

Effects of nocturnal-diurnal sleep patterns and social participation factors on memory-related disorders in middle-aged and older adults

Keywords

social participation, CHARLS, sleep patterns, memory-related disorders, middle-aged and older adults

Abstract

Introduction

To investigate whether there is an association between nocturnal-daytime sleep patterns as well as social participation and memory-related categories of diseases in the middle-aged and elderly population in China.

Material and methods

This study was based on data from the Peking University China Health and Aging Tracking Survey (CHARLS), and a total of 19,023 study participants were analyzed in the year 2018 according to the inclusion and exclusion criteria. Logistic regression models were used to explore the relationship between different night-time-daytime sleep patterns, social participation, and memory-related categories of disorders step by step with adjustments for different covariates, and the odds ratios (ORs) and 95% confidence intervals (95% CI) were obtained.

Results

Logistic regression models gradually adjusted for confounding with nighttime sleep duration >5 h and daytime sleep duration ≤30 min group, Going to a sport, social, or other club activity and social participation as the protective factors of memory-related class of diseases. Stratified analyses found that among males, people living in rural areas, and married people, the group with >5 h of nighttime sleep and ≤30 min of daytime sleep was a protective factor for memory-related classes of diseases. Among women, people living in urban areas, and married people, social participation was a protective factor for memory-related diseases.

Conclusions

Nighttime sleep time >5 h, daytime sleep time ≤30 min group, and social participation are independent protective factors for memory-related class disorders in older adults.

Keywords: sleep patterns, social participation, memory-related disorders, CHARLS, middle-aged and older adults

1 **Title page**

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5 **Abstract:**

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27 independent protective factors for memory-related class disorders in older adults.

28 **Keywords:** CHARLS; Middle-aged and older adults; Nighttime sleep duration; Social
29 participation; Memory-related disorders

30 **Introduction**

31 Dementia, brain atrophy, and Parkinson's disease, which are increasingly prevalent as
32 populations age[1, 2]. These conditions lead to significant cognitive decline, loss of independence,
33 and heightened healthcare costs, thus imposing a substantial economic burden on society[3]. Current
34 diagnostic and therapeutic strategies predominantly revolve around cognitive assessments,
35 pharmacological interventions, and lifestyle modifications; however, these approaches often do not
36 substantially alter disease trajectory or enhance long-term patient outcomes. Consequently, there is
37 an urgent need to investigate alternative protective factors that may mitigate the risk of developing
38 these debilitating conditions.

39 Memory-related disorders such as Alzheimer's disease, cerebral atrophy, and Parkinson's
40 disease are closely linked to sleep duration in middle-aged and older adults. Numerous studies have

41 focused on the association between nighttime sleep duration or daytime sleep duration and memory-
42 related disorders[4-6]. However, there is a gap in the literature regarding the combined impact of
43 both nighttime and daytime sleep patterns on memory-related disorders in this population. It
44 highlighted the need for further research to explore the specific contributions of nocturnal and
45 diurnal sleep patterns to these disorders.

46 Social participation refers to maintaining social relationships and engaging in social activities.
47 It is one of the three major social policies advocated by the World Health Organization for active
48 aging[7, 8]. Social participation has been shown to have significant effects on both physical and
49 mental health outcomes in middle-aged and older adults[9]. Maintaining cognitive function and
50 preventing cognitive decline is a key focus in the field of aging research. Research has suggested
51 that social participation may play a protective role in cognitive function and memory-related
52 disorders in older adults. Engaging in social activities and maintaining social connections may help
53 to reduce the risk of cognitive decline and improve overall cognitive function[10, 11]. However,
54 more research is needed to explore the specific effects of social participation on memory-related
55 disorders in middle-aged and older adults.

56 This observational study aims to investigate the effects of nocturnal-diurnal sleep patterns and
57 social participation on memory-related disorders in middle-aged and older adults. By examining the
58 relationship between social participation and cognitive functioning, this study seeks to provide
59 valuable insights into the potential benefits of maintaining social relationships and engaging in
60 social activities for cognitive health in later life.

61 **Method:**

62 The dataset utilized in this investigation originates from the China Health and Retirement
63 Longitudinal Study (CHARLS), which was executed by Peking University[12]. This extensive
64 nationwide survey program was initiated by the National Development Research Institute (NDRI)
65 at Peking University and encompassed 150 counties across 28 provinces, including autonomous
66 regions and municipalities directly governed by the central authority of China. In summary,
67 participants aged middle-aged and older were randomly chosen for the baseline survey, which was
68 conducted from June 2011 to March 2012, employing a sampling methodology where the
69 probability of selection was directly proportional to the population size. For the purposes of this
70 analysis, data from the year 2018 were utilized. The CHARLS project received ethical approval
71 from the Ethics Committee of the Faculty of Medicine at Peking University. Furthermore, the
72 Biomedical Ethics Review Board of Peking University granted approval for the CHARLS study
73 (IRB00001052-11015), and all participants provided their written informed consent. The dataset
74 and associated information pertinent to this study are accessible for download via the CHARLS
75 project website.

76 Nighttime sleep duration was assessed by the question, “During the past month, how many
77 hours of actual sleep did you get on average per night?”, which specifically refers to actual sleep
78 duration rather than time spent in bed. Daytime napping was measured by the question, “During the
79 past month, how long did you take a nap after lunch?”. To evaluate the combined effects, sleep
80 patterns were categorized into four groups: (1) short night sleep (≤ 5 h) with short nap (≤ 30 min);
81 (2) short night sleep (≤ 5 h) with long nap (> 30 min); (3) normal night sleep (> 5 h) with short nap
82 (≤ 30 min); and (4) normal night sleep (> 5 h) with long nap (> 30 min). The 5-hour threshold for
83 nighttime sleep was adopted as it is a cutoff for increased risks of chronic diseases (e.g.,
84 hypertension and multimorbidity) among middle-aged and older Chinese populations[13]. The 30-

85 minute nap cutoff was used to distinguish "power naps" from prolonged naps associated with
86 adverse metabolic outcomes[14]. For social activity participation, respondents were asked whether
87 they had engaged in any of the following eight activities in the past month: visiting friends and
88 relatives, board games and recreation, unpaid help, physical activity, club activity, voluntary and
89 charitable activities, unpaid care for others, and continuing education. Participation in any of the
90 eight activities was categorized as participation in social activities, and ultimately, participation in
91 social activities was divided into two categories, i.e., non-participation in social activities and
92 participation in social activities[15]. Memory-related disorders, encompassing Alzheimer's disease,
93 brain atrophy, and Parkinson's disease, were identified based on physician-diagnosed conditions.
94 Following established protocols in large-scale epidemiological cohorts, participants were asked
95 whether a healthcare professional had ever informed them of having these specific conditions. This
96 self-reported medical diagnosis approach is a standard metric in the CHARLS database. Extensive
97 validation studies have demonstrated that such self-reported data maintain high consistency with
98 objective clinical records and medical claims[16-19].

99 The exclusion criteria applied in this study were as follows: Participants with missing
100 information on memory related diseases(n = 420). Participants with missing information on social
101 participation (n = 15). Participants with missing information on age (n = 64). Aged <45 years (n =
102 292). Absence of data regarding covariates (n = 1). Consequently, a final cohort comprising 19024
103 participants who satisfied these criteria was established (see Fig. 1). This study adhered to the cohort
104 study reporting guidelines set forth by the Strengthening the Reporting of Observational Studies in
105 Epidemiology (STROBE).

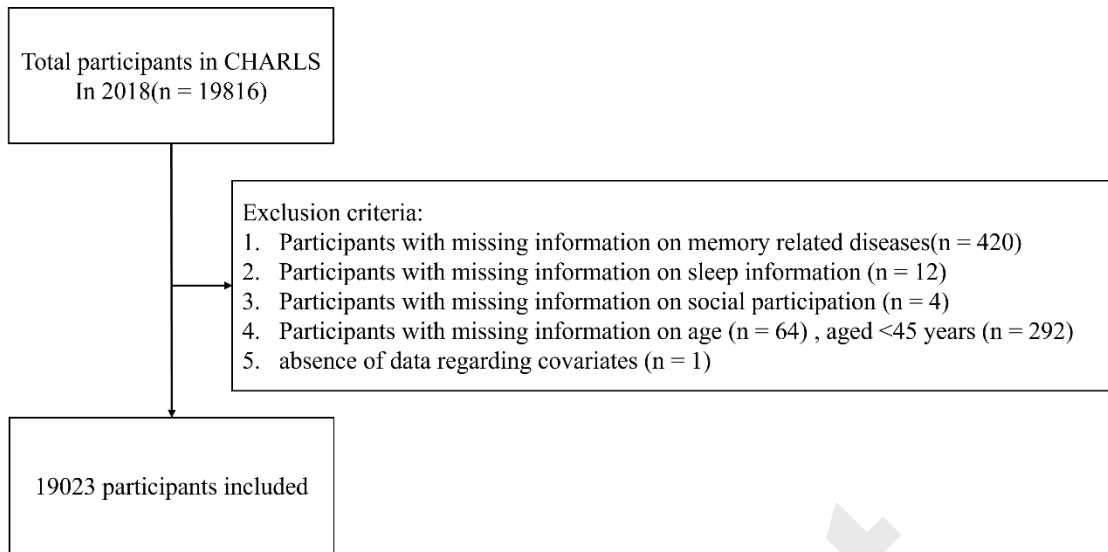
106 Continuous variables were expressed as the mean \pm standard deviation, while categorical
107 data were represented as counts accompanied by percentages. Statistical analyses were performed
108 utilizing EmpowerStats (X&Y Solutions, Inc., Boston, MA) and R software version 3.6.1. A two-
109 sided alpha level of 0.05 was established. Comparisons between groups of continuous variables
110 were carried out using the T-test, and chi-squared tests were employed for categorical variables.

111 Logistic regression modeling was used to explore the associations of the nocturnal and daytime
112 sleep patterns and social participation for having a memory-related disorder. Multiple models were
113 constructed, each adjusted for a different set of covariates, to gain insight into how these covariates
114 affect the observed associations. Subgroup analyses were used to examine the associations of the
115 nocturnal and daytime sleep patterns and social participation with memory-related disorders across
116 sociodemographic characteristics. To assess the potential impact of unmeasured confounding (e.g.,
117 past medical history or exercise habits), E-value analysis was performed for variables that showed
118 significant associations in the logistic regression models. The E-value is defined as the minimum
119 strength of association that an unmeasured confounder would need to have with both the exposure
120 and the outcome, conditional on the measured covariates, to fully explain away the observed risk
121 ratio[20]. A higher E-value indicates that a stronger unmeasured confounder would be required to
122 nullify the observed effect, thereby reflecting the robustness of the findings against potential omitted
123 variable bias

124

125 **Result**

126 **Figure 1. Flowchart of participant selection**



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Table 1. Patient Characteristics

Characteristic	Memory related diseases		P-value
	No	Yes	
Age (Mean \pm SD)	61.422 \pm 10.026	69.422 \pm 10.309	<0.001
Gender (%)			0.366
Female	9,716 (52.287%)	221 (50.113%)	
Male	8,866 (47.713%)	220 (49.887%)	
Living region (%)			0.920
Urban Community	7,460 (40.146%)	176 (39.909%)	
Rural Village	11,122 (59.854%)	265 (60.091%)	
Marital status (%)			<0.001
Non-married	2,697 (14.514%)	126 (28.571%)	
Married	15,885 (85.486%)	315 (71.429%)	
Smoke status (%)			0.195
No	10,686 (57.507%)	240 (54.422%)	
Yes	7,896 (42.493%)	201 (45.578%)	
Education status (%)			0.002
Illiterate	4493 (24.179%)	139 (31.519%)	
Primary school	8396 (45.184%)	185 (41.950%)	
Middle school or above	5,693 (30.637%)	117 (26.531%)	
Drinking status (%)			<0.001
No	12,204 (65.676%)	345 (78.231%)	
Yes	6,378 (34.324%)	96 (21.769%)	
Social participation (%)			<0.001
No	9,459 (50.904%)	271 (61.451%)	
Yes	9,124 (49.096%)	170 (38.549%)	

Interacting with friends			
No	12288 (66.129%)	317 (71.882%)	0.012
Yes	6294 (33.871%)	124 (28.118%)	
Playing Mahjong, chess, or cards			
No	15470 (83.253%)	386 (87.528%)	0.017
Yes	3112 (16.747%)	55 (12.472%)	
Providing help to family, friends, or neighbors who did not pay for the help			
No	15978 (85.986%)	396 (89.796%)	0.022
Yes	2604 (14.014%)	45 (10.204%)	
Going to a sport, social, or other club activity			
No	17343 (93.332%)	430 (97.506%)	<0.001
Yes	1239 (6.668%)	11 (2.494%)	
Participating in a community-related organization			
No	18128 (97.557%)	432 (97.959%)	0.588
Yes	454 (2.443%)	9 (2.041%)	
Doing volunteer or charity work			
No	18259 (98.262%)	436 (98.866%)	0.335
Yes	323 (1.738%)	5 (1.134%)	
The nocturnal and daytime sleep patterns (%)			
[1]nighttime sleep(≤5 h) with daytime napping(≤30 min)	3,739 (20.1%)	113 (25.6%)	<0.001
[2]nighttime sleep(≤5 h)	2,197 (11.8%)	78 (17.7%)	

with daytime napping(> 30 min)

[3]nighttime sleep(> 5 h)

6,668 (35.9%)

115 (26.1%)

with daytime napping(≤30 min)

[4]nighttime sleep(> 5 h)

5,979 (32.2%)

135 (30.6%)

with daytime napping(> 30 min)

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131 In table 1, we compared the differences between patients diagnosed with memory-related
132 disorders and undiagnosed individuals. *P*-values denote the statistical significance of differences
133 between participants with and without memory-related diseases. The data show that the mean age
134 was higher in the diagnosed group compared to undiagnosed group, a difference that was significant
135 ($P < 0.001$). Although the sex ratio did not differ significantly between the two groups ($P = 0.366$),
136 married status presented a significant difference, with the unmarried proportion being higher in the
137 diagnosed group ($P < 0.001$). In addition, the proportion of illiteracy was higher in the diagnostic
138 group, while the proportion of literacy above middle school was lower, which was significant ($P =$
139 0.006). Regarding drinking habits, non-drinkers accounted for a larger proportion in the diagnostic
140 group ($P < 0.001$). Regarding social participation, the proportion of those who did not participate in
141 social activities was significantly higher in the diagnostic group ($P < 0.001$). Finally, among the
142 variables related to sleep duration and distribution, individuals who slept ≤ 5 hours at night and
143 napped ≤ 30 minutes during the day were more represented in the diagnostic group, a
144 characteristic that was also significant ($P < 0.001$). Interacting with friends, playing mahjong, chess,
145 or playing cards, providing assistance to family, friends, or neighbors who have not paid for help,
146 and participating in sports, social, or other club activities show significant differences between
147 groups.
148

149 **Table 2. Association between the nocturnal and daytime sleep patterns and memory related diseases**

The nocturnal and daytime sleep patterns	Model 1			Model 2			Model 3		
	OR ¹	(95% CI ¹)	<i>P</i> -value	OR ¹	(95% CI ¹)	<i>P</i> -value	OR ¹	(95% CI ¹)	<i>P</i> -value
Nighttime sleep(≤5 h) with daytime napping(≤30 min)	Reference			Reference			Reference		
Nighttime sleep(≤5 h) with daytime napping(> 30 min)	1.175	(0.877, 1.576)	0.280	1.113	(0.827, 1.498)	0.480	1.117	(0.830, 1.504)	0.465
Nighttime sleep(> 5 h) with daytime napping(≤30 min)	0.571	(0.439, 0.742)	<0.001**	0.672	(0.514, 0.877)	0.003**	0.666	(0.510, 0.871)	0.003**
Nighttime sleep(> 5 h) with daytime napping(> 30 min)	0.747	(0.580, 0.962)	0.024*	0.776	(0.599, 1.005)	0.055	0.783	(0.605, 1.015)	0.065

150

Notes: **P* < 0.05, ***P* < 0.01; ¹OR = Odds Ratio, CI = Confidence Interval; Model 1 : no covariates were adjusted; Model 2 : adjusted for age, gender, and region; Model 3 : adjusted for age, gender, region, marital status, education, smoke status, and drinking status.

151 In Table 2, the group with nighttime sleep ≤ 5 h and daytime napping ≤ 30 min was used as the
152 reference. The results indicated that the pattern of nighttime sleep > 5 h with daytime napping ≤ 30
153 min was significantly associated with a reduced risk of memory-related diseases. Specifically, this
154 group presented an OR of 0.571 (95% CI: 0.439–0.742, $P < 0.001$) in Model 1 without adjustment
155 for covariates. After adjusting for age, gender, and living region in Model 2, the association
156 remained significant with an OR of 0.672 (95% CI: 0.514–0.877, $P = 0.003$). In the fully adjusted
157 Model 3, which further accounted for marital status, education, smoking, and drinking habits, the
158 protective effect persisted with an OR of 0.666 (95% CI: 0.510–0.871, $P = 0.003$).

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Table 3. Association between social participation and memory related diseases

Characteristics	Model 1			Model 2			Model 3		
	OR ¹	(95% CI ¹)	<i>P</i> -value	OR ¹	(95% CI ¹)	<i>P</i> -value	OR ¹	(95% CI ¹)	<i>P</i> -value
Social participation									
No		Reference			Reference			Reference	
Yes	0.650	(0.536, 0.790)	<0.001**	0.767	(0.630, 0.934)	0.008**	0.779	(0.639, 0.950)	0.014*
Interacting with friends									
No		Reference			Reference			Reference	
Yes	0.764	(0.619, 0.942)	0.012*	0.859	(0.695, 1.063)	0.162	0.875	(0.707, 1.083)	0.220
Playing Mahjong, chess, or cards									
No		Reference			Reference			Reference	
Yes	0.708	(0.533, 0.942)	0.018*	0.798	(0.597, 1.067)	0.128	0.803	(0.599, 1.075)	0.141
Providing help to family, friends, or neighbors who did not pay for the help									
No		Reference			Reference			Reference	
Yes	0.697	(0.511, 0.952)	0.023*	0.985	(0.717, 1.352)	0.924	1.023	(0.744, 1.406)	0.890
Going to a sport, social, or other club activity									
No		Reference			Reference			Reference	
Yes	0.358	(0.196, 0.653)	<0.001**	0.462	(0.252, 0.848)	0.013*	0.464	(0.253, 0.853)	0.013*

Notes: **P* < 0.05, ***P* < 0.01; ¹OR = Odds Ratio, CI = Confidence Interval; Model 1 : no covariates were adjusted; Model 2 : adjusted for age, gender, and region; Model 3 : adjusted for age, gender, region, marital status, education, smoke status, and drinking status.

160 Table 3 assesses the impact of social participation and its specific components on the risk of
161 memory-related disorders across three models. Social participation showed a consistent and
162 significant protective effect in all models. In the unadjusted model (Model 1), the OR for the social
163 participation group compared to the reference group was 0.650 (95% CI: 0.536–0.790, $P < 0.001$).
164 After adjusting for age, gender, and region in Model 2, the protective effect remained significant
165 with an OR of 0.767 (95% CI: 0.630–0.934, $P = 0.008$). In the fully adjusted Model 3, which further
166 accounted for marital status, education, and smoking and drinking status, the OR was 0.779 (95%
167 CI: 0.639–0.950, $P = 0.014$). Furthermore, among the specific types of activities, going to a sport,
168 social, or other club activity demonstrated the most substantial protective effect. In Model 1, this
169 activity was associated with a significant reduction in risk (OR: 0.358, 95% CI: 0.196–0.653, $P <$
170 0.001). This association persisted in the fully adjusted Model 3, with an OR of 0.464 (95% CI:
171 0.253–0.853, $P = 0.013$). These findings consistently suggest that both overall social participation
172 and specific engagement in club activities are independent protective factors against memory-
173 related disorders. Sensitivity analysis using E-values yielded robust results for the primary
174 significant predictors. Specifically, the E-values for nighttime sleep (> 5 h) with short napping
175 (≤ 30 min), social participation, and club activity were 2.37, 1.88, and 3.73, respectively. These
176 relatively high E-values suggest that any unmeasured confounding factors would need to have
177 substantial association strengths (ranging from 1.88 to 3.73) with both the exposures and the
178 memory-related disorders to negate the observed findings.
179

180

Table 4. Subgroup analysis of nighttime and daytime sleep patterns, social participation in memory related diseases.

Subgroup Characteristic	Nighttime sleep(> 5 h) with daytime napping(≤30 min)	Social participation	Going to a sport, social or other club activity
	OR ¹ (95% CI ¹) P-value	OR ¹ (95% CI ¹) P-value	OR ¹ (95% CI ¹) P-value
Gender (%)			
Female	0.706 (0.488, 1.020) 0.064	0.670 (0.505, 0.889) 0.006**	0.485 (0.236, 0.997) 0.049*
Male	0.609 (0.413, 0.900) 0.013*	0.900 (0.680, 1.191) 0.460	0.398 (0.126, 1.260) 0.117
Living region (%)			
Urban Community	0.678 (0.437, 1.051) 0.083	0.659 (0.479, 0.907) 0.011*	0.342 (0.139, 0.844) 0.020
Rural Village	0.659 (0.470, 0.925) 0.016*	0.879 (0.682, 1.134) 0.321	0.677 (0.298, 1.540) 0.353
Marital status (%)			
Non-married	1.065 (0.646, 1.754) 0.806	0.847 (0.583, 1.229) 0.382	0.640 (0.198, 2.071) 0.457
Married	0.551 (0.401, 0.755) <0.001**	0.756 (0.598, 0.957) 0.020*	0.415 (0.204, 0.847) 0.016*

181

Notes: * $P < 0.05$, ** $P < 0.01$; ¹OR = Odds Ratio, CI = Confidence Interval; covariates were adjusted.

182

183

184 Table 4 elucidates the findings of the stratified analyses conducted. It was observed that social
185 participation serves as a protective factor against memory-related diseases within the female
186 demographic ($P = 0.006$), the urban community ($P = 0.011$), as well as among married individuals
187 ($P = 0.020$). The study also found that sleeping more than five hours at night while taking no more
188 than a thirty-minute nap during the day was protective against memory-related disorders in a group
189 of male($P = 0.012$), Rural Village($P = 0.016$), and married individuals($P < 0.001$). Women living in
190 urban areas and married individuals who participate in sports clubs have a lower risk of developing
191 memory related diseases.

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193 Discussion

194 Understanding memory-related diseases is crucial due to their profound impact on individuals
195 and society. Our research offers advantages such as using a well-defined population and reliable
196 measures of sleep and social participation. Key findings from this study reveal that adequate
197 nighttime sleep and limited daytime sleep, along with active social involvement, may serve as
198 protective factors against dementia, brain atrophy, and Parkinson's disease.

199 By examining variables such as age, marital status, educational level, drinking habits, social
200 engagement, and sleep patterns, the study seeks to identify significant distinctions that could
201 enhance early detection and intervention strategies. The findings highlight the potential for targeted
202 approaches in addressing the complex interplay of these factors in the context of memory disorders,
203 thereby contributing to the broader understanding of risk factors and promoting timely therapeutic
204 interventions. Previous research has often focused on singular aspects, such as age or education
205 level, without considering the interconnectedness of various social determinants. Our findings,
206 particularly the significant associations between lower literacy rates, increased age, and diminished
207 social participation with the prevalence of memory disorders, align with the conclusions drawn by
208 other studies that highlight the multifactorial nature of cognitive decline in aging populations[21].

209 The implications of our findings hold significant potential for clinical practice and policy
210 formulation. Identifying the demographic and lifestyle characteristics that predispose individuals to
211 memory-related disorders can guide targeted screening efforts, ultimately leading to earlier
212 diagnosis and intervention. For instance, the observed higher prevalence of memory disorders
213 among individuals with lower literacy rates emphasizes the necessity for educational initiatives
214 aimed at improving cognitive resilience. Furthermore, the strong association between poor sleep
215 patterns and cognitive impairments suggests that sleep hygiene interventions could be integrated
216 into treatment plans for patients diagnosed with memory disorders, potentially mitigating cognitive
217 decline[22]. These findings underline the importance of adopting a holistic approach to treatment,
218 which acknowledges the interplay between lifestyle factors and cognitive health.

219 Nighttime sleep duration and daytime nap duration have a significant effect on cognitive
220 function in older adults. Particularly in the Chinese elderly population, researchers found a U-shaped
221 relationship between nighttime sleep duration and mild cognitive impairment (MCI)[23]. These
222 findings emphasize the importance of appropriate nighttime sleep versus daytime naps in preventing
223 and delaying cognitive decline. Sleep of more than 5 hours at night ensures the integrity of the
224 underlying sleep cycle, especially slow wave sleep (SWS), which plays a key role in synaptic steady
225 state regulation and metabolic waste removal[24]. Control of daytime nap time in less than 30
226 minutes has a clear cognitive compensatory advantage. Studies have confirmed that ≤ 30 minutes of
227 nap can effectively relieve subjective drowsiness, improve positive mood, and not trigger significant
228 sleep. Improved memory encoding capability in inert conditions[25]. The combination effectively
229 relieves daytime sleep stress without interfering with the output of endogenous circadian rhythms.
230 This strategy both reduces the accumulative risk of beta-amyloid and optimizes neurophysiological
231 stability, thus Knowledge functions produce synergistic protective effects[26]. Consistent with
232 existing literature recommending regular, adequate nocturnal sleep and limited daytime nap duration
233 for older adults[27], our findings reinforce these patterns within the Chinese context. Notably,
234 compared to Western studies that emphasize sleep continuity, Chinese older adults exhibit a higher
235 prevalence of short daytime napping, a behavior likely influenced by deeply rooted cultural habits.
236 Furthermore, the protective effects of social participation observed in this study reveal significant

237 disparities across gender and urban-rural settings. The unique family structures and social models
238 inherent to China underline the critical importance of localized interventions. Such findings suggest
239 that public health strategies must move beyond universal guidelines to incorporate socio-cultural
240 and demographic nuances to effectively combat memory-related diseases.

241 Social engagement, another important factor, has also been suggested to have a protective
242 effect against memory disorders. Social engagement provides cognitive stimulation and emotional
243 support, which helps maintain and enhance cognitive functioning in older adults. Studies have
244 shown that older adults with high levels of social engagement have slower rates of cognitive decline
245 and are associated with a lower risk of memory-based disorders[28, 29]. Among men, people living
246 in rural areas, and married people, nighttime sleep duration of more than 5 hours combined with
247 daytime naps of no more than 30 minutes was recognized as a protective factor for memory-related
248 classes of disorders. This finding echoes previous research that proper nighttime sleep and daytime
249 naps have a positive impact on cognitive functioning[30-32]. Social engagement was found to be a
250 protective factor for memory-related disorders among women, urban-dwelling populations, and
251 married individuals. This may be related to the ability of social engagement to provide cognitive
252 stimulation and emotional support, thereby maintaining and enhancing cognitive functioning.
253 Physical activity is the most important modifiable risk factor in preventing Alzheimer's disease and
254 related dementia. Those who actively participate in sports clubs are more likely to maintain physical
255 activity, thereby reducing their risk of developing memory related diseases[33-35]. Existing
256 research suggests that gender and living environment may regulate the health effects of sleep and
257 social engagement through bio-psycho-social mechanisms. Women are more susceptible to social
258 support buffer stress[36]. Urban residents are dense in social resources, so that social participation
259 is more effective, thus playing a stronger protective role[37].

260 Despite the insights provided, several limitations must be acknowledged. First, the cross-
261 sectional design precludes the establishment of temporal causality between sleep patterns, social
262 participation, and memory-related disorders. Future research utilizing longitudinal cohorts is
263 warranted to more rigorously validate these causal pathways. Second, variables were derived from
264 self-reported data, which may introduce recall or social desirability bias. Incorporating objective
265 measures, such as polysomnography or clinical diagnostic records, in future studies would further
266 enhance the precision and reliability of the findings.

267 Nevertheless, these findings emphasize that protective factors for memory-related disorders
268 may vary significantly across populations and lifestyle habits. Therefore, these differences must be
269 integrated into the development of public health prevention strategies to ensure interventions are
270 effectively targeted. For instance, for men and rural-dwelling older adults, priority should be given
271 to optimizing sleep quality and daytime napping habits. Conversely, for women and urban-dwelling
272 older adults, fostering social participation and community engagement may yield more substantial
273 neuroprotective benefits.

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276 **Ethical approval**

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278 **Conflict of interest**

279 The authors declare no conflict of interest.

280

281

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Sleep patterns (night sleep >5h,
daytime nap ≤30min)



Social participation
(elderly people socializing, club
activities, exercise)



**Memory-related
disorders**

Sleep pattern and social participation as protective factors

Preprint