effect-path c'	0.005 (–0.001 to 0.011)	1.004 (0.998–1.012)	1.207	0.278
Direct effect-path c	0.001 (–0.005 to 0.009)	1.001 (0.995–1.006)	0.363	0.734

Adjusted for age, gender, smoking, alcohol consumption, Social interaction and chronic diseases. Path ab coefficient represents 5,000 bootstrapped samples and BCa 95% CIs. Path c': the direct effect of the exposure variable (sleep) on the outcome (cognitive frailty), holding the effect of depression constant. Path c: the total effect of exposure (sleep) on the outcome (cognitive frailty). BCa, bias-corrected and accelerated; OR, odds ratio.

higher risk for cognitive frailty. In contrast, the Leng report showed that men who took excessive naps per day were 66% more likely to have cognitive impairment than those who took 30-min nap (61). In Li's study, moderate nappers had better overall cognitive performance than those who did not nap or who took long naps (62), which is consistent with the results of this study.

Exploring the mechanisms by which sleep deprivation affects cognitive function is essential to reduce its harm. Complete sleep deprivation (TSD), as an extreme manifestation of sleep deprivation, is a valid model for comprehensively studying how sleep deprivation impairs cognition (63). Numerous studies have shown that TSD impairs executive function. Honn et al. (64) revealed that TSD impairs feedback blunting in cognitive flexibility tasks. Aidman et al. (65) conducted a study showing that TSD impairs three executive

functions, namely cognitive flexibility, inhibitory control, and working memory capacity. In addition, Cain et al. (66) found that TSD affected the performance of the Stroop task through an increase in overall response time. In addition, some studies have shown that female cognitive function is less susceptible to the effects of TSD (67, 68), which may be explained by gonadal hormone status (69). Taken together, these studies suggest that cognitive function is negatively affected by TSD. There was a significant interaction among sleep quality, sleep duration and nap duration on cognitive frailty. Specifically, After controlling for confounding factors such as gender, age, education level, and nighttime sleep duration, participants with good sleep quality had a significantly increased risk of excessive napping for cognitive frailty. No association was found between napping and cognitive frailty among participants with poor sleep

quality, highlighting the importance of modifiable behavioral factors in preventing cognitive frailty. Individuals with short sleep duration are at higher risk for cognitive weakness, regardless of whether they have good or poor sleep quality. However, individuals with longer sleep duration did not show a similar increased risk. Excessive naps (> 90 min) were associated with increased risk for cognitive frailty in the longer sleep duration subgroup, whereas no significant association was found between daytime naps and cognitive frailty in the shorter and moderate sleep groups. Many researchers have reported that excessive napping during the day may lead to cognitive decline in older adults with or without dementia. Excessive daytime napping has been suggested to increase the risk of Alzheimer's dementia and is associated with deterioration of cognitive function (70–72).

The underlying mechanisms underlying the association between sleep disturbance and cognitive weakness in patients are poorly understood. The mediation analysis used in this study may provide epidemiological evidence that mediating effects through depression may be an underlying mechanism that influences sleep in relation to cognitive vulnerability. Sleep disturbances (e.g., insomnia and poor sleep quality) are often combined with cognitive weakness in old age. Previous literature has shown that sleep disturbances may lead to depression and reduce physical inactivity (73), thereby increasing the risk of negative health outcomes.

In conclusion, this study systematically reveals for the first time the synergistic effect between sleep quality, nighttime sleep duration, and napping behavior on cognitive frailty in the older adults population. Different from previous studies that only focused on a single sleep factor, we broke through the limitations of traditional logistic regression by introducing interaction terms and hierarchical analysis, and constructed a multi-dimensional dynamic model including sleep quality, duration, and napping. Our stratified analysis revealed a significant association between sleep disorders (including long naps, poor sleep quality, short sleep duration, and long sleep duration) and cognitive frailty in Chinese older adults. For example, the interaction effect analysis of 'sleep quality \times nap duration' ($p = 0.042$) revealed the moderating effect of the temporal distribution of sleep behavior on cognitive frailty. Participants with good sleep quality who took naps for more than 90 min had a 123% increased risk of cognitive frailty and an OR of 2.23 (95% CI: 1.72 ~ 2.86). These results suggest that a holistic assessment of sleep patterns, rather than a single indicator, is key to predicting cognitive health in older adults. Unlike traditional linear assumptions, our study found a U-shaped association between sleep duration and cognitive function, with optimal nighttime sleep duration being 6–9 h, while napping for more than 90 min may have negative effects. In contrast to the public health recommendation that napping is universally beneficial, this study highlighted the 'double-edged sword' effect of napping, with a significant cognitive protection effect at nighttime sleeps for 6–9 h and naps for 61–90 min (OR = 0.65). This calls for a re-examination of the universality of sleep guidelines for the older adults.

This study provides a theoretical basis and practical tools for the early prevention and control of cognitive weakness in the older adults by revealing the complex interaction between sleep quality, duration and napping. We call for multidimensional sleep assessment to be included in the core content of geriatric health management, and to develop a hierarchical intervention strategy for

high-risk groups to delay cognitive decline and improve quality of life.

There are also some limitations to this study. First of all, the evaluation scale used in the CHARLS database is a subjective evaluation scale, and no objective evaluation measures are used for its indicators. Secondly, the CHARLS sleep duration and sleep quality evaluation method was self-report, and there was a large degree of subjectivity in the self-evaluation of sleep duration and sleep quality. In the future, a cohort study is planned to carry out a prospective study to further analyze the combined effects of different levels, intensities and sleep duration on cognitive impairment in the older adults through objectively measured physical activity data, in order to provide more evidence-based basis for lifestyle intervention for cognitive impairment in the older adults.

5 Conclusion

Chinese older adults with poor sleep quality are at a 67% higher risk of cognitive weakness, and individuals with less than 6 h of sleep are at a 48% higher risk of cognitive weakness; No napping and excessive napping, the risk of cognitive debilitation increased by 23 and 69%, respectively. There is an additive interaction between sleep duration and quality and daytime napping on cognitive frailty in the older adults.

Data availability statement

The original contributions presented in the study are included in the article/[Supplementary material](#), further inquiries can be directed to the corresponding author.

Ethics statement

The studies involving humans were approved by the Biomedical Ethics Committee of Peking University (IRB00001052-11015). The studies were conducted in accordance with the local legislation and institutional requirements. The participants provided their written informed consent to participate in this study.

Author contributions

JQ: Methodology, Software, Writing – original draft, Writing – review & editing. XY: Methodology, Software, Writing – original draft, Writing – review & editing. QZ: Formal analysis, Funding acquisition, Investigation, Resources, Visualization, Writing – review & editing.

Funding

The author(s) declare that financial support was received for the research and/or publication of this article. This study was supported by Major Research Plan of National Natural Science Foundation of China (Grant No.92163213), General Program of National Natural Science Foundation of China (Grant No. 81970085), Tianjin science

and technology plan project (Grant No. 21JCZDJC00940), Tianjin health science and technology projects (Grant No. TJWJ2022XK001), Tianjin Key Medical Discipline (Specialty) Construction Project (Grant No. TJYXZDXK-006A), National Key Research and Development Program of China (No. 2023YFC3605200), Major project of Strategic Research and Consulting Project of the Chinese Academy of Engineering (No.2023-DFZD-58).

Acknowledgments

We gratefully thank China Health and Retirement Longitudinal Study (CHARLS) team for providing high-quality, nationally representative data.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

References

- Foreman KJ, Marquez N, Dolgert A. Forecasting life expectancy, years of life lost, and all-cause and cause-specific mortality for 250 causes of death: reference and alternative scenarios for 2016–40 for 195 countries and territories. *Lancet*. (2018) 392:2052–90. doi: 10.1016/S0140-6736(18)31694-5
- Office of the Leading Group for the Seventh National Population Census of the State Council. Main data of the 7th National Population Census in 2020. Beijing: China Statistics Press (2021).
- Won CW, Lee Y, Kim S, Yoo J, Kim M, Ng TP, et al. Modified criteria for diagnosing "cognitive frailty". *Psychiatry*. (2018) 15:839–42. doi: 10.30773/pi.2018.05.22
- Okoro CA, Hollis ND, Cyrus AC, Griffin-Blake S. Prevalence of disabilities and health care access by disability status and type among adults—United States, 2016. *MMWR Morb Mortal Wkly Rep*. (2018) 67:882–7. doi: 10.15585/mmwr.mm6732a3
- Avila-Funes JA, Amieva H, Barberger-Gateau P, Le Goff M, Raoux N, Ritchie K, et al. Cognitive impairment improves the predictive validity of the phenotype of frailty for adverse health outcomes: the three-city study. *J Am Geriatr Soc*. (2009) 57:453–61. doi: 10.1111/j.1532-5415.2008.02136.x
- Kelaiditi E, Cesari M, Canevelli M, van Kan GA, Ousset PJ, Gillette-Guyonnet S, et al. Cognitive frailty: rationale and definition from an (I.A.N.A./I.A.G.G.) international consensus group. *J Nutr Health Aging*. (2013) 17:726–34. doi: 10.1007/s12603-013-0437-5
- Shimada H, Lee S, Makizako H, Doi T, Chen LK, Arai H. Cognitive frailty predicts incident dementia among community-dwelling older people. *J Clin Med*. (2018) 7:250. doi: 10.3390/jcm7090250
- Ng PT, Feng L, Shwe M. Frailty in older persons: multisystem risk factors and the frailty risk index (FRI). *J Am Med Dir Assoc*. (2014) 15:635–42. doi: 10.1016/j.jamda.2014.03.008
- Esteban-cornejo I, Cabanas-sánchez V, Rodriguez-artalejo F. Cognitive frailty and mortality in a national cohort of older adults: the role of physical activity. *Mayo Clin Proc*. (2019) 94:1–10. doi: 10.1016/j.mayocp.2018.10.027
- Ruan Q, Xiao F, Gong K. Prevalence of cognitive frailty phenotypes and associated factors in a community-dwelling elderly population. *J Nutr Health Aging*. (2020) 24:172–80. doi: 10.1007/s12603-019-1286-7
- Ma L, Zhang L, Zhang Y. Cognitive frailty in China: results from China comprehensive geriatric assessment study. *Front Med*. (2021) 4:174. doi: 10.3389/fmed.2017.00174
- Navarro-Pardo E, Facal D, Campos-Magdaleno M. Prevalence of cognitive frailty, do psychosocial-related factors matter? *Brain Sci*. (2020) 10:968. doi: 10.3390/brainsci10120968
- Qu X, Dai L, Zhang P. Meta-analysis of influencing factors of cognitive frailty in the elderly. *Modern Clinic Nurs*. (2022) 21:6–10. doi: 10.1111/ijn.13306
- Kim H, Awata S, Watanabe Y. Cognitive frailty in community-dwelling older Japanese people: prevalence and its association with falls. *Geriatr Gerontol Int*. (2019) 19:647–53. doi: 10.1111/ggi.13685
- Liu YT, Fan JY, Zhao HM. Research progress on the status and influencing factors of cognitive frailty in the elderly. *J Nurs*. (2023) 34:101–5. doi: 10.1111/psys.1-3087

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Supplementary material

The Supplementary material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fpubh.2025.1596965/full#supplementary-material>

- Yin Y, Chen S, Song T, Zhou Q, Shao Y. Cognitive load moderates the effects of Total sleep deprivation on working memory: evidence from event-related potentials. *Brain Sci*. (2023) 13:898. doi: 10.3390/brainsci13060898
- Song T, Du F, Xu L, Peng Z, Wang L, Dai C, et al. Total sleep deprivation selectively impairs motor preparation sub-stages in visual search task: evidence from lateralized readiness potentials. *Front Neurosci*. (2023) 17:989512. doi: 10.3389/fnins.2023.989512
- Krittanawong C. Association between short and long sleep durations and cardiovascular outcomes: a systematic review and meta-analysis. *Eur Heart J Acute Cardiovasc Care*. (2019) 8:762–70. doi: 10.1177/2048872617741733
- Long JM. Sleep duration and health outcomes: a systematic review, meta-analysis and meta-regression. *Sleep Med Rev*. (2018) 39:25–36. doi: 10.1016/j.smrv.2017.06.011
- Wang M, Xiang X, Zhao Z, Liu Y, Cao Y, Guo W, et al. Association between self-reported napping and risk of cardiovascular disease and all-cause mortality: a meta-analysis of cohort studies. *PLoS One*. (2024) 19:266. doi: 10.1371/journal.pone.0311266
- Li M, Wang N, Dupre ME. Association between the self-reported duration and quality of sleep and cognitive function among middle-aged and older adults in China. *J Affect Disord*. (2022) 304:20–7. doi: 10.1016/j.jad.2022.02.039
- Liu WY, Wu WW. Advance in correlation between sleep disorder and frailty in the elderly (review). *Chin J Rehabil Theory Pract*. (2020) 26:1435–8. doi: 10.3969/j.issn.1006-9771.2020.12.011
- Zhang YT, Chen K, Xu YX. Research progress of first night effect of polysomnography monitoring. *Nat Med J Chin*. (2024) 104:2590–4. doi: 10.3760/cma.j.cn112137-20231219-01410
- Ricciardiello A, Mckinnon AC, Mowszowski L. Assessing sleep architecture and cognition in older adults with depressive symptoms attending a memory clinic. *J Affect Disord*. (2024) 348:35–43. doi: 10.1016/j.jad.2023.12.032
- Kim JH, Elkhadem AR, Duffy JF. Circadian rhythm sleep wake disorders in older adults. *Sleep Med Clin*. (2022) 17:241–52. doi: 10.1016/j.jsmc.2022.02.003
- Li Y, Sahakian BJ, Kang J. The brain structure and genetic mechanisms underlying the nonlinear association between sleep duration, cognition and mental health. *Nat Aging*. (2022) 2:425–37. doi: 10.1038/s43587-022-00210-2
- Xiang GZ, Su XJ, Jia JX. Advances in understanding the mechanism of baicalin in ameliorating cognitive dysfunction associated with sleep disorders. *Int J Geriatr*. (2024) 5:482–5. doi: 10.3969/j.issn.1674-7593.2024.04.018
- Lü X, Yang XY, Zhou ZK. Neural oscillation and its application in mechanistic study of neuropsychiatric disorders. *Acta Physiol Sin*. (2022) 74:657–68.
- Conolee LS, Stevenson AJ, Muñoz Maniega S. DNA methylation and protein markers of chronic inflammation and their associations with brain and cognitive aging. *Neurology*. (2021) 97:e2340–52.
- Yang E, Wang SQ, Luo D. Study on the combined effects of sleep duration and physical activity on cognitive impairment in Chinese elderly. *Chin J Prev Control Chronic Dis*. (2024) 32:741–5. doi: 10.16386/j.cjpcd.issn.1004-6194.2024.10.004

31. Sun F, Wang TL, Wu N. Research on the relation-ship between physical activity cognition and sleep quality in the working population: n analysis based on the mediating effect of social support. *Mod Prev Med.* (2024) 51:3114–8. doi: 10.20043/j.cnki.MPM.202405301
32. Arpi M, Fernandes M, Mercuri NB. Sleep biomarkers for predicting cognitive decline and Alzheimer's dis-ease: a systematic review of longitudinal studies. *J Alzheimers Dis.* (2024) 97:121–43. doi: 10.3233/JAD-230933
33. Lowe CJ, Safati A, Hall PA. The neurocognitive consequences of sleep restriction: a meta-analytic review. *Neurosci Biobehav Rev.* (2017) 80:86. doi: 10.1016/j.neubiorev.2017.07.010
34. Philip P, Akerstedt T. Transport and industrial safety, how are they affected by sleepiness and sleep restriction? *Sleep Med Rev.* (2006) 10:347–56. doi: 10.1016/j.smrv.2006.04.002
35. Lim J, Dinges DF. Sleep deprivation and vigilant attention. *Ann N Y Acad Sci.* (2008) 1129:305–22. doi: 10.1196/annals.1417.002
36. McMahon WR, Ftouni S, Drummond SPA, Maruff P, Lockley SW, Rajaratnam SMW. The wake maintenance zone shows task dependent changes in cognitive function following one night without sleep. *Sleep.* (2018) 41:148. doi: 10.1093/sleep/zsy148
37. Gibbings A, Ray LB, Berberian N, Nguyen T, Zandi Shahidi A, Owen AM. EEG and behavioural correlates of mild sleep deprivation and vigilance. *Clin Neurophysiol.* (2020) 132:45–55. doi: 10.1016/j.clinph.2020.10.010
38. Stepan ME, Altmann EM, Fenn KM. Effects of total sleep deprivation on procedural place keeping: more than just lapses of attention. *J Exp Psychol.* (2020) 149:800–6. doi: 10.1037/xge0000717
39. Boardman JM, Bei B, Mellor A, Anderson C, Sletten TL, Drummond SPA. The ability to self-monitor cognitive performance during 60h total sleep deprivation and following 2nights recovery sleep. *J Sleep Res.* (2017) 27:12633. doi: 10.1111/jsr.12633
40. Deliens G, Bukowski H. The impact of sleep deprivation on visual perspective taking. *J Sleep Res.* (2018) 27:175–83. doi: 10.1111/jsr.12595
41. Liu H, Xiang Z, Yang M, Sun J, Juanjuan P, Wangquan X, et al. The mediating role of daily living ability and sleep in depression and cognitive function based on a structural equation model. *BMC Geriatr.* (2025) 25:1–10. doi: 10.1186/s12877-025-05871-3
42. Dong P, Cheng C, Yin W, Li Z, Shi Y, Gao M, et al. Frailty as a mediator between sleep quality and cognitive impairment among the rural older adults: a cross-sectional study. *BMC Geriatr.* (2025) 25:1–9. doi: 10.1186/s12877-024-05657-z
43. Aditi S, Singh SK, Jaiswal AK, Verma M. Is there a ubiquitous association between sleep disorder and frailty? *BMC Geriatr.* (2023) 23:1–11. doi: 10.1186/s12877-023-04148-x
44. Ruiz-Castell M, Makovski TT, Bocquet V, Stranges S. Sleep duration and multimorbidity in Luxembourg: results from the European health examination survey in Luxembourg, 2013–2015. *BMJ Open.* (2019) 8:e26942. doi: 10.1136/bmjopen-2018-026942
45. Watson NF, Badr MS, Belenky G, Bliwise DL, Buxton OM, Buysse D, et al. Recommended amount of sleep for a healthy adult: a joint consensus statement of the American Academy of sleep medicine and Sleep Research Society. *J Clin Sleep Med.* (2015) 38:843–4. doi: 10.5665/sleep.4716
46. Sun H, Qin K, Zou CF, Wang HH, Lu C, Chen W, et al. The association of nighttime sleep duration and quality with chronic kidney disease in middle-aged and older Chinese: a cohort study. *Sleep Med.* (2021) 86:25–31. doi: 10.1016/j.sleep.2021.08.007
47. Fried LP, Tangen CM, Walston J. Frailty in older adults: evidence for a phenotype. *J Gerontol A Biol Sci Med Sci.* (2001) 56:146–56. doi: 10.1093/gerona/56.3.m146
48. Cao L, Zhao Z, Ji C. Association between solid fuel use and cognitive impairment: a cross-sectional and follow-up study in a middle-aged and older Chinese population. *Environ Int.* (2021) 146:106–251. doi: 10.1016/j.envint.2020.106251
49. Zhou Y, Chen Z, Shaw I. Association between social participation and cognitive function among middle- and old-aged Chinese: a fixed-effects analysis. *J Glob Health.* (2020) 10:801. doi: 10.7189/jogh.10.020801
50. Chan J, Chan TK, Kwok T. Cognitive training interventions and depression in mild cognitive impairment and dementia: a systematic review and meta-analysis of randomized controlled trials. *Age Ageing.* (2020) 49:738–47. doi: 10.1093/ageing/afaa063
51. Yuan M, Xu C, Fang Y. The transitions and predictors of cognitive frailty with multi-state Markov model: a cohort study. *BMC Geriatr.* (2022) 22:550. doi: 10.1186/s12877-022-03220-2
52. Liu H, Lee D, Zhao X. Longitudinal impact of frailty states and sleep duration on subsequent depressive symptoms of older adults. *J Am Geriatr Soc.* (2021) 69:1003–11. doi: 10.1111/jgs.16999
53. Crimmins EM, Kim JK, Langa KM. Assessment of cognition using surveys and neuropsychological assessment: the health and retirement study and the aging, demographics, and memory study. *J Gerontol B Psychol Sci Soc Sci.* (2011) 66:162–71. doi: 10.1093/geronb/gbr048
54. Meng Q, Wang H, Strauss J. Validation of neuropsychological tests for the China health and retirement longitudinal study harmonized cognitive assessment protocol. *Int Psychogeriatr.* (2019) 31:1709–19. doi: 10.1017/S1041610219000693
55. Luo Y, Pan X, Zhang Z. Productive activities and cognitive decline among older adults in China: evidence from the China health and retirement longitudinal study. *Soc Sci Med.* (2019) 229:96–105. doi: 10.1016/j.socscimed.2018.09.052
56. Baniak LM, Yang K, Choi J. Long sleep duration associated with increased frailty risk in older community-dwelling adults. *J Aging Health.* (2020) 32:42–51. doi: 10.1177/0898264318803470
57. Guo L, An L, Luo F. Social isolation, loneliness and functional disability in Chinese older women and men: a longitudinal study. *Age Ageing.* (2021) 50:1222–8. doi: 10.1093/ageing/afaa271
58. Aliberti MJR, Cenzer IS, Smith AK. Assessing risk for adverse outcomes in older adults: the need to include both physical frailty and cognition. *J Am Geriatr Soc.* (2019) 67:477–83. doi: 10.1111/jgs.15683
59. Fougere B, Daumas M, Lilamand M. Association between frailty and cognitive impairment: cross sectional data from Toulouse frailty day hospital. *J Am Med Dir Assoc.* (2017) 18:990. doi: 10.1177/1533317518791401
60. Kong L, Zhao H, Liu Y. Cognitive frailty of elderly patients with type 2 diabetes mellitus in the community and its influencing factors. *J Nurs.* (2020) 35:89–92. doi: 10.3870/j.issn.1001-4152.2020.07.089
61. Leng Y, Redline S, Stone KL, Ancoli-Israel S, Yaffe K. Objective napping, cognitive decline, and risk of cognitive impairment in older men. *Alzheimers Dement.* (2019) 15:1039–47. doi: 10.1016/j.jalz.2019.04.009
62. Li J, Cacchione PZ, Hodgson N, Riegel B, Keenan BT, Scharf MT. Afternoon napping and cognition in Chinese older adults: findings from the China health and retirement longitudinal study baseline assessment. *J Am Geriatr Soc.* (2017) 65:373–80. doi: 10.1111/jgs.14368
63. Song T, Yu K, Wang L, Xu L, Xu M, Peng Z, et al. Total sleep deprivation triggers greater activation in the parietal brain in the visual working memory updating processes: An event-related potentials study. *Front Neurosci.* (2022) 16:736437. doi: 10.3389/fnins.2022.736437
64. Honn KA, Hinson JM, Whitney P, Van Dongen HPA. Cognitive flexibility: a distinct element of performance impairment due to sleep deprivation. *Accid Anal Prev.* (2018) 126:191–7. doi: 10.1016/j.aap.2018.02.013
65. Aidman E, Jackson SA, Kleitman S. Effects of sleep deprivation on executive functioning, cognitive abilities, metacognitive confidence, and decision making. *Appl Cogn Psychol.* (2018) 33:188–200. doi: 10.1002/acp.3463
66. Cain SW, Silva EJ, Chang A, Ronda JM, Duffy JF. One night of sleep deprivation affects reaction time, but not interference or facilitation in a Stroop task. *Brain Cogn.* (2011) 76:0–42. doi: 10.1016/j.bandc.2011.03.005
67. Binks PG, Waters WF, Hurry M. Short-term total sleep deprivations does not selectively impair higher cortical functioning. *Sleep.* (1999) 22:328–34. doi: 10.1093/sleep/22.3.328
68. Corsi-Cabrera M, Sanchez AI, del-Rio-Portilla Y, Villanueva Y, Perez-Garci E. Effect of 38 h of total sleep deprivation on the waking EEG in women: sex differences. *Int J Psychophysiol.* (2003) 50:213–24. doi: 10.1016/s0167-8760(03)00168-5
69. Hajali V, Andersen ML, Negah SS, Sheibani V. Sex differences in sleep and sleep loss-induced cognitive deficits: the influence of gonadal hormones. *Horm Behav.* (2019) 108:50–61. doi: 10.1016/j.yhbeh.2018.12.013
70. Li P, Gao L, Yu L, Zheng X, Ulsa MC, Yang HW. Daytime napping and Alzheimer's dementia: a potential bidirectional relationship. *Alzheimers Dement.* (2023) 19:158–68. doi: 10.1002/alz.12636
71. Overton M, Sindi S, Basna R, Elmståhl S. Excessive sleep is associated with worse cognition, cognitive decline, and dementia in mild cognitive impairment. *Alzheimers Dement.* (2025) 17:93. doi: 10.1002/dad2.70093
72. Cai H, Su N, Li W, Li X, Xiao S, Sun L. Relationship between afternoon napping and cognitive function in the ageing Chinese population. *Gen Psychiatr.* (2021) 34:361. doi: 10.1136/gpsych-2020-100361
73. Magee CA, Huang XF, Iverson DC, Caputi P. Examining the pathways linking chronic sleep restriction to obesity. *J Obes.* (2010) 2010:1–8. doi: 10.1155/2010/821710